



Cochrane
Library

Cochrane Database of Systematic Reviews

Foot orthoses for treating paediatric flat feet (Review)

Evans AM, Rome K, Carroll M, Hawke F

Evans AM, Rome K, Carroll M, Hawke F.
Foot orthoses for treating paediatric flat feet.
Cochrane Database of Systematic Reviews 2022, Issue 1. Art. No.: CD006311.
DOI: [10.1002/14651858.CD006311.pub4](https://doi.org/10.1002/14651858.CD006311.pub4).

www.cochranelibrary.com

TABLE OF CONTENTS

ABSTRACT	1
PLAIN LANGUAGE SUMMARY	2
SUMMARY OF FINDINGS	4
BACKGROUND	15
OBJECTIVES	16
METHODS	16
RESULTS	19
Figure 1.	20
Figure 2.	25
Figure 3.	28
Figure 4.	28
Figure 5.	29
Figure 6.	30
DISCUSSION	30
AUTHORS' CONCLUSIONS	33
ACKNOWLEDGEMENTS	33
REFERENCES	35
CHARACTERISTICS OF STUDIES	40
DATA AND ANALYSES	71
Analysis 1.1. Comparison 1: Custom foot orthoses (CFOs) versus shoes for asymptomatic flat feet, Outcome 1: Proportion without pain	72
Analysis 1.2. Comparison 1: Custom foot orthoses (CFOs) versus shoes for asymptomatic flat feet, Outcome 2: Withdrawal due to adverse events	72
Analysis 2.1. Comparison 2: Prefabricated foot orthoses (PFOs) versus shoes in asymptomatic flat feet, Outcome 1: Proportion without pain	73
Analysis 2.2. Comparison 2: Prefabricated foot orthoses (PFOs) versus shoes in asymptomatic flat feet, Outcome 2: Withdrawal due to adverse events	73
Analysis 3.1. Comparison 3: CFOs versus PFOs in asymptomatic flat feet, Outcome 1: Pain	74
Analysis 3.2. Comparison 3: CFOs versus PFOs in asymptomatic flat feet, Outcome 2: Withdrawal due to adverse events	74
Analysis 4.1. Comparison 4: CFOs versus shoes in juvenile idiopathic arthritis (JIA), Outcome 1: Pain	75
Analysis 4.2. Comparison 4: CFOs versus shoes in juvenile idiopathic arthritis (JIA), Outcome 2: Function	75
Analysis 4.3. Comparison 4: CFOs versus shoes in juvenile idiopathic arthritis (JIA), Outcome 3: Quality of life	76
Analysis 4.4. Comparison 4: CFOs versus shoes in juvenile idiopathic arthritis (JIA), Outcome 4: Treatment success (gait parameters)	76
Analysis 4.5. Comparison 4: CFOs versus shoes in juvenile idiopathic arthritis (JIA), Outcome 5: Withdrawal due to adverse events	76
Analysis 5.1. Comparison 5: PFOs versus shoes in JIA, Outcome 1: Pain	77
Analysis 5.2. Comparison 5: PFOs versus shoes in JIA, Outcome 2: Function	78
Analysis 5.3. Comparison 5: PFOs versus shoes in JIA, Outcome 3: Quality of life	78
Analysis 5.4. Comparison 5: PFOs versus shoes in JIA, Outcome 4: Treatment success (Timed walking)	78
Analysis 5.5. Comparison 5: PFOs versus shoes in JIA, Outcome 5: Withdrawal due to adverse events	79
Analysis 6.1. Comparison 6: CFOs versus PFOs in JIA, Outcome 1: Pain	79
Analysis 6.2. Comparison 6: CFOs versus PFOs in JIA, Outcome 2: Function	80
Analysis 6.3. Comparison 6: CFOs versus PFOs in JIA, Outcome 3: Quality of life	80
Analysis 6.4. Comparison 6: CFOs versus PFOs in JIA, Outcome 4: Treatment success (timed walking)	80
Analysis 6.5. Comparison 6: CFOs versus PFOs in JIA, Outcome 5: Withdrawal due to adverse events	81
Analysis 7.1. Comparison 7: PFOs versus shoes in symptomatic flat feet, Outcome 1: Function	82
Analysis 7.2. Comparison 7: PFOs versus shoes in symptomatic flat feet, Outcome 2: Quality of life	83
Analysis 8.1. Comparison 8: CFOs versus shoes in DCD flat feet, Outcome 1: Gait parameters	83
Analysis 9.1. Comparison 9: Anti-pronation taping versus sham taping, Outcome 1: Balance test (SEBT) at 4 weeks	85
Analysis 9.2. Comparison 9: Anti-pronation taping versus sham taping, Outcome 2: Agility test (Illinois Agility Test) at 4 weeks ..	85
Analysis 9.3. Comparison 9: Anti-pronation taping versus sham taping, Outcome 3: Vertical jump height (cm)	85

Analysis 10.1. Comparison 10: Neuromuscular electrical stimulation (NMES) versus sham NMES, Outcome 1: Navicular height (mm)	87
Analysis 10.2. Comparison 10: Neuromuscular electrical stimulation (NMES) versus sham NMES, Outcome 2: Staheli's arch index (mm)	87
Analysis 10.3. Comparison 10: Neuromuscular electrical stimulation (NMES) versus sham NMES, Outcome 3: Calcaneal inclination angle (degrees)	88
Analysis 10.4. Comparison 10: Neuromuscular electrical stimulation (NMES) versus sham NMES, Outcome 4: Talus second metatarsal angle (degrees)	88
Analysis 10.5. Comparison 10: Neuromuscular electrical stimulation (NMES) versus sham NMES, Outcome 5: Talo-navicular coverage angle	89
ADDITIONAL TABLES	90
APPENDICES	97
FEEDBACK	104
WHAT'S NEW	106
HISTORY	106
CONTRIBUTIONS OF AUTHORS	106
DECLARATIONS OF INTEREST	106
SOURCES OF SUPPORT	106
DIFFERENCES BETWEEN PROTOCOL AND REVIEW	107
INDEX TERMS	107

[Intervention Review]

Foot orthoses for treating paediatric flat feet

Angela M Evans¹, Keith Rome², Matthew Carroll³, Fiona Hawke⁴

¹Discipline of Podiatry, School of Allied Health, Human Services and Sport, La Trobe University, Melbourne, Australia. ²Division of Rehabilitation & Occupation Studies, AUT University, Auckland 1020, New Zealand. ³Department of Podiatry, School of Clinical Sciences, Auckland University of Technology, Auckland, New Zealand. ⁴School of Health Sciences, Faculty of Health and Medicine, The University of Newcastle, Ourimbah, Australia

Contact: Angela M Evans, angela.evans@latrobe.edu.au.**Editorial group:** Cochrane Musculoskeletal Group.**Publication status and date:** New search for studies and content updated (no change to conclusions), published in Issue 1, 2022.**Citation:** Evans AM, Rome K, Carroll M, Hawke F. Foot orthoses for treating paediatric flat feet. *Cochrane Database of Systematic Reviews* 2022, Issue 1. Art. No.: CD006311. DOI: [10.1002/14651858.CD006311.pub4](https://doi.org/10.1002/14651858.CD006311.pub4).

Copyright © 2022 The Cochrane Collaboration. Published by John Wiley & Sons, Ltd.

ABSTRACT

Background

Paediatric flat feet are a common presentation in primary care; reported prevalence approximates 15%. A minority of flat feet can hurt and limit gait. There is no optimal strategy, nor consensus, for using foot orthoses (FOs) to treat paediatric flat feet.

Objectives

To assess the benefits and harms of foot orthoses for treating paediatric flat feet.

Search methods

We searched CENTRAL, MEDLINE, and Embase to 01 September 2021, and two clinical trials registers on 07 August 2020.

Selection criteria

We identified all randomised controlled trials (RCTs) of FOs as an intervention for paediatric flat feet. The outcomes included in this review were pain, function, quality of life, treatment success, and adverse events. Intended comparisons were: any FOs versus sham, any FOs versus shoes, customised FOs (CFOs) versus prefabricated FOs (PFOs).

Data collection and analysis

We followed standard methods recommended by Cochrane.

Main results

We included 16 trials with 1058 children, aged 11 months to 19 years, with flexible flat feet. Distinct flat foot presentations included asymptomatic, juvenile idiopathic arthritis (JIA), symptomatic and developmental co-ordination disorder (DCD). The trial interventions were FOs, footwear, foot and rehabilitative exercises, and neuromuscular electrical stimulation (NMES). Due to heterogeneity, we did not pool the data. Most trials had potential for selection, performance, detection, and selective reporting bias. No trial blinded participants. We present the results separately for asymptomatic (healthy children) and symptomatic (children with JIA) flat feet.

The certainty of evidence was very low to low, downgraded for bias, imprecision, and indirectness.

Three comparisons were evaluated across trials: CFO versus shoes; PFO versus shoes; CFO versus PFO.

Asymptomatic flat feet

1. *CFOs versus shoes* (1 trial, 106 participants): low-quality evidence showed that CFOs result in little or no difference in the proportion without pain (10-point visual analogue scale (VAS)) at one year (risk ratio (RR) 0.85, 95% confidence interval (CI) 0.67 to 1.07); absolute decrease (11.8%, 95% CI 4.7% fewer to 15.8% more); or on withdrawals due to adverse events (RR 1.05, 95% CI 0.94 to 1.19); absolute effect (3.4% more, 95% CI 4.1% fewer to 13.1% more).

2. *PFOs versus shoes* (1 trial, 106 participants): low to very-low quality evidence showed that PFOs result in little or no difference in the proportion without pain (10-point VAS) at one year (RR 0.94, 95% CI 0.76 to 1.16); absolute effect (4.7% fewer, 95% CI 18.9% fewer to 12.6% more); or on withdrawals due to adverse events (RR 0.99, 95% CI 0.79 to 1.23).

3. *CFOs versus PFOs* (1 trial, 108 participants): low-quality evidence found no difference in the proportion without pain at one year (RR 0.93, 95% CI 0.73 to 1.18); absolute effect (7.4% fewer, 95% CI 22.2% fewer to 11.1% more); or on withdrawal due to adverse events (RR 1.00, 95% CI 0.90 to 1.12).

Function and quality of life (QoL) were not assessed.

Symptomatic (JIA) flat feet

1. *CFOs versus shoes* (1 trial, 28 participants, 3-month follow-up): very low-quality evidence showed little or no difference in pain (0 to 10 scale, 0 no pain) between groups (MD -1.5, 95% CI -2.78 to -0.22). Low-quality evidence showed improvements in function with CFOs (Foot Function Index - FFI disability, 0 to 100, 0 best function; MD -18.55, 95% CI -34.42 to -2.68), child-rated QoL (PedsQL, 0 to 100, 100 best quality; MD 12.1, 95% CI -1.6 to 25.8) and parent-rated QoL (PedsQL MD 9, 95% CI -4.1 to 22.1) and little or no difference between groups in treatment success (timed walking; MD -1.33 seconds, 95% CI -2.77 to 0.11), or withdrawals due to adverse events (RR 0.58, 95% CI 0.11 to 2.94); absolute difference (9.7% fewer, 20.5% fewer to 44.8% more).

2. *PFOs versus shoes* (1 trial, 25 participants, 3-month follow-up): very low-quality evidence showed little or no difference in pain between groups (MD 0.02, 95% CI -1.94 to 1.98). Low-quality evidence showed no difference between groups in function (FFI-disability MD -4.17, 95% CI -24.4 to 16.06), child-rated QoL (PedsQL MD -3.84, 95% CI -19 to 11.33), or parent-rated QoL (PedsQL MD -0.64, 95% CI -13.22 to 11.94).

3. *CFOs versus PFOs* (2 trials, 87 participants): low-quality evidence showed little or no difference between groups in pain (0 to 10 scale, 0 no pain) at 3 months (MD -1.48, 95% CI -3.23 to 0.26), function (FFI-disability MD -7.28, 95% CI -15.47 to 0.92), child-rated QoL (PedsQL MD 8.6, 95% CI -3.9 to 21.2), or parent-rated QoL (PedsQL MD 2.9, 95% CI -11 to 16.8).

Authors' conclusions

Low to very low-certainty evidence shows that the effect of CFOs (high cost) or PFOs (low cost) versus shoes, and CFOs versus PFOs on pain, function and HRQoL is uncertain. This is pertinent for clinical practice, given the economic disparity between CFOs and PFOs. FOs may improve pain and function, versus shoes in children with JIA, with minimal delineation between costly CFOs and generic PFOs.

This review updates that from 2010, confirming that in the absence of pain, the use of high-cost CFOs for healthy children with flexible flat feet has no supporting evidence, and draws very limited conclusions about FOs for treating paediatric flat feet.

The availability of normative and prospective foot development data, dismisses most flat foot concerns, and negates continued attention to this topic. Attention should be re-directed to relevant paediatric foot conditions, which cause pain, limit function, or reduce quality of life. The agenda for researching asymptomatic flat feet in healthy children must be relegated to history, and replaced by a targeted research rationale, addressing children with indisputable foot pathology from discrete diagnoses, namely JIA, cerebral palsy, congenital talipes equino varus, trisomy 21 and Charcot Marie Tooth. Whether research resources should continue to be wasted on studying flat feet in healthy children that do not hurt, is questionable. Future updates of this review will address only relevant paediatric foot conditions.

PLAIN LANGUAGE SUMMARY

Foot orthoses for treating flat feet in children

Review question

What are the benefits and harms of using foot orthoses (shoe inserts) to treat flat feet in children?

Background

Children with flat feet have a lower foot arch. When the child is standing, the foot arch looks flat against the floor, and may roll inwards, and even touch the floor. Sometimes, flat feet can cause pain, or change the way a child walks.

There are many types of non-surgical treatments for flat feet, but unless painful, most children do not need any treatment.

Foot orthoses (FOs) or shoe inserts, muscle stretching, footwear selection, physical activity modification, and reducing body weight, may be part of an overall foot and activity management. The short-term use of medication for pain and inflammation may be prescribed.

Study characteristics

This Cochrane Review is current to September 2021. There are 16 studies (1058 children, aged 11 months to 19 years) including three groups - healthy children with painless flat feet; children with arthritis and painful flat feet; others (developmental coordination disorder; painful flat feet). The studies were conducted across the USA, Australia, India, Iran, Turkey, UK, and Republic of Korea. We found information about footwear, exercises, and different types of foot orthoses.

Results:

Comparing custom foot orthoses (CFOs) to shoes in painless flat feet:

Proportion without pain (1 trial, 106 children) at 12 months:

12% fewer children with CFOs were without pain (ranging from 26% fewer to 5.5% more)

67 out of 100 children were without pain with CFOs compared to 79 children out of 100 with shoes.

Withdrawal due to side effects (3 trials, 211 children):

3% more children with CFOs withdrew from treatment due to side effects (ranging from 4% fewer to 13% more) .

72 out of 100 children withdrew from treatment with CFOs compared with 69 children out of 100 with shoes.

Comparing prefabricated foot orthoses (PFOs) to shoes in painless flat feet

Proportion without pain (1 trial, 106 children) at 12 months:

5% fewer children with PFOs were without pain (ranging from 18.9% fewer to 12.6% more)

74 out of 100 children were without pain with PFOs compared to 79 out of 100 children with shoes.

Withdrawal due to side effects (4 trials, 338 children):

0.7% fewer children with PFOs withdrew from treatment due to side effects (ranging from 15.2% fewer to 16.6% more).

71 out of 100 children withdrew from treatment with PFOs compared with 72 out of 100 children with shoes.

Comparing CFOs to PFOs in painless flat feet

Proportion without pain (1 trial, 106 children) at 12 months:

7% fewer children with CFOs were without pain (ranging from 22.2% fewer to 11.1% more)

68 out of 100 children were without pain with CFOs compared to 74 out of 100 children with PFOs.

Withdrawal due to side effects (1 trial, 118 children):

0% fewer children withdrew from treatment with CFOs due to side effects (ranging from 9.2% fewer to 11% more).

91 out of 100 children withdrew from treatment with CFOs compared with 91 out of 100 children with PFOs.

Function, quality of life, treatment success and side effects were not reported in these trials

Quality of the evidence

In healthy children with painless flat feet, low to very low-quality evidence shows that compared to shoes, CFOs and PFOs result in no difference in the proportion without pain or withdrawal due to side effects from treatment. The quality of the evidence is very low to low, weakening conclusions. We downgraded the certainty of the evidence because the trials were poorly conducted and there were not enough data.

SUMMARY OF FINDINGS

Summary of findings 1. Customised foot orthoses compared to shoes in children with asymptomatic flat feet

Customised foot orthoses compared to shoes in children with asymptomatic flat feet

Patient or population: children with asymptomatic flat feet

Setting: outpatient hospital clinic

Intervention: customised foot orthoses (CFO)

Comparison: shoes

Outcomes	Relative effect (95% CI)	Anticipated absolute effects* (95% CI)			Certainty of the evidence (GRADE)	What happens
		With shoes (N = 52)	With CFOs (N = 54)	Difference (absolute)		
Pain (measured as proportion with pain) follow-up: 12 months N° of participants: 106 (1 RCT)	RR 0.85 (0.67 to 1.07)	78.8%	67% (52.8% to 84.4%)	11.8% fewer (4.7% fewer to 15.8% more)	⊕⊕⊕⊕ Low ^{a,b}	CFOs may result in little to no difference in the proportion of children reporting pain (absolute reduction of 11.8% (4.7% fewer to 15.8% more))
Function or disability	-	-	-	-	-	not reported
Quality of life	-	-	-	-	-	not reported
Treatment success	-	-	-	-	-	not reported
Withdrawal due to adverse events follow-up: 3 months to 4 months N° of participants: 211 (3 RCTs)	RR 1.05 (0.94 to 1.19)	68.9%	72.3% (64.7% to 82%)	3.4% more (4.1% fewer to 13.1% more)	⊕⊕⊕⊕ Low ^{a,b}	The evidence suggests that CFOs result in little to no difference in withdrawal due to adverse events (absolute effect 3.4% more (4.1% fewer to 13.1% more))
Adverse events	-	-	-	-	-	not reported
Serious adverse events	-	-	-	-	-	not reported

***The risk in the intervention group** (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: Confidence interval; **RR:** Risk ratio

GRADE Working Group grades of evidence

High certainty: we are very confident that the true effect lies close to that of the estimate of the effect

Moderate certainty: we are moderately confident in the effect estimate; the true effect is likely to be close to the estimate of the effect, but there is a possibility that it is substantially different

Low certainty: our confidence in the effect estimate is limited; the true effect may be substantially different from the estimate of the effect

Very low certainty: we have very little confidence in the effect estimate; the true effect is likely to be substantially different from the estimate of effect

^aDowngraded for bias (participants, parents, and examiners were aware of treatment, which may have impacted self-reported outcomes; subgroup analysis of those with pain was conducted (post hoc))

^bDowngraded for imprecision due to wide confidence intervals including both an increase and decrease in the effect estimate

Summary of findings 2. Prefabricated foot orthoses compared to shoes in children with asymptomatic flat feet

Prefabricated foot orthoses compared to shoes in children with asymptomatic flat feet

Patient or population: children with asymptomatic flat feet

Setting: outpatient hospital clinic

Intervention: prefabricated foot orthoses (PFO)

Comparison: shoes

Outcomes	Relative effect (95% CI)	Anticipated absolute effects* (95% CI)			Certainty of the evidence (GRADE)	What happens
		With shoes (N = 52)	With PFOs (N = 54)	Difference (absolute)		
Pain (measured as proportion with pain) follow-up: 12 months N ^o of participants: 106 (1 RCT)	RR 0.94 (0.76 to 1.16)	78.8%	74.1% (59.9 to 91.5)	4.7% fewer (18.9% fewer to 12.6% more)	⊕⊕⊕⊕ Low ^{a,b}	PFOs likely result in little to no difference in the proportion of children reporting pain, absolute reduction 4.7% (18.9% fewer to 12.6% more)
Function or disability	-	-	-	-	-	not reported
Quality of life	-	-	-	-	-	not reported
Treatment success	-	-	-	-	-	not reported

Withdrawal due to adverse events	RR 0.99 (0.79 to 1.23)	72.3%	71.6% (57.1% to 88.9%)	0.7% fewer (15.2% fewer to 16.6% more)	⊕⊕⊕⊕ Very low ^{a,b,c}	We are uncertain of the effects of PFOs on withdrawal due to adverse events. Absolute reduction 0.7% (15.2 fewer to 16.6 more)
follow-up: 12 months						
N ^o of participants: 338 (4 RCTs)						
Adverse events	-	-	-	-	-	not reported
Serious adverse events	-	-	-	-	-	not reported

***The risk in the intervention group** (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: Confidence interval; **RR:** Risk ratio

GRADE Working Group grades of evidence

High certainty: we are very confident that the true effect lies close to that of the estimate of the effect

Moderate certainty: we are moderately confident in the effect estimate; the true effect is likely to be close to the estimate of the effect, but there is a possibility that it is substantially different

Low certainty: our confidence in the effect estimate is limited; the true effect may be substantially different from the estimate of the effect

Very low certainty: we have very little confidence in the effect estimate; the true effect is likely to be substantially different from the estimate of effect

^aDowngraded for bias, (performance, attrition, other bias), participants, parents, and examiners not blinded; pain only assessed post hoc, as subgroup analysis; high attrition in some trials (notably [Gould 1989](#))

^bDowngraded for imprecision; wide 95% CI for intervention

^cDowngraded for indirectness; variably aged participant samples between studies

Summary of findings 3. Custom foot orthoses compared to prefabricated foot orthoses for children with asymptomatic flat feet

Custom foot orthoses compared to prefabricated foot orthoses for children with asymptomatic flat feet

Patient or population: children with asymptomatic flat feet

Setting: outpatient clinics

Intervention: customised foot orthoses (CFO)

Comparison: prefabricated foot orthoses (PFO)

Outcomes	Relative effect (95% CI)	Anticipated absolute effects* (95% CI)			Certainty of the evidence (GRADE)	What happens
		With PFOs (N = 54)	With CFOs (N = 54)	Difference (absolute)		

Pain (measured as proportion with pain) follow-up: 12 months Nº of participants: 108 (1 RCT)	RR 0.93 (0.73 to 1.18)	74%	68% (51.9% to 85.2%)	7.4% fewer (22.2% fewer to 11.1% more)	⊕⊕○○ Low ^{a,b}	CFOs likely results in little to no difference in the proportion of children reporting pain. Absolute reduction 7.4% (22.2 % fewer to 11.1 % more)
Function or disability	-	-	-	-	-	not reported
Quality of life	-	-	-	-	-	not reported
Treatment success	-	-	-	-	-	not reported
Withdrawal due to adverse events follow up: 12 months Nº of participants: 118 (1 RCT)	RR 1.00 (0.90 to 1.12)	91.5%	91.5% (82.4% to 100%)	0.0% fewer (9.2% fewer to 11% more)	⊕⊕○○ Low ^{a,b}	The evidence suggests that CFOs do not increase or reduce withdrawal due to adverse events.
Adverse events	-	-	-	-	-	not reported
Serious adverse events	-	-	-	-	-	not reported

***The risk in the intervention group** (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: Confidence interval; **RR:** Risk ratio

GRADE Working Group grades of evidence

High certainty: we are very confident that the true effect lies close to that of the estimate of the effect

Moderate certainty: we are moderately confident in the effect estimate; the true effect is likely to be close to the estimate of the effect, but there is a possibility that it is substantially different

Low certainty: our confidence in the effect estimate is limited; the true effect may be substantially different from the estimate of the effect

Very low certainty: we have very little confidence in the effect estimate; the true effect is likely to be substantially different from the estimate of effect

^aDowngraded for bias, (performance, other bias), participants, parents, and examiners not blinded; pain only assessed post hoc, as subgroup analysis

^bDowngraded for imprecision; wide 95% CI for CFO as intervention

Summary of findings 4. Custom foot orthoses compared to shoes in children with juvenile idiopathic arthritis and flat feet

Custom foot orthoses compared to shoes in children with juvenile idiopathic arthritis and flat feet

Patient or population: children with juvenile idiopathic arthritis (JIA) and flat feet

Setting: outpatient rheumatology clinics
Intervention: custom foot orthoses (CFO)
Comparison: shoes

Outcomes	Relative effect (95% CI)	Anticipated absolute effects* (95% CI)			Certainty of the evidence (GRADE)	What happens
		With shoes (N = 13)	With CFOs (N = 15)	Difference		
Pain (measured on 0 to 10-point VAS; lower = less pain) follow-up: 3 months N° of participants: 28 (1 RCT)		The mean pain with shoes was 2.82 points	The mean pain with CFOs was 1.32 points	MD 1.5 points lower (2.78 points lower to 0.22 points lower)	⊕⊕⊕⊕ Very low ^{a,b,c}	CFOs likely results in little to no difference in pain.
Function or disability (measured on 0 to 100-point FFI; 0 = no disability) follow-up: 3 months N° of participants: 28 (1 RCT)		The mean FFI score with shoes was 34.15 points	The mean FFI score with CFOs was 15.6 points	MD 18.55 points lower (34.42 points lower to 2.68 points lower)	⊕⊕⊕⊕ Low ^{a,b}	CFOs may result in a clinically important improvement in function or disability.
Quality of life (child-rated) (measured on 0 to 100-point PedsQL; higher score = better QoL) follow-up: 3 months N° of participants: 25 (1 RCT)		The mean child-rated PedQL score with shoes was 59.78 points	The mean child-rated PedQL score with CFOs was 47.68 points	MD 12.1 points higher (1.6 points lower to 25.8 points higher)	⊕⊕⊕⊕ Low ^{a,c}	CFOs may result in a clinically important improvement in child-rated QoL.
Quality of life (parent-rated) (measured on 0 to 100-point PedsQL; higher score = better QoL) follow up: 3 months N° of participants: 26 (1 RCT)		The mean parent-rated PedQL score with shoes was 55.95 points	The mean parent-rated PedQL score with CFOs was 46.94 points	MD 9.01 points higher (4.08 points lower to 22.1 points higher)	⊕⊕⊕⊕ Low ^{a,c}	CFOs may result in a clinically important improvement in parent-rated QoL.

Treatment success (measured on the 50FTW (seconds)) follow-up: 3 months N° of participants: 28 (1 RCT)		The mean time for the 50FTW with shoes was 8.36 seconds	The mean time for the 50FTW with CFOs was 7.03 seconds	MD 1.33 seconds less (2.77 seconds less to 0.11 seconds more)	⊕⊕⊕⊕ Low ^{a,c}	CFOs likely result in little to no difference in timed walking.
Withdrawal due to adverse events follow-up: N° of participants: 28 (1 study)	RR 0.58 (0.11 to 2.94)	23.1%	13.4% (2.5% to 67.8%)	absolute difference 9.7% fewer (20.5% fewer to 44.8% more)	⊕⊕⊕⊕ Low ^{a,c}	CFOs likely result in little to no difference in withdrawals due to adverse events. Absolute reduction 9.7% (20.5% fewer to 44.8% more)
Adverse events	-	-	-	-	-	not reported
Serious adverse events	-	-	-	-	-	not reported

***The risk in the intervention group** (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: Confidence interval; **FFI:** Foot Function Index; **50FTW:** 50-Foot Timed Walk; **MD:** mean difference; **PedsQL:** Pediatric quality of life inventory; **RR:** Risk ratio; **VAS:** visual analogue scale; **QoL:** quality of life

GRADE Working Group grades of evidence

High certainty: we are very confident that the true effect lies close to that of the estimate of the effect

Moderate certainty: we are moderately confident in the effect estimate; the true effect is likely to be close to the estimate of the effect, but there is a possibility that it is substantially different

Low certainty: our confidence in the effect estimate is limited; the true effect may be substantially different from the estimate of the effect

Very low certainty: we have very little confidence in the effect estimate; the true effect is likely to be substantially different from the estimate of effect

^aDowngraded for bias; single blinded, children and their parents knew which treatment they had, which may have affected the assessment of pain

^bDowngraded for indirectness; only short-term outcomes (3 months); FFI not validated in children; PedsQL has no foot-related data

^cDowngraded for imprecision; small sample size and wide CI including both an increase and decrease in the effect estimate

Summary of findings 5. Prefabricated foot orthoses compared to shoes in children with juvenile idiopathic arthritis and flat feet

Prefabricated foot orthoses compared to shoes in children with juvenile idiopathic arthritis and flat feet

Patient or population: children with juvenile idiopathic arthritis and flat feet

Setting: outpatient rheumatology clinics

Intervention: prefabricated foot orthoses (PFO)

Comparison: shoes

Outcomes	Relative effect (95% CI)	Anticipated absolute effects* (95% CI)			Certainty of the evidence (GRADE)	What happens
		With shoes (N = 12)	With PFOs (N = 12)	Difference		
Pain (measured on 0 to 10-point VAS; lower = less pain) follow-up: 3 months Nº of participants: 25 (1 RCT)		The mean pain with shoes was 2.82 points	The mean pain with PFOs was 2.84 points	MD 0.02 points higher (1.94 points lower to 1.98 points higher)	⊕⊕⊕⊕ Very low ^{a,b,c}	PFOs likely result in little to no difference in pain.
Function or disability (measured on 0 to 100-point FFI; 0 = no disability) follow-up: 3 months Nº of participants: 25 (1 RCT)		The mean FFI score with shoes was 34.15 points	The mean FFI score with PFOs was 38.32 points	MD 4.17 points lower (24.4 points lower to 16.06 points higher)	⊕⊕⊕⊕ Low ^{a,c}	PFOs likely result in little to no difference in function or disability.
Quality of life (child-rated) (measured on 0 to 100-point PedsQL; higher score = better QoL) follow up: 3 months Nº of participants: 22 (1 RCT)		The mean child-rated PedQL score with shoes was 59.78 points	The mean child-rated PedQL score with PFOs was 37.99 points	MD 3.84 points on PedsQL lower (19.01 lower to 11.33 higher)	⊕⊕⊕⊕ LOW ^{1,3}	PFOs likely results in little to no difference in child-rated QoL.
Quality of life (parent-rated) (measured on 0 to 100-point PedsQL; higher score = better QoL) follow-up: 3 months Nº of participants: 22 (1 RCT)		The mean parent-rated PedQL score with shoes was 55.95 points	The mean parent-rated PedQL score with PFOs was 56.59 points	MD 0.64 points lower (13.22 points lower to 11.94 points higher)	⊕⊕⊕⊕ Low ^{a,c}	PFOs likely results in little to no difference in parent-rated QoL.
Treatment success (measured on the 50FWT (seconds)) follow-up: 3 months		The mean time for the 50FWT with shoes was 8.36 seconds	The mean time for the 50FWT with PFOs was 7.98 seconds	MD 0.38 seconds lower (1.9 seconds lower to 1.14 seconds higher)	⊕⊕⊕⊕ Low ^{a,c}	PFOs likely results in little to no difference in timed walking.



Nº of participants: 25 (1 RCT)						
Withdrawal due to adverse events follow-up:	RR 0.72 (0.14 to 3.61)	23.1%	16.6% (3.2% to 83.3%)	absolute difference - 6.5% less (19.8% less to 60.2% more)	-	PFOs likely results in little to no difference in withdrawals due to adverse events. Absolute reduction 6.5% (19.8% fewer to 60.2% more)
Nº of participants: 25 (1 study)						
Adverse events	-	-	-	-	-	not reported
Serious adverse events	-	-	-	-	-	not reported

*The risk in the intervention group (and its 95% confidence interval) is based on the assumed risk in the comparison group and the relative effect of the intervention (and its 95% CI).

CI: Confidence interval; FFI: Foot Function Index; 50FWT: 50-Foot Timed Walk; MD: mean difference; PedsQL: Pediatric quality of life inventory; RR: Risk ratio; VAS: visual analogue scale; QoL: quality of life

GRADE Working Group grades of evidence

High certainty: we are very confident that the true effect lies close to that of the estimate of the effect

Moderate certainty: we are moderately confident in the effect estimate; the true effect is likely to be close to the estimate of the effect, but there is a possibility that it is substantially different

Low certainty: our confidence in the effect estimate is limited; the true effect may be substantially different from the estimate of the effect

Very low certainty: we have very little confidence in the effect estimate; the true effect is likely to be substantially different from the estimate of effect

^aDowngraded for bias; single blinded; children and their parents knew which treatment they had, which may have affected their assessment of pain

^bDowngraded for indirectness; only short-term outcomes (3 months); FFI not validated in children; PedsQL had no foot-related data

^cDowngraded for imprecision; small sample size

Summary of findings 6. Custom foot orthoses compared to prefabricated foot orthoses in children with juvenile idiopathic arthritis and flat feet

Custom foot orthoses compared to prefabricated foot orthoses in children with juvenile idiopathic arthritis and flat feet

Patient or population: children with juvenile idiopathic arthritis and flat feet

Setting: outpatient rheumatology clinics

Intervention: custom foot orthoses (CFO)

Comparison: prefabricated foot orthoses (PFO)

Outcomes	Relative effect (95% CI)	Anticipated absolute effects* (95% CI)	Certainty of the evidence (GRADE)	What happens
----------	--------------------------	----------------------------------------	-----------------------------------	--------------

	With PFOs (N = 41)	With CFOs (N = 46)	Difference		
Pain (measured on 0 to 10-point VAS; lower = less pain) follow-up: 3 months to 6 months Nº of participants: 87 (2 RCTs)	The mean pain with PFOs was 3.22 points	The mean pain with CFOs was 1.74 points	MD 1.48 points lower (3.23 points lower to 0.26 points higher)	⊕⊕⊕⊕ Low ^{a,b}	CFOs may result in little to no difference in pain.
Function or disability (measured on 0 to 100-point FFI; 0 = no disability) follow-up: 3 months Nº of participants: 27 (1 RCT)	The mean FFI score with PFOs was 29.9 points	The mean FFI score with CFOs was 15.6 points	MD 14.38 points lower (30.22 points lower to 1.46 points higher)	⊕⊕⊕⊕ Low ^{a,b}	CFOs may result in little to no difference in function.
Quality of life (child-rated) (measured on 0 to 100-point PedsQL; higher score = better QoL) follow-up: 3 months to 6 months Nº of participants: 83 (2 RCTs)	The mean child-rated PedQL score with PFOs was 55.94 points	The mean child-rated PedQL score with CFOs was 64.58 points	MD 8.64 points higher (3.9 points lower to 21.18 points higher)	⊕⊕⊕⊕ Low ^{a,b}	CFOs may result in a small improvement in child-rated QoL.
Quality of life (parent-rated) (measured on 0 to 100-point PedsQL; higher score = better QoL) follow up: 3 months to 6 months Nº of participants: 84 (2 RCTs)	The mean parent-rated PedQL score with PFOs was 55.31 points	The mean parent-rated PedQL score with CFOs was 58.25 points	MD 2.94 points higher (11 points lower to 16.88 points higher)	⊕⊕⊕⊕ Low ^{a,b}	CFOs may result in little to no difference in parent-rated QoL.
Treatment success (measured on the 50FWT (seconds)) follow-up: 3 months Nº of participants: 27 (1 RCT)	The mean time for the 50FWT with PFOs was 7.98 seconds	The mean time for the 50FWT with CFOs was 7.03 seconds	MD 0.95 seconds lower (1.88 seconds lower to 0.02 seconds lower)	⊕⊕⊕⊕ Low ^{a,b}	CFOs may result in little to no difference in timed walking

Withdrawal due to adverse events	RR 0.80 (0.13 to 4.87)	4.9%	3.9% (0.6% to 23.8%)	1.0% fewer (4.2% fewer to 18.9% more)	⊕⊕⊕⊕ Low ^{a,b}	CFOs may result in little difference in withdrawals due to adverse events.
Follow-up: N ^o of participants: 87 (2 RCTs)						
Adverse effects	-	-	-	-	-	not reported
Serious adverse events	-	-	-	-	-	not reported

***The risk in the intervention group** (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: Confidence interval; **FFI:** Foot Function Index; **50FWT:** 50-Foot Timed Walk; **MD:** mean difference; **PedsQL:** Pediatric quality of life inventory; **RR:** Risk ratio; **VAS:** visual analogue scale; **QoL:** quality of life

GRADE Working Group grades of evidence

High certainty: we are very confident that the true effect lies close to that of the estimate of the effect

Moderate certainty: we are moderately confident in the effect estimate; the true effect is likely to be close to the estimate of the effect, but there is a possibility that it is substantially different

Low certainty: our confidence in the effect estimate is limited; the true effect may be substantially different from the estimate of the effect

Very low certainty: we have very little confidence in the effect estimate; the true effect is likely to be substantially different from the estimate of effect

^aDowngraded for bias; single blinded; children and their parents knew which treatment they had, which may have affected the assessment of pain

^bDowngraded for imprecision due to wide 95% CIs

Summary of findings 7. Prefabricated foot orthoses compared to shoes in children with symptomatic flat feet

Prefabricated foot orthoses compared to shoes in children with symptomatic flat feet

Patient or population: children with symptomatic flat feet

Setting: outpatient hospital clinic

Intervention: prefabricated foot orthoses (PFO)

Comparison: shoes

Outcomes	Relative effect (95% CI)	Anticipated absolute effects* (95% CI)			Certainty of the evidence (GRADE)	What happens
		With shoes (N = 26)	With PFOs (N = 26)	Difference		
Pain	-	-	-	-	-	not reported

Function or disability (global function assessed with 0 to 100-point PODCI; higher scores = better functioning) follow-up: mean 12 weeks Nº of participants: 52 (1 RCT)		The mean POD-CI score with shoes was 0.7 points	The mean POD-CI score with PFOs was 3.7 points	MD 3 points higher (2.28 points higher to 3.72 points higher)	⊕⊕⊕⊕ Low ^{a,b}	The evidence suggests that PFOs results in little to no difference in function
Quality of life (measured on 0 to 100-point PedsQL; higher score = better QoL) follow-up: mean 12 weeks Nº of participants: 52 (1 RCT)		The mean PedQL score with shoes was -1.1 points	The mean PedQL score with PFOs was 2.9 points	MD 1.8 points higher (1.07 points higher to 2.53 points higher)	⊕⊕⊕⊕ Low ^{a,b}	The evidence suggests that PFOs results in little to no difference in quality of life
Treatment success	-	-	-	-	-	not reported
Withdrawal due to adverse events	-	-	-	-	-	not reported
Adverse effects	-	-	-	-	-	not reported
Serious adverse events	-	-	-	-	-	not reported

***The risk in the intervention group** (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: confidence interval; **MD:** mean difference; **PODCI:** Pediatrics Outcomes Data Collection Instrument; **RR:** risk ratio

GRADE Working Group grades of evidence

High certainty: we are very confident that the true effect lies close to that of the estimate of the effect

Moderate certainty: we are moderately confident in the effect estimate; the true effect is likely to be close to the estimate of the effect, but there is a possibility that it is substantially different

Low certainty: our confidence in the effect estimate is limited; the true effect may be substantially different from the estimate of the effect

Very low certainty: we have very little confidence in the effect estimate; the true effect is likely to be substantially different from the estimate of effect

^aDowngraded for bias (participants and parents aware of treatment received; selective reporting of outcomes, as the published study included more outcomes than were listed in the trial registry (ClinicalTrials.gov [NCT02414087](https://clinicaltrials.gov/ct2/show/study/NCT02414087)))

^bDowngraded for imprecision due to small sample size, small effects across scaled outcome measures

BACKGROUND

Description of the condition

Despite decades of attention and scrutiny (Aharonson 1992; Brooks 1991; Mereday 1972; Staheli 1987), the paediatric flat foot remains a quandary for clinicians, researchers, and parents alike. It is established that some flat feet are associated with pain (Rome 2010), but not all flat feet are painful or debilitating. Debate about pre-emptive treatment for flat feet in children has been misguided (Bresnahan 2009; D'Amico 2009; Evans 2008; Harris 2010). Hence, it is important to clarify whether any form of treatment is indicated, for children with flat feet, which are not painful. Prevalence estimates for paediatric flatfoot vary broadly. It has been reported as 44% in children aged three to six years, and 24% in children aged six years or older (Pfeiffer 2006); 70% in children aged three to four years, and 40% by five to eight years (Daneshmandi 2011); 23.5% in seven to 14 year olds (Yoosefinejad 2014); and between 2.2% and 12.3% in children aged four to 13 years (Garcia-Rodriguez 1999).

In a study of 835 school children in Austria, Pfeiffer 2006 reported that 10% of children with flat feet were wearing foot orthoses, whilst only 1% were deemed pathological, indicating a marked over use of foot orthoses. Yan 2013 reported flatfoot in 90% of 100 normal Chinese children in Beijing, aged less than two years, and just 4% at age 10 years. Whilst different methods of assessment were used, this trend crosses both ethnicity and age groups, and is now further reinforced by both normative and prospective findings (Gijon-Nogueron 2019; Martinez-Nova 2018).

What has often failed to be appreciated, is the developing morphology of the paediatric foot structure, i.e. from flat to less flat across the first decade of life, with some variation (Bresnahan 2009; Evans 2008; Wenger 1989). The definition for flatfoot, whilst not universal, does find agreement across authors on the position of the heel (everted – valgoid), and the medial longitudinal foot arch (flat – convex (Capello 1998; Evans 2008; Staheli 1987; Wenger 1989)). What is universal, and reasonable, is concern about pain and functional limitation that may occur with some children who have flat feet, to potentially diminish mobility, independence, and quality of life.

Markers of benign versus pathological paediatric flat feet have been identified, and assist with predicting the later symptomatic cases in older children (Evans 2021). The three markers are: (1) valgus heel, seen clinically as a greater than 10 degree resting calcaneal stance position (RCSP (Kerr 2015)); (2) talo-navicular joint coverage angle, seen clinically (on x-ray) as a greater than 35 degree medial talar head exposure (Moraleda 2011; Yan 2013); and (3) ankle range reduction or 'equinus', seen clinically as a less than 30 degree weight-bearing lunge (Kim 2017).

Recent normative data, based on over 3000 healthy children, has shown that paediatric foot posture has a wide normal range across childhood, with the average Foot Posture Index (FPI) equal to +4(3), such that FPI scores within the range of +1 to +7 (1 standard deviation) include 68% of children (Gijon-Nogueron 2019). Further, prospective data from over 1000 healthy children followed for three years, showed that at each age point, foot posture 'centralised'. This means that there were fewer pronated (flat) and highly pronated (flatter) foot types as age increased, with slight increases in supinated foot types. The greatest prospective shift was the

increase of normal foot types as age increased (Martinez-Nova 2018).

Flat feet are also commonly seen in children with diagnoses associated with indisputable risk of foot pathology, e.g. juvenile idiopathic arthritis (JIA), with a prevalence of approximately 1:5000 births; cerebral palsy, with a prevalence of approximately 1:400 births; and trisomy 21/Down syndrome, with a prevalence of approximately 1:900 births. However, the most significant paediatric foot pathologies are not actually flat feet, but feet with a high arch (cavus morphology), as affect children with congenital talipes equino varus (CTEV – prevalence approximates 1:1000 births), heritable motor neuropathies such as Charcot Marie Tooth (CMT – prevalence approximates 1:3300 births), infections such as Poliomyelitis.

Description of the intervention

Treatment options for paediatric flatfoot vary from non-surgical to surgical approaches (Klaue 1997). The latter are rare, and are usually pursued only after failure of non-invasive management, or for rigid flatfoot presentations (Bauer 2015). Non-surgical interventions include advice, FOs (foot orthoses), stretching or strengthening exercises, footwear type and modifications, and less commonly, neuromuscular electrical stimulation (NMES), serial casting, weight reduction, analgesic and anti-inflammatory medications (Blitz 2010).

Whilst FOs, per se, include a range of physical appliances, there are important distinctions between custom or bespoke FOs (CFOs), prefabricated FOs (PFOs), and customised PFOs (CPFOs).

How the intervention might work

The basic premise for foot orthoses as a treatment for paediatric flatfoot, is to promote a stable foot posture that allows an efficient gait function. By distributing forces and loads across the weight-bearing, body-carrying foot, joint ranges can be used effectively for gait function, without harmful stresses and strains. Paired with footwear, which is known to influence both foot stability and gait discretely, FOs are the treatment mainstay for paediatric flatfoot (Wegener 2011).

Other treatment options, such as stretching (e.g. calf musculature for ankle dorsiflexion range), specific muscle strengthening, and core muscle strengthening, have often been regarded as 'adjunctive' to FOs. Footwear acts as supplementary 'whole child' treatment directives in children with muscular hypotonia who also require strength and balance physical therapy (Rigoldi 2012; Weber 2014).

Footwear is frequently overlooked, and yet influences FOs as the basic 'housing' structure. All footwear, but especially athletic footwear, that is constructed as a structure external to the foot in gait, will influence foot functioning. Cursorily, athletic footwear is intentionally designed to stabilise the flat foot, co-ordinate with the rectus foot, and cushion the high arch foot (Evans 2015). Hence the use of any FOs must only be subsequent to evaluation of footwear effects. This applies not only to the paediatric flat foot, but to all clinical gait evaluation, and always prior to any consideration of FOs use.

Why it is important to do this review

Most cases of paediatric flat feet fall within the range of normal findings (Gijon-Nogueron 2019), yet there is a lack of confidence in both primary and specialist care (Carli 2012). Further, children's flat feet improve as they grow during their first 10 years (Gijon-Nogueron 2019; Martinez-Nova 2018).

There is clear need for accurate guidance from robust scientific evidence. In this era of over-diagnosis of disease (Moynihan 2012; Moynihan 2014; Moynihan 2017), overmedicalisation of regular range phenomena, and unnecessary treatment of normal variation (Evans 2017; Evans 2021). The cost to public health systems of unnecessary consultations, and unnecessary treatment is not insubstantial. Screening children for flat feet is unfounded, and both logically and economically refuted (Evans 2012; Wilson 1968). Most paediatric flatfoot presentations reveal flexible feet that are pain-free. However, a flat foot that is either painful or rigid is not a normal finding, and requires both diagnosis and effective treatment. Specific subgroups of children have conditions known to predispose them to foot pain, such as JIA, increased joint laxity (e.g. Ehlers-Danlos, Down, or Marfan syndromes), and tarsal coalitions. These may present overtly or covertly, and should be part of a differential diagnosis for painful paediatric flatfoot (Evans 2009; Evans 2021).

This review is an update of an earlier Cochrane Review, which found limited evidence from three trials on the use of non-surgical interventions for the treatment of paediatric flatfoot (New Reference). Considering that a burden of paediatric flat feet is not universally demonstrable, and early identification has not been found to be beneficial, this review aims to provide answers as to whether a continued concern regarding flat feet in healthy children is necessary (Evans 2012).

OBJECTIVES

To assess the efficacy (benefits and harms) of foot orthoses as treatment for paediatric flat feet versus no treatment or other treatment.

METHODS

Criteria for considering studies for this review

Types of studies

All randomised controlled trials (RCTs) and pseudo-randomised controlled clinical trials (CCTs; using methods of allocating participants to a treatment that are not strictly random, for example date of birth, hospital record number, or alternation).

Types of participants

Since there is no universally accepted definition for paediatric flatfoot, flat feet or flatfoot are the terms used to describe a recognisable clinical foot morphology that involves several adjacent joints of the foot (Harris 2004). We included trials involving children younger than 16 years of age with a diagnosis of flat feet. Studies of various soft tissue diseases, and pain due to tendinitis, were eligible for inclusion, provided that the flat foot pain results were presented separately. Studies in which participants had heel pain, stress fractures of the metatarsals, ankle fractures, rheumatoid foot pathology, diabetic foot, or

neuromuscular conditions were also eligible for inclusion, provided all children, or an identified subgroup of children, had flat feet.

We included trials that included children with asymptomatic flat feet, juvenile idiopathic arthritis (JIA; where flatfoot is a common clinical feature (Henry 2008)), or other clinical concerns.

Types of interventions

Interventions included rigid, semi-rigid, or soft foot orthoses (FO), corrective footwear; strengthening exercises, stretching exercises, activity modification; manipulation; serial casting; weight reduction; anti-inflammatory medication; anti-pronatory strapping; neuromuscular electrical stimulation (NMES); and educational advice to children or their parents or guardians, and compared FOs versus sham or no intervention (control), or other non-surgical interventions for paediatric flat feet.

Comparison were made with other interventions, and with no treatment (with deference to usual footwear in some trials). Epidemiological data regarding normal foot posture across childhood has provided a context for better clinical appreciation of normal variation, thus discouraging narrow intervention criteria (Gijon-Nogueron 2019).

We excluded studies involving surgical interventions.

Types of outcome measures

Major outcomes

The following major outcomes will frame this review and are reported in the summary of findings tables: [Summary of findings 2](#); [Summary of findings 3](#); [Summary of findings 4](#); [Summary of findings 5](#); [Summary of findings 6](#).

1. Pain was considered as proportion and group means, with most interest on change from baseline to intervention, and comparative intervention effects. Pain was stipulated as being reported by the child, parent, or carer. This outcome was only applicable in studies involving symptomatic participants.
2. Function or disability indices of the foot
3. Quality of life measures
4. Treatment success (e.g. measured by a participant or proxy-reported global impression of clinical change, as being very much improved, improved, or similar). The parameters of treatment success could include score changes, functional change, with changes measured as means, proportions, and variance.
5. Proportion of withdrawals for each trial group, both intervention and control
6. Proportion with adverse events, as reported
7. Proportion with serious adverse events, as reported

Minor outcomes

Goniometric measurements, x-rays, or those that were collated in a gait laboratory (both kinetic and kinematic data).

Search methods for identification of studies

Electronic searches

We searched:

- Cochrane Central Register of Controlled Trials (CENTRAL; 2021, Issue 9) in the Cochrane Library (searched 1 September 2021; see [Appendix 1](#));
- MEDLINE Ovid (July 2009 to 1 September 2021; see [Appendix 2](#))
- Embase via Ovid (July 2009 to 1 September 2021; see [Appendix 3](#))
- World Health Organization International Clinical Trials Registry Platform (WHO ICTRP; apps.who.int/trialsearch; searched 7 August 2020);
- US National 7 August of Health Ongoing Trials Register ClinicalTrials.gov (www.clinicaltrials.gov; searched 7 August 2020).

We started the search in 2009 to allow overlap with the search from [Evans 2010](#).

In MEDLINE, we combined subject specific search terms and free text words with the optimum search strategy for randomised trials described by [Lefebvre 2008](#). We adapted the search strategy for the other databases

Searching other resources

We complemented the electronic search by checking reference lists of relevant articles for additional studies reported in published papers, scientific meetings, and personal communication.

Data collection and analysis

Selection of studies

One of the review authors (KR) assisted by the Cochrane Musculoskeletal Group's Information Specialist at the editorial base, carried out the initial searches. Pairs of review authors (FH and AE; KR and FH; KR and AE) independently assessed potentially eligible trials for inclusion; they resolved any disagreement through discussion. Titles of journals and names of authors or supporting institutions were not masked at any stage.

We conducted this systematic review in accordance with the PRISMA statement guidelines ([Moher 2015](#)).

Data extraction and management

Three authors (AE, KR, FH) independently extracted data using a pre-piloted form. They resolved any disagreement through discussion, using electronic communications techniques.

We extracted the following study characteristics:

1. Methods: study design, total duration of study, details of any 'run-in' period, number of study centres and location, study setting, withdrawals, and date of study
2. Participants: N, mean age, age range, sex, disease duration (JIA), severity of condition, diagnostic criteria, important baseline data, inclusion criteria, and exclusion criteria
3. Interventions: intervention, comparison, concomitant medications, and excluded medications
4. Outcomes: primary and secondary outcomes specified and collected, and time points reported
5. Characteristics of the design of the trial as outlined below in the [Assessment of risk of bias in included studies](#) section.
6. Notes: funding sources for trials, and notable declarations of interest from authors.

When a trial included more than one measure for an outcome, we adopted a pre-specified hierarchy, as follows: pain measures, gait and function, health-related quality of life (HRQoL; 1. child, 2. parent proxy), with precedence given to validated outcome measures where multiple measures were available in a trial - as reflects clinical and parent concerns and clinical care priorities.

Our decision for selecting data to extract, included:

- if both final values and change from baseline values were reported for the same outcome, we extracted final values;
- if both unadjusted and adjusted values for the same outcome were reported, we extracted unadjusted values
- if data were analysed on the basis of intention-to-treat (ITT), we extracted the sample treated with custom foot orthoses (CFO; this did not differ for outcomes assessing benefits or harms);
- If multiple time points were reported, we extracted final time points

Main comparisons

Note that in this review there are multiple populations and comparisons, and we have stated these in order of clinical importance:

1. Custom foot orthoses (CFO) versus shoes (CFOs are the most expensive FOs, and shoes are known to alter gait and foot mobility, as reported in the systematic review from [Wegener 2011](#))
2. Prefabricated foot orthoses (PFO) versus shoes (children with JIA have indisputable disease, and frequently present with debilitating foot pain ([Fellas 2017a](#)))
3. CFO versus PFO (both are common interventions and comprise usual care for common concerns about flatfoot in children ([Pfeiffer 2006](#)). However, the cost ratio approximates 4:1 (CFO:PFO), and the justification for CFO use has been both questioned ([Evans 2008](#)), and defended ([Bresnahan 2009](#); [D'Amico 2009](#)).

Assessment of risk of bias in included studies

Two review authors (FH, AE) independently assessed the risk of bias of each included trial against key criteria: random sequence generation; allocation concealment; blinding of participants, personnel; blinding of outcome assessment - separately for subjective self-reported outcomes, such as pain and function, and objective outcomes, such as adverse events; incomplete outcome data; selective outcome reporting; and other sources of bias (e.g. follow-up times, participant maturation). This is in accordance with methods for RoB 1 recommended by The Cochrane Collaboration ([Higgins 2017](#)). Review authors resolved disagreements by consensus.

We classified each potential source of bias as high, low, or unclear risk, and provided a quote from the study report, together with a justification for our judgement in the risk of bias table. When information on risk of bias related to unpublished data or correspondence with a trialist, we noted this in the risk of bias table. We presented the figures generated by RoB 1 to provide summary assessments of the risk of bias.

Measures of treatment effect

We analysed dichotomous data as risk ratios (RR) with 95% confidence intervals (CI). We calculated mean differences (MD) and 95% CI for continuous outcomes measured on the same scale, and standardised mean difference (SMD), if different scales were used to measure an outcome, and 95% CIs. We entered data presented as a scale with a consistent direction of effect across studies. SMDs were back-translated to a typical scale (e.g. 0 to 10 for pain) by multiplying the SMD by a typical among-person standard deviation (e.g. the standard deviation of the control group at baseline from the most representative trial; as per Chapter 6 of the *Cochrane Handbook* (Higgins 2020b)).

In the 'Comments' column of the summary of findings table, we reported the absolute percent difference, the number needed to treat number for an additional beneficial outcome (NNTB), and the number needed to treat for an additional harmful outcome (NNTH). We calculated the absolute percent change from the difference in the risks between the intervention and control group using GRADEpro GDT, and expressed as a percentage (GRADEpro GDT). The NNTB or NNTH is only provided for dichotomous outcomes that show a clinically significant difference). We calculated the NNTB or NNTH from the control group event rate and the relative risk, using the Visual Rx NNT calculator (Cates 2016). The minimal clinically important differences (MCID) were 0.9 points for pain, measured on a 10-point visual analogue scale (VAS); and 7 points for disability, measured on the 100-point disability subscale of the Foot Function Index (FFI), as calculated by Landorf 2008.

Unit of analysis issues

When multiple trial arms were reported in a single trial, we included only the relevant arms. If two comparisons (e.g. CFOs versus shoes, and CFOs versus sham orthoses) were combined in the same meta-analysis, we halved the control group to avoid double-counting. We clarified the presence of more than two intervention groups in the Characteristics of included studies table.

Dealing with missing data

We contacted investigators to verify key study characteristics and request missing numerical outcome data, when indicated (e.g. when a study was identified as abstract only, or when data were not available for all participants). Any assumptions and imputations used to handle missing data were reported explicitly in the Characteristics of included studies table, and the effect of assumptions or imputations was explored with sensitivity analyses.

For dichotomous outcomes (e.g. number of withdrawals due to adverse events), we calculated the withdrawal rate using the number of participants randomised to the group as the denominator.

For continuous outcomes (e.g. mean pain), we calculated the MD or SMD based on the number of participants analysed at that time point. If the number of participants analysed was not presented for each time point, we used the number of participants randomised to each group at baseline.

When possible, we computed missing standard deviations from other statistics, such as standard errors, confidence intervals, or P values, according to the methods recommended in the *Cochrane Handbook for Systematic Reviews of Interventions* (Deeks 2020). If

we were unable to calculate standard deviations from the available data, we imputed them (e.g. from other studies in the meta-analysis).

Assessment of heterogeneity

We assessed clinical and methodological diversity of participants, interventions, outcomes, and study characteristics for the included studies, to determine whether a meta-analysis was appropriate. We assessed statistical heterogeneity by visually inspecting forest plots for obvious differences in result between the studies, and using the I^2 statistic. As recommended in the *Cochrane Handbook for Systematic Reviews of Interventions*, the interpretation of an I^2 value of 0% to 40% might not be important; 30% to 60% may represent moderate heterogeneity; 50% to 90% may represent substantial heterogeneity; and 75% to 100% may represent considerable heterogeneity (Deeks 2020). If we identified substantial heterogeneity, we reported it. We planned to investigate possible causes with subgroup analyses, had data permitted.

Assessment of reporting biases

Had we been able to pool more than 10 trials, we planned to undertake formal statistical tests to investigate funnel plot asymmetry, and follow the recommendations in Section 13 of the *Cochrane Handbook for Systematic Reviews of Interventions* (Page 2020).

To assess outcome reporting bias, we checked trial protocols against published reports. For studies published after 1 July 2005, we screened the International Clinical Trials Registry Platform of the World Health Organisation for the trial protocol (apps.who.int/trialsearch). We evaluated whether selective reporting of outcomes was present.

Data synthesis

We pooled data using a random-effects model across studies for outcomes with common interventions and comparators, for participants with similar flatfoot conditions (i.e. JIA, pain, asymptomatic).

Subgroup analysis and investigation of heterogeneity

We did not plan any subgroup analyses.

Sensitivity analysis

Had there been sufficient data, we planned to examine the potential effect on results for pain and function of selection bias by restricting the analysis to studies at low risk of selection bias (adequate allocation concealment); detection bias by restricting the analysis to studies with low risk of detection bias (adequate blinding of outcome assessor - the participants for these outcomes) and those with imputed missing data.

Interpreting results and reaching conclusions

We followed the guidelines provided in Chapter 15 of the *Cochrane Handbook for Systematic Reviews of Interventions* when interpreting results, and we were aware of distinguishing lack of evidence of effect from lack of effect (Schunemann 2020b). We based our conclusions only on findings from the quantitative or narrative synthesis of included studies for this review. We avoided making recommendations for practice, and our implications for research

suggested priorities for future research and outlined remaining uncertainties in this area.

Summary of findings and assessment of the certainty of the evidence

GRADE and Summary of findings tables

We collated seven summary of findings (SoF) tables for the following comparisons:

- custom foot orthoses (CFO) compared to shoes in asymptomatic flat feet ([Summary of findings 1](#));
- prefabricated foot orthoses (PFO) compared to shoes in asymptomatic flat feet ([Summary of findings 2](#));
- CFOs compared to PFOs in asymptomatic flat feet ([Summary of findings 3](#));
- CFOs compared to shoes in children with juvenile idiopathic arthritis (JIA; [Summary of findings 4](#));
- PFOs compared to shoes in children with JIA ([Summary of findings 5](#));
- CFOs compared to PFOs in children with JIA ([Summary of findings 6](#));
- PFOs compared to shoes in symptomatic flat feet ([Summary of findings 7](#)).

Summary of finding tables provide key information concerning the quality of evidence, the magnitude of effect of the interventions examined, and the sum of available data on the major outcomes ([Types of outcome measures](#)), as recommended Chapter 14 of the *Cochrane Handbook for Systematic Reviews of Interventions* ([Schünemann 2020a](#)).

We used the five GRADE considerations (study limitations, consistency of effect, imprecision, indirectness, and publication bias) to assess the quality of a body of evidence comprised of

the studies that contributed data to the meta-analyses for the prespecified major outcomes. We reported the quality of evidence as high, moderate, low, or very low. We used [GRADEpro GDT](#) software to prepare and present the SoF tables ([GRADEpro GDT](#)). We justified all decisions to downgrade the quality of the evidence using footnotes. We provided the NNTB or NNTH and absolute percent change in the 'Comments' column of the SoF table, as described in the 'Measures of treatment effect' section above.

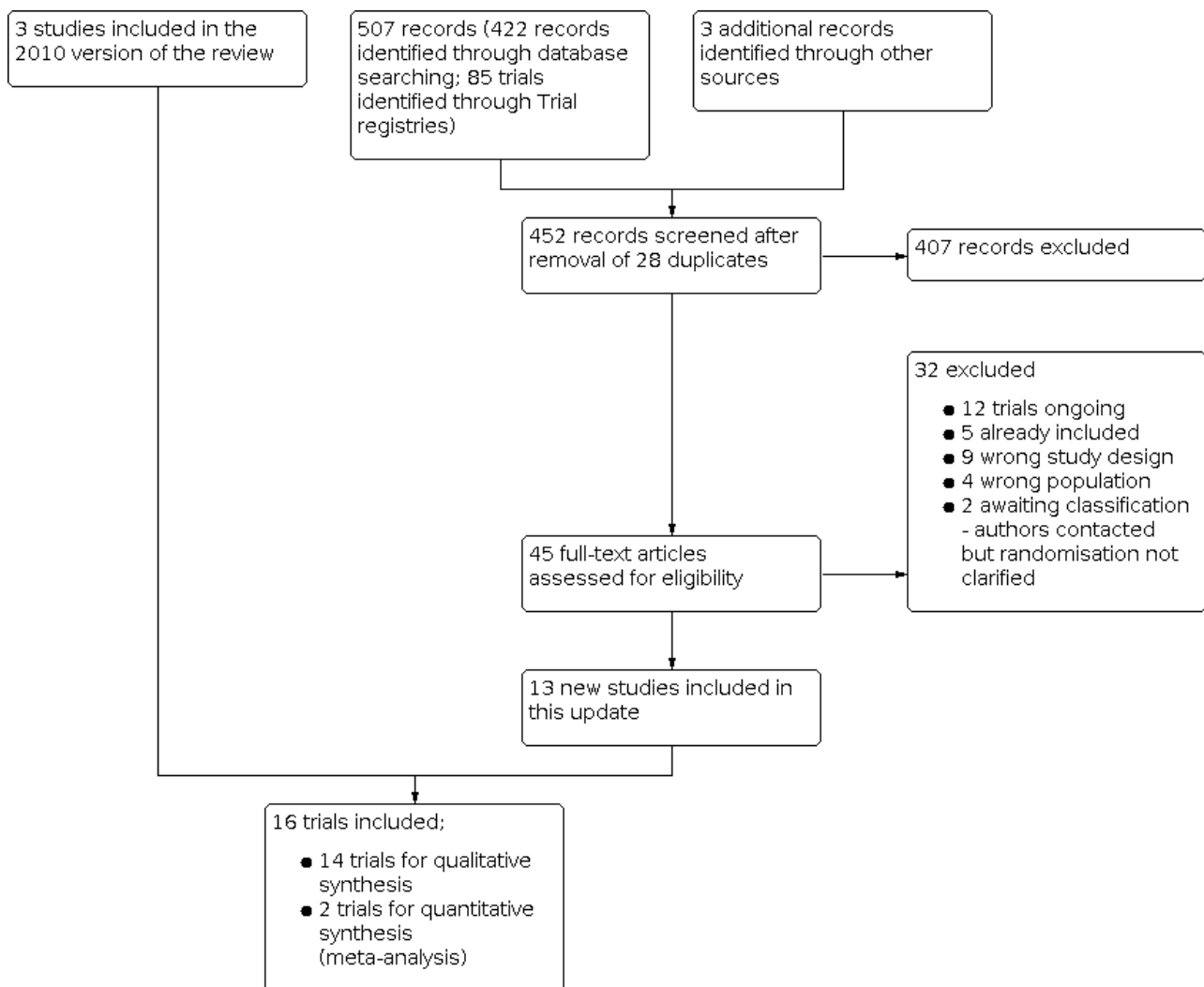
RESULTS

Description of studies

Results of the search

Our updated search retrieved 422 records (66 from CENTRAL, 252 from MEDLINE, and 104 from Embase). Our search of WHO ICTRP retrieved 38 citations; ClinicalTrials.gov retrieved 47 citations, giving a total of 507 records. We removed 28 duplicates, and screened 452 records. We excluded 407 records, and retrieved 45 for full-text review for possible inclusion. Of the 45 full-text articles, we excluded a further 13 ([Figure 1](#)) ([Benedetti 2011](#); [Ford 2017](#); [Hill 2020](#); [Hurd 2010](#); [MacKenzie 2012](#); [Mosca 2010](#); [Okamura 2020](#); [Perhamre 2011](#); [Perhamre 2012](#); [Pothrat 2013](#); [Riccio 2009](#); [Uden 2017](#); [Yung 2011](#)) because of wrong study design and wrong population; 5 were trials already included ([Ahn 2017](#); [Kanatli 2016](#); [Wenger 1989](#); [Whitford 2007](#); [Pandey 2013](#)). Additional details of these excluded studies are reported in the [Characteristics of excluded studies](#) Table. Twelve ongoing studies were identified, and hence included in the table of [Characteristics of ongoing studies](#). We contacted investigators of two RCTs in order to verify key study characteristics and obtain missing numerical outcome data ([Pandey 2013](#); [Sinha 2013](#)). Both authors were contacted via email, and unfortunately this met with no success in gaining the required data (answers to queries, missing data, clarification of randomisation, and SDs). These studies were included in the table of [Characteristics of studies awaiting classification](#).

Figure 1. Study flow diagram for the trial search



Thus, 13 studies met the final eligibility criteria for inclusion in this review update (Abd-Elmonem 2021; Aboutorabi 2013; Ahn 2017; Asgaonkar 2012; Bok 2016; Coda 2014; Gould 1989; Hsieh 2018; Jafarnezhadgero 2018; Kanatli 2016; Khamooshi 2017; Morrison 2013; Solanki 2020). However, two studies established only immediate effects of interventions, with no follow up data, and hence were regarded for description, but not for analysis (Aboutorabi 2013; Bok 2016). Gould 1989 was not included in the 2010 review, due to absence of a discrete control group, which was a criterion at that time (Rome 2010). Three studies were included from the previous version of the review (Powell 2005; Wenger 1989; Whitford 2007). Hence, a total of 16 studies were included in this review update.

Included studies

We included a total of 16 randomised controlled trials (RCT), details of which are included in the Characteristics of included studies tables and Table 1; Table 2; and Table 3.

Design

Ten of the 16 randomised RCTs included a no treatment control group (Abd-Elmonem 2021; Asgaonkar 2012; Hsieh 2018;

Jafarnezhadgero 2018; Kanatli 2016; Khamooshi 2017; Morrison 2013; Solanki 2020; Wenger 1989; Whitford 2007). The studies were conducted in nine countries (Australia, USA, UK, Iran, Egypt, Turkey, Republic of Korea, India, Taiwan) and were English language publications. The duration of trials ranged from four weeks (Solanki 2020), to five years (Gould 1989). The trials were parallel designs; seven had multiple arms (Aboutorabi 2013; Bok 2016; Gould 1989; Khamooshi 2017; Powell 2005; Wenger 1989; Whitford 2007), seven had a single intervention versus control group (Abd-Elmonem 2021; Asgaonkar 2012; Hsieh 2018; Jafarnezhadgero 2018; Kanatli 2016; Morrison 2013; Solanki 2020), and two trials comparing two single interventions (Ahn 2017; Coda 2014).

Most studies only included data measured at baseline and final time points; four reported multiple time points (Gould 1989; Solanki 2020; Wenger 1989; Whitford 2007). Two studies investigated immediate effects only and provided no follow-up data; we included them in the review, but not in the analyses (Aboutorabi 2013; Bok 2016).

Participants

The 16 studies included a total of 1058 children who completed the trials, and were aged from 11 months to 19 years, were included in the review. The inclusion and exclusion criteria used in all studies were clearly, if variably, defined. The majority of trials were in healthy children with flexible flat feet, with a range of inclusion criteria, and age groups. The inclusion criteria for trials of children with juvenile idiopathic arthritis (JIA) were: lower limb involvement, failed treatment with foot orthoses, ability to walk 15 metres, disease modifying anti-rheumatic drugs (DMARDs) for six months or more (Coda 2014); foot pain for > 1 month but < 2 years (Powell 2005). One trial with 22 boys included those with a diagnosis of developmental co-ordination disorder (DCD), and a Foot Posture Index (FPI) > +4 (Morrison 2013). Four trials recruited children with symptomatic flat feet (Bok 2016; Coda 2014; Hsieh 2018; Powell 2005).

Two groups, defined only by diagnosis, included:

1. Asymptomatic flat feet in healthy children

Ten trials (N = 805) assessed the effect of a number of comparisons on healthy children with asymptomatic, flexible flat feet. They collected data at baseline, and a number of defined time points. Whilst exhibiting obvious diversity, these trials represented the most children, a range of intervention modalities, common clinical practice presentations, and thus, the bulk of potentially relevant evidence (Abd-Elmonem 2021; Ahn 2017; Asgaonkar 2012; Gould 1989; Jafarnezhadgero 2018; Kanatli 2016; Khamooshi 2017; Solanki 2020; Wenger 1989; Whitford 2007).

Wenger 1989 evaluated 131 children, aged 1 to 6 years, with clinically diagnosed pes planus and no pain. Bilateral pes planus was diagnosed by visual observation of the valgus position of the heel, and the low appearance of the arch upon weight bearing.

Gould 1989 assessed 125 normal toddlers (beginner walkers) aged 11-14 months. Gender of children was not mentioned until the 5 year-old results, in 25/50 Group 1 participants (15M, 10F).

Whitford 2007 evaluated 178 children, aged 7 to 11 years, with clinically diagnosed pes planus and no pain. Bilateral pes planus was diagnosed by the assessment of calcaneal eversion in RCSP, and by the navicular drop test. The navicular drop test measures the extent of excessive foot pronation in static stance.

Asgaonkar 2012 recruited 80 children with flat feet, aged between five and 15 years. It is not especially clear, but appears that 139/894 of the children initially screened had flat feet, from which a random 80/139 were enrolled in the study. Uneven groups of 45 treatment and 35 controls were formed at random. Flat foot assessment was based on inked footprints, and the ratio of midfoot: foot width.

Jafarnezhadgero 2018 recruited 30 boys, aged 8 to 12 years. The boys were randomised into two same-size groups (N=15). Flat foot was assessed using navicular drop, arch height index, and resting calcaneal stance position.

Kanatli 2016 evaluated 45 children (33 boys, 12 girls) with mean age of 39.5 months (17-72 months), with diagnosis of moderate flexible flat foot.

Khamooshi 2017 assessed 60 female aged 9 to 13 year old, with good general health and flat feet.

Ahn 2017 investigated 40 children who were aged 10 years (4.5), with flexible flat feet.

Solanki 2020 investigated 44 children (sex unspecified) with mean age indicated as "students from 8th to 9th standard" (estimated age of 12 to 14 years), with clinically assessed 'hyperpronated' (flat) feet, as diagnosed by 'too many toes sign', navicular height, calcaneal angle.

Abd-Elmonem 2021 evaluated 72 typically developing children aged from 7 to 12 years (31 boys, 35 girls at completion) with asymptomatic flexible flat feet, as a clinically diagnosed.

2. Children with JIA, and other clinical concerns

Two studies (N = 108) included children diagnosed with JIA, flat feet, and foot pain (Coda 2014; Powell 2005). Powell 2005 evaluated 48 children, aged 5 to 19 years, diagnosed with JIA with pes planus and foot pain. Coda 2014 recruited 60 children with JIA, aged 10.64 (3.84) years, controls: 11.17 (3.51) years (controls: 6M: 23F; treatment group 9M: 22F).

Painful flatfoot: Hsieh 2018 evaluated 52 six-year-old children (28 males and 24 females) with symptomatic flexible flat feet.

Morrison 2013 investigated 22 boys with a diagnosis of DCD, who had flexible flat feet, a FPI > +4, and were aged 6 to 11 years.

Immediate effects only, were reported for two studies (Bok 2016; Aboutorabi 2013). Bok 2016 examined 21 children with flexible flat feet and foot pain, aged 8 to 13 years. Aboutorabi 2013 assessed 30 children with asymptomatic flexible flat feet, and a matched control group (N=20), with mean age 7.76 years. The two immediate effects trials, with data derived only at baseline time points were not included in analyses.

Overall, four RCTs were deemed too anomalous for inclusion in the main sub-groups (Hsieh 2018; Morrison 2013; Bok 2016; Aboutorabi 2013)

Interventions

The 16 included studies included a range of interventions: custom foot orthoses (CFO), prefabricated foot orthoses (PFO), CPFOs (customised PFOs), valgus insoles, inverted FOs, heel cups, University of California Biomechanics Laboratory (UCBL) supports, orthopaedic shoes (straight last, straight last plus cookie, with medial arch strengthening), foot exercises, foot plus core exercises, anti-pronation taping, neuromuscular electrical stimulation (NMES). Definitions of foot orthoses types, i.e. CFOs, PFOs, and CPFOs can be found in Table 1.

Reporting of intervention according to groups

1. Asymptomatic flat feet in healthy children

Wenger 1989 randomised each child into one of four groups: (1) orthopaedic shoes with no corrective features, the shoes were normal in contour and contained a steel shank; (2) shoes with a Thomas heel, a long medial counter and a navicular pad; (3) shoes with a Thomas heel, a long medial counter and a heel cup; (4) shoes and customised UCBL plastic foot orthoses. The trial ran for 3 years.

Gould 1989 randomised the children across four treatment groups: (Gp 1) straight last shoes (Gp 2) Gp 1 shoes plus long arch cookies, (Gp 3) orthopaedic shoes with long counters, solid shanks, Thomas heels, 0.3cm inside heel wedges, (Gp 4) Gp 3 shoes plus thin longitudinal arch support. The trial ran for 4 years.

Whitford 2007 randomised each child into one of three groups: (1) CFOs made from a rigid thermoplastic material with a vinyl cover; (2) PFOs made from a semi-rigid thermoplastic material with a standard intrinsic heel postings of 4° and a 5mm metatarsal rise; (3) no treatment. All children with shortened ankle joint plantar flexors were taught how to conduct calf muscle stretches at home and the researchers discussed suitable shoes with the parents. The trial ran for 12 months.

In **Asgaonkar 2012** the intervention group wore valgus PFO or insoles for 1 year, and the control group received no treatment.

Jafarnezhadgero 2018 randomised each child to either CFO treatment or sham insoles for a 4 month trial. All participants wore the same shoes from trial commencement.

Kanatli 2016 compared custom made orthopaedic shoes (N=21), with a control group using the outcomes of Joint laxity, AI, x-ray angles over 34.6 +/- 10.9 months (2-5 years). Shoes were renewed every 6 months in the treatment group, when x-rays were also taken.

Ahn 2017 compared two CFO designs (N=40), Over 1 year, in children of mean age 10.14 (4.99) years.

Khamooshi 2017 compared foot exercises versus foot and core exercises with an untreated control group, in an 8 week trial for girls (N=60) aged 9 to 13 years.

Solanki 2020 randomised each child (N=44) to either rigid taping and conventional therapy (strengthening exercises, Faradic foot bath), versus sham taping and same conventional therapy, in a 4 week trial for children aged approximately 12 to 14 years, of unspecified sex. **Solanki 2020**

Abd-Elmonem 2021 compared corrective (foot strength) exercises and NMES, versus corrective (foot strength) exercises and sham NMES (0 current), in a 4 month trial, for 72 children, aged 7 to 12 years (31 boys, 35 girls at completion).

2. Children diagnosed with JIA, flat feet and foot pain, or other clinical concerns

Powell 2005 randomised each child into one of three intervention groups: (1) custom-made semi-rigid orthotics made of metal-particle reinforced plastic with shock-absorbing posts; (2) prefabricated shoe inserts made from flat neoprene; and (3) new supportive athletic shoes with a medial longitudinal arch and shock-absorbing insoles. All children received new athletic shoes at the beginning of the study. This trial ran for 3 months.

Coda 2014 randomised each child into two groups, with CPFOS versus control PFOs over a 6 month period.

Painful flexible flat foot: one study evaluated young children with painful flexible flat foot, N=52 (**Hsieh 2018**) aged 6 years, and compared customised PFOs (N=26), with a control group using the

outcomes of physical activity, function and quality of life over 12 weeks.

Developmental coordination disorder (DCD): one study assessed boys diagnosed with DCD (**Morrison 2013**). All participants completed a 7 week rehabilitation programme, with a treatment group using CFOs from the start (N=9, mean age 8 years) and a control group who received CFOs at completion of the 7 week trial (N=5, mean age 6.5 years).

Two studies assessed only immediate effects of shoes and FOs (**Aboutorabi 2013**, **Bok 2016**), and no follow up data. **Bok 2016** randomised children with painful flat foot in to one of four groups: (1) Sport shoes only (2) CFOs with no inverted angle (3) CFOs with a 15° inverted angle (4) CFOs with a 30° inverted angle. The other study randomised 30 children with asymptomatic flat foot, into 2 groups medical shoe versus a regular shoe with PFOs (**Aboutorabi 2013**).

Footwear advice or provision varied across trials. Six trials supplied the footwear, which was also analysed as an intervention (**Powell 2005**; **Wenger 1989**, **Gould 1989**; **Kanatli 2016**, **Aboutorabi 2013**, **Jafarnezhadgero 2018**). The supplied footwear varied from athletic footwear (**Powell 2005**, **Jafarnezhadgero 2018**) to medical/orthopaedic footwear (ankle-high boots). Footwear was advised in three studies (**Bok 2016**, **Morrison 2013**, **Whitford 2007**), and otherwise footwear was specified, and so presumably the participant's usual shoes were used. **Wenger 1989** provided a pedorthotist for all follow-up visits to ensure the corrective shoes were fitted according to the standards and specifications of the Prescription Footwear Association.

Both **Coda 2014** and **Jafarnezhadgero 2018** utilised sham foot orthoses for the control groups.

Whitford 2007 prescribed calf muscle stretches to all children who required stretching, irrespective of the study group, and a seven week rehabilitation programme was directed for both intervention and control groups in boys with DCD (**Morrison 2013**).

Outcomes

Overall, the 16 included trials used a wide range of outcome measures, with the majority measuring pain, function, health-related quality of life and foot X-rays; see **Table 2**; **Table 4**; and **Table 5**.

Trial durations and follow-up ranged from immediate (**Aboutorabi 2013**; **Bok 2016**), to five years (**Gould 1989**; **Kanatli 2016**). Follow-up ranged from four weeks to three years (approximately a two-year average) in the majority of trials in healthy children with flexible flat feet; three to six months in the JIA studies; and over seven weeks in the DCD study.

Major outcomes

Pain

Five studies measured pain on VAS (**Asgaonkar 2012**; **Coda 2014**; **Powell 2005**; **Whitford 2007**).

Studies which measured pain using a continuous outcome measure (VAS), reported the proportions of participants with or without pain at follow up (**Coda 2014**, **Powell 2005**). These studies were specific to children with JIA.

Wenger 1989 stated that parents reported a reduction in pain symptoms in children with flat feet across the four groups (corrective shoes, heel cup, UCBL inserts, and the control group) but no data were provided.

Hsieh 2018 measured pain using parented reported Paediatric outcome data collection instrument (PODCI), in children recruited with painful flat feet. PODCI (The Pediatric Outcomes Data Collection Instrument, Daltroy 1998) measures capability primarily, and includes a paediatric version to be completed by a parent and an adolescent version that can be completed by the parent, child, or both (Klepper 2011).

Whitford 2007 made similar report, with pain at baseline and trial completion, reported as participant percentages. This was a subgroup, as participant recruitment targeted asymptomatic, healthy children.

Whitford 2007 measured mean pain as a continuous outcome with VAS, but only reported the proportion with pain and the proportion with no pain (introducing a possible reporting bias). We extracted proportion with no pain, from the percentage data reported.

Hsieh 2018 did not directly measure pain, but pain/comfort are a psychometric property of the PODCI outcome measure used (Daltroy 1998).

Function

Overall, seven studies measured function (Aboutorabi 2013; Bok 2016; Hsieh 2018; Morrison 2013; Powell 2005; Solanki 2020; Whitford 2007). Four studies directly measured function (Powell 2005, Whitford 2007, Hsieh 2018; Solanki 2020). Five studies used gait as a measure of function (Aboutorabi 2013; Asgaonkar 2012; Bok 2016; Khamooshi 2017; Morrison 2013).

Powell 2005 measured function on the Foot Function index (FFI; Budiman 1991). The FFI was developed to measure the impact of foot pathology on function in terms of pain, disability and activity restriction.

Four studies measured timed walking (Powell 2005, Morrison 2013, Aboutorabi 2013, Hsieh 2018), using different methods, i.e. the six minute walk test (Morrison 2013), timed up and go test (Hsieh 2018), 50 feet walk (Powell 2005), step velocity (Aboutorabi 2013). The six minute walk test (6MWT) is validated (New Reference), with reference data for both healthy and disease paediatric status.

One study (Solanki 2020) assessed balance and agility, using vertical jump height (VJH) Montalvo 2021, single excursion balance test (SEBT) Gribble, 2012, Illinois agility test (IAT) Kutlu 2018.

Health-related quality of life

Four studies investigated HRQoL (Coda 2014; Hsieh 2018; Powell 2005; Whitford 2007).

Three studies measured health-related quality of life using the PedsQL which is a well validated measure of many aspects of HRQoL (Coda 2014; Hsieh 2018; Powell 2005), with multiple domains, and reference disease data (Varni 1996; Varni 2002). Whitford 2007 used as proxy, the Self Perception Profile for Children (with six subscales for scholastic competence, social acceptance, behavioural conduct, physical appearance, athletic competence, global self-worth).

Treatment success

No studies stipulated criteria for treatment success. However, four studies used gait parameters (Asgaonkar 2012; Gould 1989; Jafarnezhadgero 2018; Powell 2005).

Proportion of withdrawals due to adverse events

Five studies reported withdrawals due to adverse events (Wenger 1989; Whitford 2007; Asgaonkar 2012; Gould 1989; Powell 2005).

Proportion with adverse events

None if the included studies reported the proportion of children with adverse events, or serious adverse events.

Proportion with serious adverse events

Not reported in any of the included studies.

Minor outcomes

Five studies used radiographic imaging (Abd-Elmonem 2021; Ahn 2017; Gould 1989; Kanatli 2016; Whitford 2007).

Six studies measured gait assessment (Aboutorabi 2013; Asgaonkar 2012; Hsieh 2018; Jafarnezhadgero 2018; Morrison 2013; Powell 2005).

Two studies used plantar pressure measures (Bok 2016; Morrison 2013). Three studies used footprint measures (Gould 1989; Kanatli 2016; Khamooshi 2017).

Single studies used clinical photos (Wenger 1989), centre of pressure analysis (Aboutorabi 2013), joint range laxity (Kanatli 2016), and resting calcaneal stance position (RCSP (Ahn 2017)).

Two studies assessed physical expenditure (Asgaonkar 2012; Whitford 2007).

Two studies assessed clinical impressions (Wenger 1989; Whitford 2007).

Three studies recorded parent feedback (Wenger 1989, Coda 2014, Hsieh 2018).

Table 1 collates description of the 16 included trials.

Table 2 is the trial group comparison matrix.

Excluded studies

We excluded thirteen studies after full-text screening, either because of wrong study design or wrong population. Nine studies were non-randomised clinical trials (Benedetti 2011; Ford 2017; Hill 2020; Hurd 2010; MacKenzie 2012; Mosca 2010; Pothrat 2013; Riccio 2009; Uden 2017). Four studies did not include children (Yung 2011), did not address flat feet (Perhamre 2011; Perhamre 2012), or involved adult participants (Okamura 2020).

Additional details of the excluded studies are reported in the Characteristics of excluded studies table.

Ongoing studies

We identified 12 ongoing studies. Details are included in the table of Characteristics of ongoing studies.

Studies awaiting classification

We contacted investigators of two studies to verify key study characteristics and obtain missing numerical outcome data (Pandey 2013; Sinha 2013). We contacted both authors via email; but did not obtain the required data (answers to queries, missing data, clarification of randomisation, and standard deviations).

Details of both studies are included in the table of [Characteristics of studies awaiting classification](#).

Risk of bias in included studies

The summary of risk of bias is presented in [Figure 2](#). We did not judge any of the 16 included trials at low risk of bias across all domains. Most trials were at risk of selection, performance, detection, and selective reporting biases.

Figure 2. Risk of bias summary: review authors' judgements about each risk of bias item for each included study

	Random sequence generation (selection bias)	Allocation concealment (selection bias)	Blinding of participants and personnel (performance bias): All outcomes	Blinding of outcome assessment (detection bias) - self-reported outcomes (e.g., pain, function)	Blinding of outcome assessment (detection bias) - objective outcomes	Incomplete outcome data (attrition bias): All outcomes	Selective reporting (reporting bias)	Other bias
Abd-Elmonem 2021	+	+	?	+	+	+	+	+
Aboutorabi 2013	?	?	-	-	?	?	?	?
Ahn 2017	?	?	-	-	?	?	?	?
Asgaonkar 2012	?	?	-	-	-	-	?	?
Bok 2016	?	?	?	?	?	?	?	?
Coda 2014	+	?	-	+	?	+	?	?
Gould 1989	?	?	-	?	?	?	?	?
Hsieh 2018	+	+	-	-	-	+	-	+
Jafarnezhadgero 2018	?	?	+	-	+	+	?	+
Kanatli 2016	-	-	-	?	?	+	-	?
Khamooshi 2017	?	?	?	?	?	?	?	?
Morrison 2013	?	+	-	?	-	?	?	+
Powell 2005	+	+	-	-	-	?	?	?
Solanki 2020	+	+	+	+	+	+	?	+
Wenger 1989	+	?	-	+	?	?	?	?
Whitford 2007	+	?	-	-	+	+	?	?

Figure 2. (Continued)

Wenger 1989	+	?	-	+	?	?	?	?
Whitford 2007	+	?	-	-	+	+	?	?

Allocation

All trials reported that participants were randomised to an intervention, but only seven trials adequately described the method used to generate the random sequence (Abd-Elmonem 2021; Coda 2014; Hsieh 2018; Powell 2005; Solanki 2020; Wenger 1989; Whitford 2007) and are at low risk of selection bias for random sequence generation.

Three trials did not describe if allocation of treatment was concealed (Coda 2014; Wenger 1989; Whitford 2007) and are at unclear risk of bias. Only five trials adequately reported allocation concealment (Abd-Elmonem 2021; Hsieh 2018; Morrison 2013; Powell 2005; Solanki 2020) and are judged at low risk of selection bias for allocation concealment. We judged one trial at high risk of bias for both random sequence generation and allocation concealment (Kanatli 2016).

Blinding

We detected performance bias in most trials. We judged 11 out of the included 16 trials at high risk of performance bias (Aboutorabi 2013; Ahn 2017; Asgaonkar 2012; Coda 2014; Gould 1989; Hsieh 2018; Kanatli 2016; Morrison 2013; Powell 2005; Wenger 1989; Whitford 2007). We only judged two trials at low risk of performance bias (Jafarnezhadgero 2018; Solanki 2020). Due to the nature of the interventions, the participants could not be blinded.

We judged detection bias for self-reporting outcomes at low risk in four trials (Abd-Elmonem 2021; Coda 2014; Solanki 2020; Wenger 1989), and low risk for objective outcome assessment in four (Abd-Elmonem 2021; Jafarnezhadgero 2018; Solanki 2020; Whitford 2007). We judged detection bias for self-reporting outcomes at high risk in seven trials (Aboutorabi 2013; Ahn 2017; Asgaonkar 2012; Hsieh 2018; Jafarnezhadgero 2018; Powell 2005; Whitford 2007), and high risk for objective assessment in four (Asgaonkar 2012; Hsieh 2018; Morrison 2013; Powell 2005).

Incomplete outcome data

We judged seven trials at low risk of attrition bias (Abd-Elmonem 2021; Coda 2014; Hsieh 2018; Jafarnezhadgero 2018; Kanatli 2016; Solanki 2020; Whitford 2007), five of which reported the reasons participants withdrew from the study (Abd-Elmonem 2021; Coda 2014; Hsieh 2018; Kanatli 2016; Whitford 2007); two of which had no withdrawals (Jafarnezhadgero 2018; Solanki 2020). We assessed one trial at high risk of attrition bias due to an imbalance in the number of dropouts per group, and lack of reasons for withdrawal (Asgaonkar 2012). Apart from this, we judged that three studies were likely to have biased results due to participant withdrawals (Gould 1989; Morrison 2013; Wenger 1989), and it is largely unclear if the non-completing participants biased the results, as the study authors did not report to which treatment groups these participants were randomised. Gould 1989's attrition rate was unbalanced, with group 1- 25/50; group 2- 10/25; group 3- 7/25 ; group 4- 10/25 finishing the study. In total, 52/125 (42%) children completed the four year study. Likewise, considerable

attrition affected Wenger 1989 98/131 (75%) completions, and Morrison 2013 14/21 (68%) completions, reasons for attrition were not provided. Five studies provided insufficient information to permit a judgement of low risk or high risk of attrition bias, hence the risk of attrition bias was unclear (Aboutorabi 2013; Ahn 2017; Bok 2016; Khamooshi 2017; Powell 2005)

Selective reporting

We judged two trials at high risk of selective reporting bias, because of a discrepancy between the trial register and published outcomes, and the lack of follow-up data for outcomes measured at baseline (Hsieh 2018; Kanatli 2016). Hsieh 2018 provided more outcomes in the published paper than were listed in the trial registry, and Kanatli 2016 only provided baseline data for Arch index scores, no follow-up data.

We judged four studies at unclear risk of reporting bias, because of lack of gender-related data, and lack of follow-up data for outcomes measured at baseline (Bok 2016; Gould 1989; Morrison 2013; Wenger 1989). Gould 1989 stated that other lower extremity measurements (femoral, tibial torsions and knee varus/valgus) were to be subsequently reported, adding that knee valgus at age 5 years was "striking" (92.3%, of the 52/125 (42%) children who completed the 5 year trial) but without relationship to sex. In Wenger 1989 heel cord tightness and foot progression angles were only reported at baseline. Morrison 2013 assessed FPI-6 and the LLAS were used for patient selection at baseline, but not further reported. In an immediate effects trial, without follow up, Bok 2016 selected children with painful flat feet, but then omitted pain as an outcome.

We judged eight studies at unclear risk of reporting bias as there was insufficient information to permit a judgement of low risk or high risk (Aboutorabi 2013; Ahn 2017; Asgaonkar 2012; Coda 2014; Jafarnezhadgero 2018; Powell 2005; Solanki 2020; Whitford 2007). In one study, outcome data were not clearly reported (Khamooshi 2017).

Other potential sources of bias

Five trials were at low risk of other bias. We assessed that four of them had no source of other bias (Abd-Elmonem 2021; Hsieh 2018; Jafarnezhadgero 2018; Solanki 2020); the other one declared the source of funding, as well as independence from the funder for research design, conduct, and reporting (Morrison 2013). We judged all other trials at unclear risk of other bias due to insufficient information on funding sources.

Effects of interventions

See: **Summary of findings 1 Customised foot orthoses compared to shoes in children with asymptomatic flat feet**; **Summary of findings 2 Prefabricated foot orthoses compared to shoes in children with asymptomatic flat feet**; **Summary of findings 3 Custom foot orthoses compared to prefabricated foot orthoses for children with asymptomatic flat feet**; **Summary of findings**

4 Custom foot orthoses compared to shoes in children with juvenile idiopathic arthritis and flat feet; Summary of findings 5 Prefabricated foot orthoses compared to shoes in children with juvenile idiopathic arthritis and flat feet; Summary of findings 6 Custom foot orthoses compared to prefabricated foot orthoses in children with juvenile idiopathic arthritis and flat feet; Summary of findings 7 Prefabricated foot orthoses compared to shoes in children with symptomatic flat feet

The effects of interventions are discussed by diagnostic population groups, then comparisons. Due to the diversity of trials and the differences in the interventions and outcomes reported, we were only able to pool data for JIA in meta-analyses. Results of all other studies are reported separately.

1. Asymptomatic flat feet

Effects of interventions were assessed in 10 studies (Abd-Elmonem 2021; Ahn 2017; Asgaonkar 2012; Gould 1989; Jafarnezhadgero 2018; Kanatli 2016; Khamooshi 2017; Solanki 2020; Wenger 1989; Whitford 2007). Results are presented in Summary of findings 1.

Custom foot orthoses (CFOs) compared to shoes

Studies reported findings at baseline and three months (Wenger 1989), at baseline, 3 months and 12 months (Whitford 2007). Jafarnezhadgero 2018 reported CFOs versus sham FOs (both feet were fitted with the same sports shoes) at baseline and after 4 months. Aboutorabi 2013 reported an immediate effect, but given the absence of follow-up, we did not include the data in analyses.

Outcomes

- Pain* – CFOs may result in little to no difference in the proportion of children without pain (risk ratio (RR) 0.85, 95% confidence interval (CI) 0.67 to 1.07; 1 study, 106 participants; low-certainty evidence; Analysis 1.1). There was an absolute reduction of 11.8% (4.7% fewer to 15.8% more); we downgraded for bias and imprecision.
- Withdrawal due to adverse events* – CFOs result in little to no difference in withdrawal due to adverse events (RR 1.05, 95% CI 0.94 to 1.19; 3 studies, 211 participants; low-certainty evidence; Analysis 1.2). The absolute effect was 3.4% more (4.1% fewer to 13.1% more); we downgraded for bias and imprecision.
- Function, quality of life, treatment success, and adverse events were not reported.

Prefabricated foot orthoses (PFOs) compared to shoes

Four studies reported findings at three months (Wenger 1989), at 3 months and 12 months (Whitford 2007), at two years, three years, and five years of age (Gould 1989), and at one year (Asgaonkar 2012). Results are presented in Summary of findings 2.

Outcomes

- Pain* – PFOs likely result in little to no difference in the proportion of children without pain (RR 0.94, 95% CI 0.76 to 1.16; 1 study, 106 participants; low-certainty evidence; Analysis 2.1). The absolute reduction was 4.7% (18.9% fewer to 12.6% more); we downgraded for bias and imprecision.
- Withdrawals due to adverse events* – We are uncertain of the effects of PFOs on withdrawal due to adverse events (RR 0.99, 95% CI 0.79 to 1.23; 4 studies, 338 participants; very low-certainty evidence; Analysis 2.2). The absolute reduction was

0.7% (15.2% fewer to 16.6% more); we downgraded for bias, imprecision, and indirectness due to variably aged participant samples between studies.

- Function, quality of life, treatment success, and adverse events were not reported.

CFOs compared to PFOs

Studies reported findings at three months (Wenger 1989), and at 3 months and 12 months (Whitford 2007). Results are presented in Summary of findings 3.

Outcomes

- Pain* – CFOs likely result in little to no difference in pain (RR 0.93, 95% CI 0.73 to 1.18; 1 study, 108 participants; low-certainty evidence; Analysis 3.1). The absolute reduction was 7.4% (22.2% fewer to 11.1% more), we downgraded for bias and imprecision.
- Proportion of withdrawals* – CFOs result in no difference in withdrawals due to adverse events (RR 1.00, 95% CI 0.90 to 1.12; 1 study, 118 participants; low-certainty evidence; Analysis 3.2). We downgraded for bias and imprecision.
- Function, quality of life, treatment success, and adverse events were not reported.

Abd-Elmonem 2021 compared NMES and foot strengthening exercises with sham NMES and foot strengthening exercises, which did not comply with the comparison of interventions in this review.

Aboutorabi 2013 assessed immediate effects in both healthy children (N = 20) and those with flat feet (N = 30). The study reported that neither medical shoes nor PFOs showed significant gait effects in healthy children, however, the children with flat feet showed reduced centre of pressure (CoP) displacement with medical shoes (P < 0.05), and PFOs (not significant).

Ahn 2017 compared two types of CFO designs, which did not comply with the comparison of interventions in this review.

Kanatli 2016 compared an orthopaedic shoe with a control group, which did not comply with the comparison of interventions in this review.

Solanki 2020 compared anti-pronation and conventional treatment (foot strengthening exercises, Faradic foot baths), with sham taping and conventional treatment, which did not comply with the comparison of interventions in this review.

2. Flat feet in juvenile idiopathic arthritis (JIA), or other clinical concerns

Two studies assessed these populations; Coda 2014 followed up at three months and six months; Powell 2005 followed up at three months.

CFO compared to shoes

Results are presented in Summary of findings 4; one trial examined this comparison (Powell 2005).

Outcomes

- Pain* – CFOs likely result in little or no difference in pain (0 to 10 VAS, 0 = no pain) compared to shoes (MD -1.50, 95% CI -2.78 to -0.22; 1 study, 28 participants; very low-certainty

- evidence; [Analysis 4.1](#)). We downgraded for bias, imprecision, and indirectness; 1 study, 28 participants).
- Function or disability** – CFOs may result in a clinically important improvement in function or disability compared with shoes, measured with the FFI (MD -18.55, 95% CI -34.42 to -2.68; 1 study, 28 participants; low-certainty evidence; [Analysis 4.2](#); [Figure 3](#)); we downgraded for bias and imprecision.
 - Quality of life (QoL)** – CFOs may result in a clinically important improvement in child-rated QoL on the PedsQL (MD 12.10, 95% CI -1.60 to 25.80; 1 study, 25 participants; low-certainty evidence; [Analysis 4.3](#); [Figure 4](#)). CFOs may result in a clinically important improvement in parent-rated QoL on the PedsQL (MD 9.01, 95% CI -4.08 to 22.10; 1 study, 26 participants; low-certainty evidence; [Analysis 4.3](#); [Figure 4](#)); we downgraded for bias and imprecision as the 95% CIs include both an improvement and no improvement, the trend towards significance in the CFO group, with the 5-point minimally important clinical difference doubled - PedsQL/Child 12-point increase with CFO versus shoes, PedsQL/Parents 9-point increase with CFO versus shoes low-certainty evidence downgraded for bias and imprecision.
 - Treatment success** – CFOs likely result in little to no difference in timed walking on the 50-foot timed walk (50FTW; MD -1.33, 95% CI -2.77 to 0.11; 1 study, 28 participants; low-certainty evidence; [Analysis 4.4](#)); we downgraded for bias and imprecision.
 - Withdrawals due to adverse events** – CFOs likely result in little to no difference in withdrawals due to adverse events (RR 0.58, 95% CI 0.11 to 2.94; 1 study, 28 participants; low-certainty evidence; [Analysis 4.5](#); absolute reduction 9.7% (20.5% fewer to 44.8% more); we downgraded for bias and imprecision.
 - Proportion with adverse events** – none reported
 - Proportion with serious adverse events** – none reported

Figure 3. Forest plot of comparison: 4 CFOs versus shoes in JIA, outcome: 4.2 Function

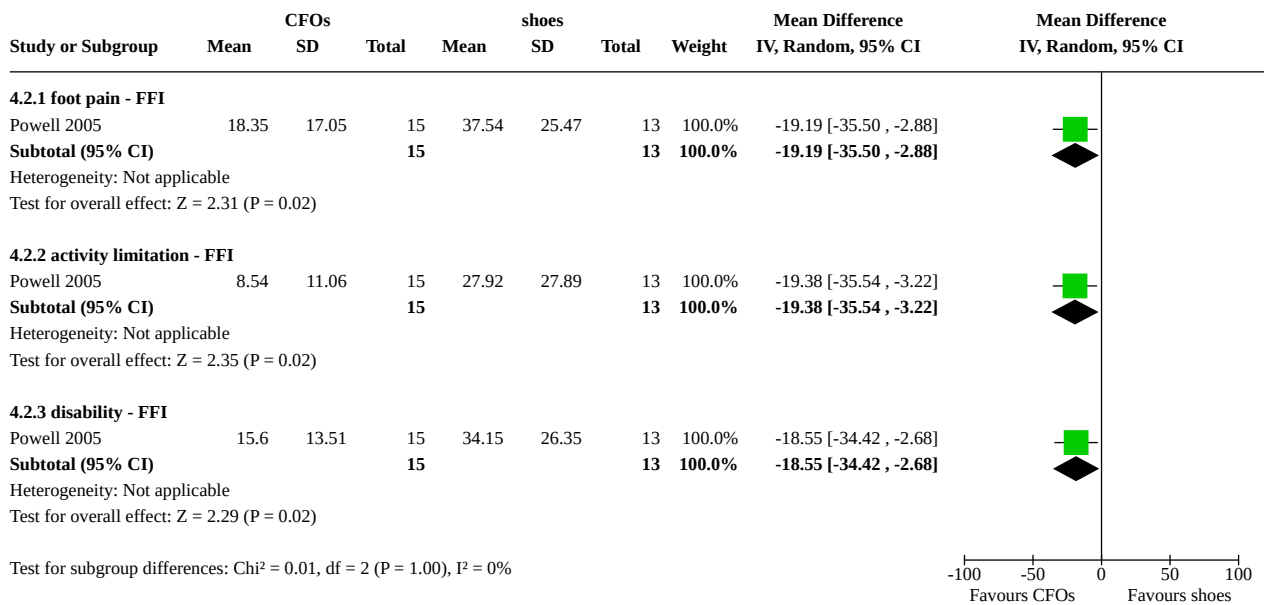
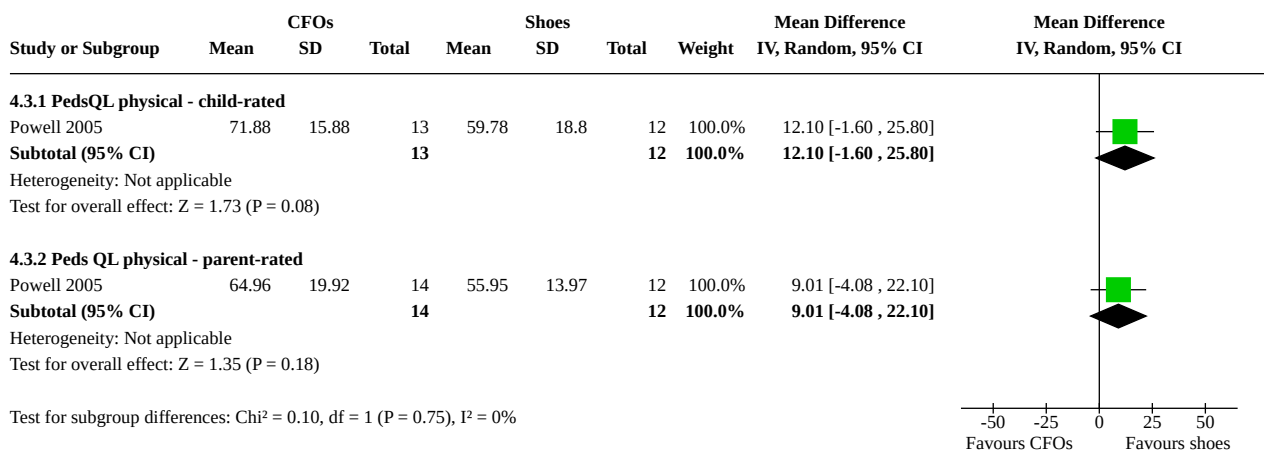


Figure 4. Forest plot of comparison: 4 CFOs versus shoes in JIA, outcome: 4.3 Quality of life



PFO compared to shoes

Results are summarised in [Summary of findings 5](#); one trial examined this comparison ([Powell 2005](#)).

Outcomes

- Pain** – PFOs likely result in little to no difference in pain (MD 0.02, 95% CI -1.94 to 1.98; 1 study, 28 participants; very low-certainty evidence; [Analysis 5.1](#)); we downgraded for bias, imprecision, and indirectness.
- Function or disability** – PFOs likely results in little to no difference in function (MD -4.17, 95% CI -24.40 to 16.06) or activity limitation between groups (MD -7.96, 95% CI -26.79 to 10.87; 1 study, 25 participants; low-certainty evidence; [Analysis 5.2](#)); we downgraded for bias and imprecision. such that the PFO group activity was less limited at 3-months, versus the shoe group, low-certainty evidence downgraded for bias and imprecision. [Powell 2005](#) also reported timed walking, which showed the PFO group slower than the shoe group over a distance of 50 feet, at 3-month follow up MD -0.38 (95% CI -1.90, 1.14).
- Quality of life** – PFOs likely results in little to no difference in child-rated QoL on the PedQL (MD -3.84, 95% CI -19.01 to 11.33; 1 study, 22 participants), indicating less pain in the PFO group low-certainty evidence downgraded for bias and imprecision. PFOs likely results in no difference in or parent-rated QoL on the PedQL (MD -0.64, 95% CI -13.22 to 11.94; 1 study, 22 participants; low-certainty evidence; [Analysis 5.3](#)); we downgraded for bias and imprecision. Overall, [Powell 2005](#) reported no significant improvement in QoL for either the PFO or the shoe group.
- Treatment success** – PFOs likely result in little to no difference in timed walking on the 50FTW (MD -0.38 seconds, 95% CI -1.9 to 1.14; 1 study, 25 participants; low-certainty evidence; [Analysis 5.4](#)); we downgraded for bias and imprecision.
- Withdrawal due to adverse events** – PFOs likely results in little to no difference in withdrawals due to adverse events (RR 0.72, 95% CI 0.14 to 3.61; 1 study, 25 participants; low-certainty evidence; [Analysis 5.5](#)); absolute reduction 6.5% (19.8% fewer to 60.2% more); we downgraded for bias and imprecision.
- Proportion with adverse events** – none reported
- Proportion with serious adverse events** – none reported

CFO compared to PFO

Results are summarised in [Summary of findings 6](#); we pooled data from two studies ([Coda 2014](#); [Powell 2005](#)).

Outcomes

- Pain** – CFOs may result in little to no difference in pain (MD -1.48, 95% CI -3.23 to 0.26; 2 studies, 87 participants; low-certainty evidence; [Analysis 6.1](#); [Figure 5](#)); we downgraded for bias and imprecision. Sensitivity analysis, performed by excluding the study that did not blind participants ([Powell 2005](#)), was not significant due to overlapping confidence intervals (MD -2.88; 95%CI -15.70 to 9.94).
- Function or disability** – CFOs may result in little to no difference in function, on the FFI (MD -14.38, 95% CI -30.22 to 1.46; 1 study, 27 participants; low-certainty evidence; [Analysis 6.2](#)); we downgraded for bias and imprecision.
- Quality of life** – CFOs may result in little to no difference in child-rated QoL, using the PedsQL (MD 8.64, 95% CI -3.90 to 21.18; 1 study, 83 participants) or parent-rated QoL, using the PedsQL (MD 2.94, 95% CI -11.00 to 16.88; 1 study, 84 participants; low quality evidence; [Analysis 6.3](#); [Figure 6](#)); we downgraded for bias and imprecision. QoL was also reported at six months by [Coda 2014](#); child-rated PedsQL scores were 89.67 (17.92) with custom-prefabricated foot orthoses (CPFOs) and 83.63 (27.14) with PFOs; parent-rated PedQL scores were 83.70 (31.5) for CPFOs and 84.47 (35.58) for PFOs, indicating marginally improved in QoL at six months rated by children, but not by parents, low-certainty evidence, downgraded for bias and imprecision.
- Treatment success** – CFOs may result in little difference in timed walking at 3 months, assessed with the 50FTW (MD -0.95 seconds, 95% CI -1.88 to -0.02; 1 study, 27 participants; low-certainty evidence; [Analysis 6.4](#)); we downgraded for bias and imprecision.
- Withdrawals due to adverse events** – CFOs may result in little difference in withdrawals due to adverse events (RR 0.80, 95% CI 0.13 to 4.87; 2 studies, 87 participants; low-certainty evidence; [Analysis 6.5](#)); we downgraded for bias and imprecision.
- Proportion with adverse events** – none reported
- Proportion with serious adverse events** – none reported

Figure 5. Forest plot of comparison: 6 CFOs versus PFOs in JIA, outcome: 6.1 Pain

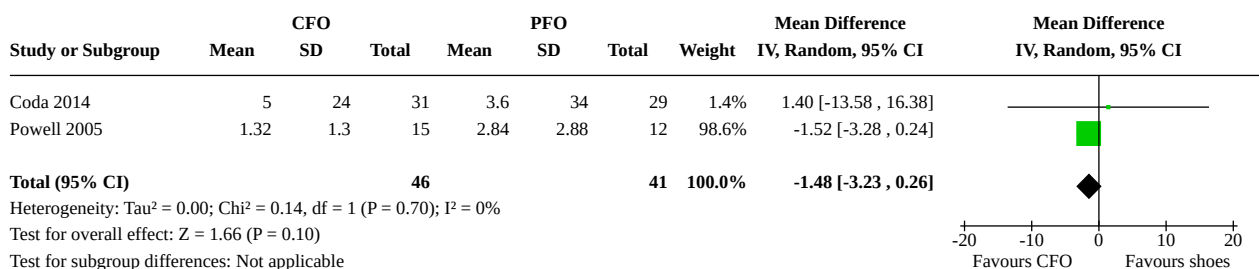
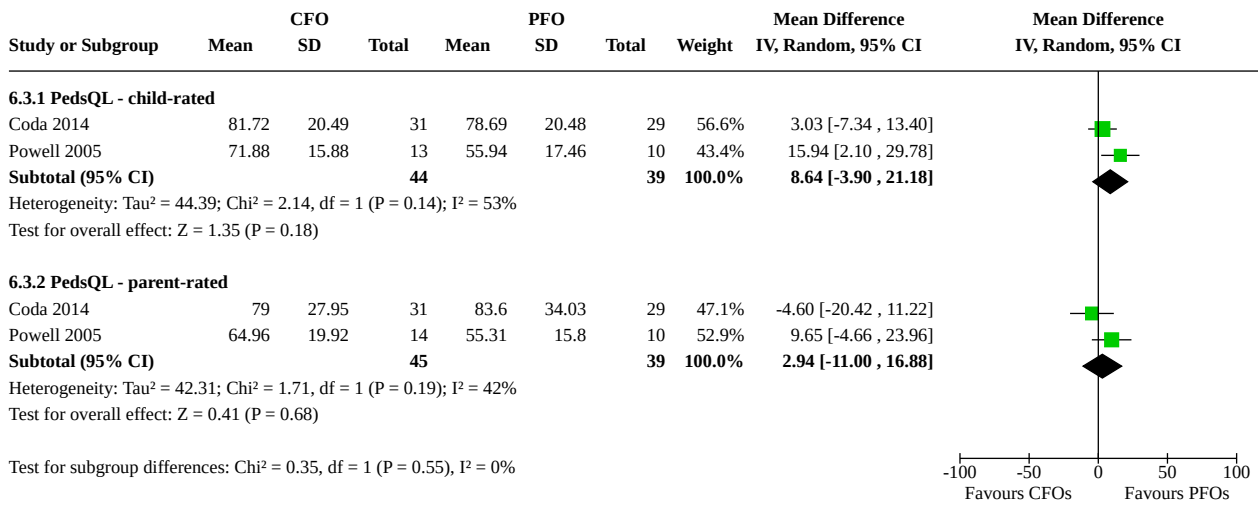


Figure 6. Forest plot of comparison: 6 CFOs versus PFOs in JIA, outcome: 6.3 Quality of life



3. Painful (symptomatic) flat feet

Assessed in two studies; Hsieh 2018 compared customised PFOs (adapted by heat gun moulding) versus no treatment and followed up at 12 weeks). Results are presented in Summary of findings 7. Bok 2016 outfitted all children with four sets of shoes with different degrees of orthotic, and measured immediately. Results are presented below.

PFO compared to shoes

Outcomes

1. Pain – not reported
2. Function or disability – PFOs result in little to no difference in function, assessed with PODCI (global function MD 3.00, 95% CI 2.28 to 3.72; 1 study, 50 participants; low-certainty evidence; Analysis 7.1); we downgraded for bias and imprecision.
3. Quality of life – PFOs result in little to no difference in QoL, assessed with the PedsQL (total score MD 1.80, 95% CI 1.07 to 2.53; 1 study, 50 participants; low-certainty evidence; Analysis 7.2); we downgraded for bias and imprecision.
4. Treatment success – not reported
5. Proportion of withdrawals – not reported
6. Proportion with adverse events – none reported
7. Proportion with serious adverse events – none reported

Bok 2016 (21 children) reported the immediate effects of treatment, three designs of CFOs, inverted at 0, 10, and 30 degrees, compared to shoes with no orthotics. Foot function was evaluated as peak pressure, maximum force, and contact area, using the Pedar in-shoe apparatus, in six foot regions (or 'masks'): medial forefoot (MF), central forefoot (CF), lateral forefoot (LF), medial midfoot (MM), lateral midfoot (LM), and rearfoot (RF).

Function or disability – CFOs with 0, 15, or 30 degrees inversion reduced MF and RF peak pressure (P < 0.001); increased maximum mid-foot plantar pressure (especially 30 degree inversion); and increased the contact area at the MM and RF.

None of the other outcomes were reported.

4. Developmental co-ordination disorder (DCD)

One study of 22 British boys with DCD examined the effects of foot orthoses (Morrison 2013).

CFO compared to shoes

Outcomes

1. Function or disability – the six-minute walk test (P = 0.43), and spatio-temporal parameters (Gaitrite™ system) did not differ between the CFO (N = 9) and the control (shoe) groups (N = 5) following the seven-week rehabilitation programme. The CFO group walked a median of four metres further, and the control group walked a median of 15 metres further (1 study, 14 participants; Analysis 8.1).
2. Treatment success – no differences were found between groups for the 6MWT (P = 0.43), cadence (P = 0.019), or double-support (P = 0.042), following the seven-week rehabilitation programme (1 study, 14 participants; Analysis 8.1).
3. Withdrawals due to adverse events – data for proportion of children who withdrew due to adverse events were not specified.
4. Pain, quality of life, and adverse events – not reported

DISCUSSION

Summary of main results

The 16 randomised controlled trials (RCT) included in this updated Cochrane Review signify more than a five-fold increase in available evidence from clinical trials since the previous review in 2010, which included just three RCTs. However, whilst there is more literature to peruse and critique, there continues to be heterogeneity, which limits data pooling and precludes meta-analysis of all studies. Two of the 16 trials provided only immediate effects data, and hence, we only included their description in this review, rather than including their data in any analyses.

There are two discrete groups of RCTs investigating foot orthoses to treat the typical paediatric flat foot: (1) those addressing healthy children with asymptomatic flat feet: 10 trials (Abd-Elmonem

2021; Ahn 2017; Asgaonkar 2012; Gould 1989; Jafarnezhadgero 2018; Kanatli 2016; Khamooshi 2017; Solanki 2020; Wenger 1989; Whitford 2007), and (2) those addressing flat feet in children with juvenile idiopathic arthritis (JIA): 2 trials (Powell 2005, Coda 2014); and other clinical concerns: painful flat feet: 1 trial (Hsieh 2018); developmental co-ordination disorder (DCD): 1 trial (Morrison 2013); immediate intervention effects only: 2 trials (Aboutorabi 2013, Bok 2016).

Asymptomatic flexible flat feet in healthy children

Firstly, and most prodigiously, 10 Ten RCTs addressed the most commonly presenting paediatric foot concern of asymptomatic flat feet; interventions included shoes, foot orthoses (FOs), and exercises. Five of these 10 RCTs included a non-intervention control group (Asgaonkar 2012; Kanatli 2016; Khamooshi 2017; Wenger 1989; Whitford 2007), three trials included a sham control group (Abd-Elmonem 2021; Jafarnezhadgero 2018; Solanki 2020), and two trials intervention comparisons alone (Ahn 2017; Gould 1989). Whilst all studies address the typical presentation of flexible and asymptomatic flat feet in healthy children, there is considerable heterogeneity within the 805 initially enrolled children: age ranged from 11 months to 15 years. Initial sample sizes ranged from 30 to 178, with seven trials enrolling less than 100 participants (Ahn 2017; Asgaonkar 2012; Jafarnezhadgero 2018; Kanatli 2016; Khamooshi 2017; Solanki 2020; Abd-Elmonem 2021). One trial included girls only (N=60) (Khamooshi 2017). One trial included boys only (N=30) (Jafarnezhadgero 2018). The period of follow up ranged from 4 weeks to 5 years and study retention ranged from 42% to 100% (655/805 completions, average 81%). Interventions tested varied across the 10 studies comprising foot orthoses (heel cups, UCBL orthoses, CFOs, PFOs, CPFOs, valgus insoles, TCFOs, RFOs), shoes (straight last, straight last with cookie, orthopaedic, orthopaedic with arch support, usual footwear, supplied trial footwear, medical shoes), taping, NMES, and exercises (foot exercises, foot and core exercises). The outcome measures varied across all studies: x-rays (5), pedoscope/footprints (3), pain (2), physical cost (2), self-perception (1), motor skills (2), static measures (3), gait (1), joint laxity (1), kinematic and kinetic gait analysis (1). The included studies were conducted across 30 years, between 1989 and 2021. The most recent trials (Solanki 2020; Abd-Elmonem 2021) discontinued the demonstrably fruitless investigation of foot orthoses for asymptomatic healthy children with flat feet, and implemented foot strength as the focus of intervention.

Flat feet in children with JIA, or other clinical concerns

Two RCTs addressed children with JIA (Powell 2005; Coda 2014), and provided data in a manner to enable us to pool them in meta-analysis (Fellas 2017b). The small sample sizes of the studies made inference insufficient to guide practice, i.e. in all group comparisons, the mean difference was larger than the minimal clinically important difference (MCID) of 8 mm on a 100 mm visual analogue scale (VAS) for pain at six months, within large confidence intervals (CI). Overall results were inconclusive to support the use of FOs in JIA foot and ankle pain.

Other clinical concerns of flat feet in children with foot pain, DCD, and trials with only immediate effects comprise the remaining four trials that addressed either immediate intervention effects with no follow-up data, in small samples with and without foot pain (Aboutorabi 2013; Bok 2016); a small sample with one-third

attrition over seven weeks in British boys identified as having a DCD (Morrison 2013); or a small sample of healthy children with painful flat feet (non-comparable study groups at baseline) over 12 weeks (Hsieh 2018). Separately, or combined, these four trials offer almost nothing, and at most, a possible trend. The inclusion of a trial addressing children with flat feet and foot pain whilst non-controversial, is also limited given the control group (only) received analgesics (Hsieh 2018).

Overall completeness and applicability of evidence

As a consequence of data heterogeneity, we were unable to conduct meta-analyses for the studies addressing asymptomatic flat foot presentation in healthy children. The JIA meta-analysis, with two studies, presents limited findings due to small samples. All studies were single-blind trials, with the investigators being aware of the type of intervention received, which may have resulted in performance and detection (assessor) bias. Blinded healthcare providers may also differ from non-blinded ones in their degree of attention to the children, or in their use of alternative forms of care. Time frames varied across the trials, ranging from four weeks to five years, making comparisons difficult. The age ranges differed across studies, and therefore, it is difficult to generalise about foot orthoses as a treatment for paediatric flat feet in children, aged 11 months to 16 years. There was limited technical information about the manufacturing process and prescription of the varying types of FOs used in the trials. Three studies recruited children of single sex (boys only (Jafarnezhadgero 2018; Morrison 2013); girls only (Khamooshi 2017)), hence, it is unclear as to whether results apply to both sexes. Twelve studies recruited both sexes. One study did not specify sex (Solanki 2020). The shoes used across the studies varied, the effects of which are unclear; see Table 3. However, the use of usual footwear favours external validity, and has wider community health relevance. Adverse effects were reported in one of the 16 trials, with no adverse effects from wearing FOs and shoes (Powell 2005). However, with only one trial reporting on this variable, and given the many limitations of the data available, caution must be noted. Gait analysis was included in four studies (Asgaonkar 2012; Bok 2016; Jafarnezhadgero 2018; Morrison 2013), but two studies only assessed the immediate effects of the intervention (Aboutorabi 2013; Bok 2016). More recent trials addressed foot strength, but provided no useful clinical information, given the small samples and short trial durations in healthy, asymptomatic participants (Abd-Elmonem 2021; Solanki 2020).

The main outcome measures reported in the 16 trials, included pain, function, and quality of life. Three studies investigated the primary outcome specified by this review (pain reduction), but differentially over three months and 12 months, in children with JIA (Coda 2014; Hsieh 2018; Powell 2005). It is unclear if the Varni-Thompson Paediatric Pain questionnaire (Varni 2002), used by Powell 2005 related to symptoms in the children's feet and lower extremities only, or if whole body pain was included. One trial conducted a post hoc subgroup analysis of pain reduction for those children who reported lower-limb pain at baseline, using a 0 to 10 VAS (Whitford 2007). However, the study design did not have an a priori hypothesis to specifically test the effects of CFOs or PFOs on the treatment of painful flat feet. Asgaonkar 2012 measured pain using a 0 to 10 VAS, and Hsieh 2018 assessed pain using the PODCI (Daltry 1998). Three studies directly investigated function as an outcome (Hsieh 2018; Powell 2005; Whitford 2007), whilst a further

five studies afforded deference to gait as function (Aboutorabi 2013; Asgaonkar 2012; Bok 2016; Khamooshi 2017; Morrison 2013). The intervention effects after seven weeks in children with DCD (Morrison 2013), after 8 weeks in children with JIA (Powell 2005), after 12 weeks in children with painful flat feet (Hsieh 2018), and after 1 year in children with asymptomatic flat feet (Whitford 2007) were unsurprisingly, variable. Given the especially disparate clinical presentations, range of follow-up periods, and small sample sizes of these studies, only the substantial trial by Whitford 2007 (N = 178), warrants further comment. Whitford 2007 assessed both motor proficiency and exercise efficiency, and found no difference between the two intervention groups (CFO and PFO) versus the control group after 12 months.

Four studies investigated quality of life in children with JIA and foot pain (Coda 2014; Powell 2005), in children with asymptomatic flat feet (Whitford 2007), and in children with painful flat feet (Hsieh 2018). radiographic outcomes (x-rays) were assessed in five studies in children with pain-free flat feet (Abd-Elmonem 2021; Ahn 2017; Gould 1989; Kanatli 2016; Whitford 2007), with follow-up ranging from four months to five years (Table 5). Two trials involved very young children (11 to 14 months (Gould 1989), and 17 to 72 months (Kanatli 2016), which captured the expectedly more pronounced, and normal developmental flat foot posture and indistinct foot skeletal morphology, given the physiologic age (Evans 2012; Gijon-Nogueron 2019; Martinez-Nova 2018; Pfeiffer 2006). Unsurprisingly, the studies without control groups reported 'improvement' of one intervention group over another, with no meaningful comparison from baseline and differing sample ages and sizes (Ahn 2017; Gould 1989). Two studies with control groups found no difference in x-ray findings between groups (Kanatli 2016; Whitford 2007), however, Abd-Elmonem 2021 reported significant improvement in x-ray parameters after four months of NMES versus sham-NMES (both treatment and control groups also improved from receiving foot strength exercises).

The intervention effects after seven weeks in children with DCD (Morrison 2013), after 8 weeks in children with JIA (Powell 2005), after 12 weeks in children with painful flat feet (Hsieh 2018), and after 1 year in children with asymptomatic flat feet (Whitford 2007) were unsurprisingly, variable. Given the especially disparate clinical presentations, range of follow-up periods, and small sample sizes of these studies, only the substantial trial by Whitford 2007 (N = 178), warrants further comment. Whitford 2007 assessed both motor proficiency and exercise efficiency, and found no difference between the two intervention groups (CFO and PFO) versus the control group after 12 months. Two studies with control groups found no difference in x-ray findings between groups (Kanatli 2016; Whitford 2007), however, Abd-Elmonem 2021 reported significant improvement in x-ray parameters after four months of NMES versus sham-NMES (both treatment and control groups also improved from receiving foot strength exercises).

Quality of the evidence

We assessed the quality evidence using GRADE for the major outcomes. We rated the evidence at low or very low certainty. No study returned high quality evidence for any outcome. Two studies were found to be of moderate quality evidence (Whitford 2007, Coda 2014). The remainder of the studies were rated as low and very low quality of evidence across all outcomes. We found 12 unpublished trials (Ongoing studies), which includes seven small samples of healthy children, 2 samples of children with JIA, 1

sample of children with neuromuscular disorders, and one large sample (N=1055) registered as two trials (Characteristics of ongoing studies).

For the major outcomes in asymptomatic feet, evidence for pain, function, quality of life and withdrawals due to adverse events was downgraded to low-certainty due to possible selection, performance and detection biases, as well as imprecision due to data from mostly single studies. For the major outcomes in symptomatic feet in children with JIA, evidence for pain was downgraded to very low-certainty due to detection bias as children and their parents knew which treatment they had, which may have affected their assessment of pain; as well as due to imprecision and indirectness. Evidence was mostly of low-certainty for function, treatment success, quality of life and withdrawal due to adverse events in children with JIA, downgraded for bias and imprecision.

The GRADE findings are reported in the SoF tables Pain reduction was assessed in five trials (Asgaonkar 2012; Coda 2014; Powell 2005; Wenger 1989; Whitford 2007), improvement in gait and function was assessed in six trials (Aboutorabi 2013; Bok 2016; Khamooshi 2017; Morrison 2013; Powell 2005; Whitford 2007; Solanki 2020), and QoL in three trials (Coda 2014; Powell 2005; Whitford 2007). The findings from four trials including x-ray imaging (Kanatli 2016; Ahn 2017; Gould 1989; Wenger 1989), are included in the recent systematic review of specifically x-ray findings (Choia 2020), which similarly found low level certainty from the evidence.

Due to the poor methodological quality of the trials, and heterogeneity of the studies within this review, definite conclusions could not be made. Further, the necessity of further attention to healthy children with flexible, pain-free flat feet is now demonstrably unfounded (Kanatli 2016, Martinez-Nova 2018).

Potential biases in the review process

Two authors were involved in the previous systematic review in 2010 (Rome 2010). AE has several publications in this field. AE and KR, are co-authors on an earlier literature review (MacKenzie 2012). However, all aspects of this review have been carried out with author independence, decisions reached by consensus, and reviewed by the Cochrane Musculoskeletal Group. We followed the Cochrane methodology to reduce sources of potential bias in the review process.

Agreements and disagreements with other studies or reviews

This updated Cochrane Review expanded the previous findings of Rome 2010, which stated that low quality evidence negated conclusive evidence on the benefits of non-surgical interventions for flat feet. These findings are largely in agreement with the conclusions of a prior critical review (MacKenzie 2012).

Low quality evidence from studies addressing the asymptomatic flexible paediatric flat foot have occupied too much of the medical literature, and for too long (Evans 2021). Another systematic review, which included studies from lower levels of the evidence hierarchy, concluded: "FOs show potential as a treatment method for children with flexible pes planus" despite the poor methodological quality of studies (Dars 2018). A more recent systematic review evaluated the long-term structural effect of orthoses for paediatric flexible flatfoot and did not support its use on the medial longitudinal arch, determined by radiographic imaging (Choia 2020). Further, it was

revealed that flat feet in young children improved with growth, regardless of the type of footwear used. This is now a repeated theme across the literature (Gijon-Nogueron 2019; Kanatli 2016; Martinez-Nova 2018; Pfeiffer 2006; Wenger 1989; Whitford 2007), and there is no strong evidence that the long-term use of foot orthoses improved the structure of flat feet in children (Choi 2020).

Hence, a decade later, more trials have contributed only low level evidence. It is concerning that this situation is continued in the majority of currently registered trials. We identified several ongoing trials, yet to be completed and published (ACTRN12616001082493; CTRI/2018/07/014989; CTRI/2019/08/020925; IRCT2017082235517N1; ISRCTN14602568; ISRCTN49672274; KCT0001717; NCT02414087; NCT02633566; NCT03151538; NCT04104555; NCT04410926). Concerningly, most of these trials are designed similarly to those we included in this review, i.e. most address the use of foot orthoses as an intervention for asymptomatic paediatric flat feet, in small samples of young and healthy children.

AUTHORS' CONCLUSIONS

Implications for practice

We conclude that there is no evidence to support the efficacy of foot orthoses (FO) for children with asymptomatic (painless) flat feet. The evidence from randomised controlled trials (RCT) is thwarted by biases affecting the study quality, diverse participants, varied clinical presentations, ranging interventions, disparate outcome measures, small sample sizes, high attrition, and lack of follow-up.

Further, the evidence across four decades remains very limited, with ill-defined conclusions about foot orthoses for treating paediatric flat feet. Very-low to low-certainty evidence from two studies evaluating the effectiveness of FOs for foot and ankle pain in children with juvenile idiopathic arthritis (JIA) found inconclusive evidence on the benefits of foot orthoses on pain, child- and parent-rated quality of life, quality of life, or withdrawals due to adverse events.

Whilst less frequent, children with painful flat feet are directed to use low-cost PFOs with well-selected footwear. Footwear, a given influence for FOs, is demonstrated to alter gait and foot mobility, and hence requires regard in both clinical and research settings. Available normative foot posture data, triage principles, and wider diagnoses must be considered before any intervention occurs.

Given the lack of difference between the low cost PFOs and the expensive CFOs, and the benefits of early treatment, clinicians may consider immediate use of PFOs for JIA foot and ankle pain. JIA is indisputably consequential, and many children experience disabling foot and lower limb pathology despite best available pharmaceutical measures.

It is hoped that this review will inform health professionals, researchers, parents, and children alike, so that concerns about paediatric flat foot presentations are triaged in accordance with best available evidence.

Implications for research

This review identified low to very low-certainty evidence that the effect of foot orthoses on pain, function, and quality of life in children with asymptomatic (painless) flat feet is uncertain.

Recently, the normal, age foot posture in children was published, and shows that children are expected to have varying flat feet as a part of normal growth (without foot pain, left and right feet looking similar and flat feet should reduce with age - angelaevanspodiatrists.com.au/evidence-essentials-blog-8-june-2019/). The availability of normative and prospective foot development data, dismisses most flatfoot concerns, and negates continued attention to this topic. The agenda for researching flat feet in children should target children with indisputable foot pathology from discrete diagnoses, namely, foot pain and diagnoses of JIA; syndromes associated with hypermobility (e.g. Down, Ehlers-Danlos, Marfan, etc); neuromuscular conditions (e.g. cerebral palsy; muscular dystrophy or atrophy), and perhaps conditions with rising public health implications (e.g. physical inactivity, obesity, hypertension, diabetes). In such warranted investigations, the use of similar and validated outcome measures would allow the combination of results, and the ability to pool estimates, to obtain meaningful consensus. Any future trials in relevant cohorts must be adequately powered with participants, and evaluate whether any group differences are both real and clinically worthwhile. Short-term benefit should be established, in order to justify the considerable resources and ethical implications for lengthy studies. Follow-up periods of at least five years are needed due to lower limb and foot growth in children. This will avail observation in children that may predict any predisposition to foot and gait problems as adults. If no such differences are observed, there would be no need to treat flat feet in children, regardless of the aetiology.

Dispelling misconceptions regarding paediatric foot posture will save resources in the form of clinical consultations, the cost of unnecessary intervention, and misidentification of 'deficiency' (Evans 2021). There are two ongoing trials addressing children with JIA (ACTRN12616001082493; ISRCTN49672274), and one large scale trial for children with symptomatic flat feet (NCT04104555), but many current trials will add little to the body of evidence, and probably waste research resources.

The agenda for researching asymptomatic flat feet in healthy children is now firmly closed, as there is no justification for wasting research and healthcare resources on flat feet in healthy children that do not hurt. Instead, a new and targeted research agenda, addressing children with indisputable foot pathology and associated diagnoses is definitely indicated, and further, encouraged. Future updates of this review will address only relevant paediatric foot conditions.

ACKNOWLEDGEMENTS

The authors wish to thank Renea Johnston, Managing Editor, Cochrane Musculoskeletal Group, for updating the search strategy, and for helpful advice, and Sheila Cyril, Assistant Managing Editor, Cochrane Musculoskeletal Group for running the updated searches. We would like to thank the following people for translating articles into English: Miranda Cumpston, formerly of the Cochrane Musculoskeletal Group; Gustavo Zanoli, Cochrane Musculoskeletal Group; Anette Bluemle, German Cochrane Centre; Lorenzo Moja, Italian Cochrane Centre; and Malgorzata Bala, Department of Internal Medicine, Jaiellonian University Medical College, Krakow, Poland. We thank the peer reviewers: Dr Andrea Coda, Senior Lecturer in Podiatry, PhD, Hons (BSc) Pod. School of Health Sciences, College of Health, Medicine and Wellbeing, The University

of Newcastle. Priority Research Centre Health Behaviour (PRCHB),
Hunter Medical Research Institute (HMRI), NSW, Australia, and Dr

Helen A Banwell (Ph.D), Allied Health and Human Performance,
University of South Australia, Adelaide, South Australia.

REFERENCES

References to studies included in this review

Abd-Elmonem 2021 {published data only}

Abd-Elmonem AM, El-Negamy EH, Mahran MA, Ramadan AT. Clinical and radiological outcomes of corrective exercises and neuromuscular electrical stimulation in children with flexible flatfeet: a randomized controlled trial. *Gait & Posture* 2021;**88**:297-303.

Aboutorabi 2013 {published data only} [10.1177/0309364613496111](https://doi.org/10.1177/0309364613496111)

Aboutorabi A, Saeedi H, Kamali M, Farahmand B, Eshraghi A, Dolagh RS. Immediate effect of orthopedic shoe and functional foot orthosis on center of pressure displacement and gait parameters in juvenile flexible flat foot. *Prosthetics and Orthotics International* 2014;**38**(3):218-23.

Ahn 2017 {published data only}

Ahn SY, Bok SK, Kim BO, Park IS. The effects of talus control foot orthoses in children with flexible flatfoot. *Journal of the American Podiatric Medical Association* 2017;**107**(1):46-53. [DOI: [10.7547/15-045](https://doi.org/10.7547/15-045)]

Asgaonkar 2012 {published data only} [10.1.1.464.883](https://doi.org/10.1.1.464.883)

Asgaonkar B, Kadam P. Effectiveness of valgus insole on pain, gait parameters and physiological cost index of walking in flat feet in 5 to 15 years. *Indian Journal of Physiotherapy and Occupational Therapy* 2012;**6**(2):90-4.

Bok 2016 {published data only}

Bok SK, Lee H, Kim BO, Ahn S, Song Y, Park I. The effect of different foot orthosis inverted angles on plantar pressure in children with flexible flatfeet. *PLoS One* 2016;**11**(7):1-10.

Coda 2014 {published data only} [10.1136/archdischild-2013-305166](https://doi.org/10.1136/archdischild-2013-305166)

Coda A, Fowle PW, Davidson JE, Walsh J, Carline T, Santos D. Foot orthoses in children with juvenile idiopathic arthritis: a randomised controlled trial. *Archives of Disease in Childhood* 2014;**99**(7):649-51.

Gould 1989 {published data only}

Gould N, Moreland M, Alvarez R, Trevino S, Fenwick J. Development of the child's arch. *Foot & Ankle International* 1989;**9**:241-5.

Hsieh 2018 {published data only} [10.1097/MD.00000000000010655](https://doi.org/10.1097/MD.00000000000010655)

Hsieh RL, Peng HL, Lee WC. Short-term effects of customized arch support insoles on symptomatic flexible flatfoot in children. A randomized controlled trial. *Medicine* 2018;**97**(20):1-9.

Jafarnezhadgero 2018 {published data only} [10.1371/journal.pone.0205187](https://doi.org/10.1371/journal.pone.0205187)

Jafarnezhadgero A, Madadi-Shad M, Alavi-Mehr SM, Granacher U. The long-term use of foot orthoses affects walking kinematics and kinetics of children with flexible flat feet: A randomized controlled trial. *PLoS One* 2018;**13**(10):e0205187.

Kanatli 2016 {published data only} [10.1016/j.jos.2016.04.014](https://doi.org/10.1016/j.jos.2016.04.014)

Kanatli U, Aktas E, Yetkin H. Do corrective shoes improve the development of the medial longitudinal arch in children with flexible flat feet? *Journal of Orthopaedic Science* 2016;**21**(5):662-6.

Khamooshi 2017 {published data only}

Khamooshi R, Mohammadieh SM, Rahnema N, Zalani FR. Comparing the effects of simultaneous eight-week stretching/strengthening training with core stability exercises on the flat foot deformity of 9 to 13 year old female students. *International Journal of Musculoskeletal Pain Prevention* 2016;**1**(4):149-56.

Morrison 2013 {published data only} [10.1016/j.ridd.2013.06.012](https://doi.org/10.1016/j.ridd.2013.06.012)

Morrison S, Ferrari J, Smillie S. Assessment of gait characteristics and orthotic management in children with developmental coordination disorder: preliminary findings to inform multidisciplinary care. *Research in Developmental Disabilities* 2013;**34**(10):3197-201.

Powell 2005 {published data only}

Powell M, Seid M, Szer IS. Efficacy of custom foot orthotics in improving pain and functional status in children with juvenile idiopathic arthritis: a randomised trial. *Journal of Rheumatology* 2005;**32**:943-50.

Solanki 2020 {published data only}

Priyanka S, Murugan S. Anti pronation taping and its effect on balance, explosive power and agility of school going children with flat foot. *Indian Journal of Public Health Research & Development* 2020;**11**(7):868-73.

Wenger 1989 {published data only}

Wenger DR, Mauldin D, Spleck G, Morgan D, Lieber RL. Corrective shoes and inserts as treatment for flexible flatfoot in infants and children. *Journal of Bone and Joint Surgery* 1989;**71**(6):800-10.

Whitford 2007 {published data only}

Whitford D, Esterman A. A randomised controlled trial of two types of in-shoe orthoses in children with flexible excess pronation of the feet. *Foot & Ankle International* 2007;**28**:715-23.

References to studies excluded from this review

Benedetti 2011 {published data only} [10.3928/01477447-20101221-04](https://doi.org/10.3928/01477447-20101221-04).

Benedetti MG, Ceccarelli F, Berti L, Luciani D, Catani F, Boschi M, Giannini S. Diagnosis of flexible flatfoot in children: a systematic clinical approach. *Orthopedics* 2011;**34**(2):94.

Ford 2017 {published data only} <http://dx.doi.org/10.1016/j.fcl.2017.04.008>

Ford SE, Scannell BP. Pediatric flatfoot: pearls and pitfalls. *Foot and Ankle Clinics* 2017;**22**(3):643-56.

Hill 2020 {published data only}

Hill M, Healy A, Chockalingam N. Effectiveness of therapeutic footwear for children: a systematic review. *Journal of Foot and Ankle Research* 2020;**13**(23):1-26. [DOI: doi.org/10.1186/s13047-020-00390-3]

Hurd 2010 {published data only}

Hurd WJ, Kavros SJ, Kaufman KR. Comparative biomechanical effectiveness of over-the-counter devices for individuals with a flexible flatfoot secondary to forefoot varus. *Clinical Journal of Sport Medicine* 2010;**20**(6):428-35.

MacKenzie 2012 {published data only}

MacKenzie AJ, Rome K, Evans AM 2012 Dec 1, 32(8):830-4. The efficacy of nonsurgical interventions for pediatric flexible flat foot: a critical review. *Journal of Pediatric Orthopaedics* 2012;**32**(8):830-4.

Mosca 2010 {published data only} [10.1007/s1.1832-010-0239-9](https://doi.org/10.1007/s1.1832-010-0239-9)

Mosca V. Flexible flatfoot in children and adolescents. *Journal of Children's Orthopaedics* 2010;**4**(2):107-21.

Okamura 2020 {published data only (unpublished sought but not used)}

* Okamura K, Fukuda K, Oki S, Ono T, Tanaka S, Kanai S. Effects of plantar intrinsic foot muscle strengthening exercise on static and dynamic foot kinematics: A pilot randomized controlled single-blind trial in individuals with pes planus. *Gait Posture* 2020 Jan;**75**:40-45.

Perhamre 2011 {published data only}

Perhamre S, Lundin F, Norlin R, Klässbo M. Sever's injury; treat it with a heel cup: a randomized, crossover study with two insole alternatives. *Scandinavian Journal of Medicine & Science in Sports* 2011;**21**(6):e42-7.

Perhamre 2012 {published data only}

Perhamre S, Lundin F, Klässbo M, Norlin R. A heel cup improves the function of the heel pad in Sever's injury: effects on heel pad thickness, peak pressure and pain. *Scandinavian Journal of Medicine & Science in Sports* 2012;**22**(4):516-22.

Pothrat 2013 {published data only}

Pothrat C, Authier G, Viehweger E, Rao G. Multifactorial gait analysis of children with flat foot and hind foot valgus deformity. *Computer Methods in Biomechanics and Biomedical Engineering* 2013;**16**(Suppl 1):80-1.

Riccio 2009 {published data only}

Riccio I, Gimigliano F, Gimigliano R, Porpora G, Iolascon G. Rehabilitative treatment in flexible flatfoot: a perspective cohort study. *Musculoskeletal Surgery* 2009;**93**(3):101-7.

Uden 2017 {published data only} [10.1186/s13047-017-0218-1](https://doi.org/10.1186/s13047-017-0218-1)

Uden H, Scharfbillig R, Causby R. The typically developing paediatric foot: how flat should it be? A systematic review. *Journal of Foot and Ankle Research* 2017;**10**(37):1-17.

Yung 2011 {published data only}

Jung D-Y, Koh E-K, Kwon O-Y. Effect of foot orthoses and short-foot exercise on the cross-sectional area of the abductor hallucis muscle in subjects with pes planus: a randomized controlled trial. *Journal of Back and Musculoskeletal Rehabilitation* 2011;**24**:225-31.

References to studies awaiting assessment
Pandey 2013 {published data only}

Pandey S, Pal CP, Kumar D, Singh P. Flatfoot in Indian population. *Journal of Orthopaedic Surgery* 2013;**21**(1):32-6.

Sinha 2013 {published data only} [10.1177/230949901302100111](https://doi.org/10.1177/230949901302100111)

Sinha S, Song HR, Kim HJ, Park MS, Yoon YC, Song SH. Medial arch orthosis for paediatric flatfoot. *Journal of Orthopaedic Surgery* 2013;**21**(1):37-43.

References to ongoing studies
ACTRN12616001082493 {unpublished data only}

Effectiveness of foot orthoses in children with juvenile idiopathic arthritis: a randomised control trial. Ongoing study. not stated. Contact author for more information.

CTRI/2018/07/014989 {unpublished data only}

Effect of adhesive taping and exercises on outward rotation of foot in affected children. Ongoing study. not stated. Contact author for more information.

CTRI/2019/08/020925 {published and unpublished data}

CTRI/2019/08/020925. Bare foot walking effect on flat foot in school going children [Effects of bare foot walking on the flat foot in school going children: a randomised control trial]. www.ctri.nic.in/Clinicaltrials/pmaindet2.php?trialid=36115 (first registration 27 August 2019).

IRCT2017082235517N1 {unpublished data only} en.irct.ir/trial/26811

Effects of long-term use of arch support foot orthoses on lower limb kinematics and kinetics during walking in children with flexible flat foot. Ongoing study. 4 September 2017. Contact author for more information.

ISRCTN14602568 {unpublished data only} www.isrctn.com/ISRCTN14602568

Orthotics for treatment of symptomatic flat feet in children - the OSTRICH study. Ongoing study. 18 May 2020. Contact author for more information.

ISRCTN49672274 {unpublished data only}

Foot disease in juvenile idiopathic arthritis: foot care trial. Ongoing study. not stated. Contact author for more information.

KCT0001717 {unpublished data only}

The effects of talus control foot orthosis in children with flexible flatfoot. Ongoing study. 7 December 2015. Contact author for more information.

NCT02414087 {unpublished data only}

Therapeutic effects of customized insoles on children with flatfoot. Ongoing study. not stated. Contact author for more information.

NCT02633566 {unpublished data only}

Clinical trial of the effect of functional orthoses in children with flat feet. Ongoing study. not stated. Contact author for more information.

NCT03151538 {unpublished data only}

Effects on pes planus exercise training mixed with play on pre-school children. Ongoing study. not stated. Contact author for more information.

NCT04104555 {unpublished data only}

Orthotics for treatment of symptomatic flat feet in children - the OSTRICH study. Ongoing study. 19 September 2019. Contact author for more information.

NCT04410926 {unpublished data only}

Corrective exercises with neuromuscular electrical stimulation in children with flexible flat feet. Ongoing study. 1 September 2018 to 31 December 2019. Contact author for more information.

Additional references
Aharonson 1992

Aharonson Z, Arcan M, Steinback TV. Foot-ground pressure pattern of flexible flatfoot in children with and without correction of calcaneovalgus. *Clinical Orthopaedics and Related Research* 1992;**181**:177-82.

Bauer 2015

Bauer K, Mosca VS, Zionts LE. What's new in paediatric flatfoot? *Journal of Pediatric Orthopedics* 2016;**36**(8):865-9.

Blitz 2010

Blitz NM, Stabile RJ, Giorgini RJ, Di Domenico LA. Flexible pediatric and adolescent pes planovalgus: conservative and surgical treatment options. *Clinics in Podiatric Medicine and Surgery* 2010;**27**(1):59-77.

Bresnahan 2009

Bresnahan P. The flat-footed child – to treat or not to treat: what is the clinician to do? *Journal of American Podiatric Medical Association* 2009;**99**(3):178.

Brooks 1991

Brooks MH. Flatfeet in children. *BMJ* 1991;**302**:237.

Budiman 1991

Budiman-Mak E, Conrad KJ, Roach K. The Foot Function Index: a measure of foot pain and disability. *Journal of Clinical Epidemiology* 1991;**44**:561-70.

Capello 1998

Cappello T, Song KM. Determining treatment of flatfeet in children. *Current Opinion in Pediatrics* 1998;**10**:77-81.

Carli 2012

Carli A, Saran N, Kruijt J, Alam N, Hamdy R. Physiological referrals for paediatric musculoskeletal complaints: a costly problem that needs to be addressed. *Paediatrics and Child Health* 2012;**17**(e):93-7.

Cates 2016 [Computer program]

Visual Rx. Cates C, Version 4. Hertfordshire (UK): Dr Chris Cates, 2016. available at www.nntonline.net.

Choia 2020

Choi JY, Hong WH, Suh JS, Han JH, Lee DJ, Lee YJ. The long-term structural effect of orthoses for pediatric flexible flat foot: a systematic review. *Foot and Ankle Surgery* 2020;**26**(2):181-8. [DOI: doi.org/10.1016/j.fas.2019.01.007]

D'Amico 2009

D'Amico JC. The flat-footed child – to treat or not to treat: what is the clinician to do? *Journal of American Podiatric Medical Association* 2009;**99**(3):267-8.

Daltroy 1998

Daltroy LH, Liang MH, Fossel AH, Goldberg MJ. The POSNA pediatric musculoskeletal functional health questionnaire: report on reliability, validity, and sensitivity to change. *Journal of Pediatric Orthopaedics* 1998;**18**(5):561-71.

Daneshmandi 2011

Daneshmandi H, Azhdari F, Saki F, Daneshmandi MS. The study of lower limb alignment in athletes with and without ACL reconstruction. *Brazilian Journal of Biomotricity* 2011;**5**(4):248-54.

Dars 2018

Dars S, Uden H, Banwell HA, Kumar S. The effectiveness of non-surgical intervention (Foot Orthoses) for paediatric flexible pes planus: a systematic review: update. *PLoS One* 2018;**13**(2):e0193060.

Deeks 2020

Deeks JJ, Higgins JPT, Altman DG, editor(s). Chapter 10. Analysing data and undertaking meta-analyses. In: Higgins JPT, Thomas J, Chandler J, Cumpston M, Li T, Page MJ, et al, editor(s). *Cochrane Handbook for Systematic Reviews of Interventions* Version 6.1. Cochrane, 2020. Available from training.cochrane.org/handbook/archive/v6.1.

Evans 2008

Evans AM. The flat-footed child—to treat or not to treat: what is the clinician to do? *Journal of the American Podiatric Medical Association* 2008;**98**(5):386-93.

Evans 2009

Evans AM, Nicholson H, Zakarias N. The paediatric flat foot proforma (p-FFP): improved and abridged following a reproducibility study. *Journal of Foot and Ankle Research* 2009;**2**(1):1-8. [DOI: [10.1186/1757-1146-2-25](https://doi.org/10.1186/1757-1146-2-25)]

Evans 2012

Evans AM. Screening for foot problems in children: is this practice justifiable? *Journal of Foot and Ankle Research* 2012;**5**(1):18. [DOI: [10.1186/1757-1146-5-18](https://doi.org/10.1186/1757-1146-5-18)]

Evans 2015

Evans A, Menz H, Bourke J, Nikopoulos N, Ferrari J, Wilkinson M, et al. Foot care and people with intellectual disability. *International Journal of Child Health & Human Development* 2015;**8**(4):471-91.

Evans 2017

Evans AM. Mitigating clinician and community concerns about children's flatfeet, intoeing gait, knock knees or bow legs. *Journal of Paediatric Child Health* 2017;**53**(11):1050-3. [DOI: [10.1111/jpc.13761](https://doi.org/10.1111/jpc.13761)]

Evans 2021

Evans AM. Paediatric flatfeet – a 2020 guide for clinicians to identify the 'Boomerangs'. *Journal of the American Podiatric Medical Association* 2021;**5**:20-103. [DOI: [10.7547/20-103](https://doi.org/10.7547/20-103)]

Fellas 2017a

Fellas A, Hawke F, Santos, D, Coda A. Prevalence, presentation and treatment of lower limb pathologies in juvenile idiopathic arthritis: a narrative review. *Journal of Paediatric Child Health* 2017;**53**(9):836-40.

Fellas 2017b

Fellas A, Coda A, Hawke F. Physical and mechanical therapies for lower-limb problems in juvenile idiopathic arthritis: a systematic review with meta-analysis. *Journal of the American Podiatric Medical Association* 2017;**107**(5):399-412. [DOI: [10.7547/15-213](https://doi.org/10.7547/15-213)]

Garcia-Rodriguez 1999

Garcia-Rodriguez A, Martib-Jimenez F, Carnero-Varo M, Gomez-Garcia E, Gomez-Aracena J, Fernandez-Crehuet J. Flexible flat feet in children: a real problem? *Pediatrics* 1999;**103**:84-6.

Gijon-Nogueron 2019

Gijon-Nogueron G, Martinez-Nova A, Alfageme-Garcia P, Montes-Alguacil J, Evans AM. International normative data for paediatric foot posture assessment: a cross-sectional investigation. *BMJ Open* 2019;**9**(4):e023341. [DOI: [10.1136/bmjopen-2018-023341](https://doi.org/10.1136/bmjopen-2018-023341)]

GRADEpro GDT [Computer program]

McMaster University (developed by Evidence Prime) GRADEpro GDT. Hamilton (ON): McMaster University (developed by Evidence Prime), accessed 11 November 2021. Available at gradepro.org.

Gribble, 2012

Gribble PA, Hertel J, Plisky P. Using the Star Excursion Balance Test to assess dynamic postural-control deficits and outcomes in lower extremity injury: a literature and systematic review. *Journal of Athletic Training* 2012;**47**(3):339-57.

Harris 2004

Harris EJ, Vanore JV, Thomas JL, Kravitz SR, Mendelson SA, Mendicino RW, et al. Diagnosis and treatment of pediatric flatfoot. *The Journal of Foot and Ankle Surgery* 2004;**43**:341-73.

Harris 2010

Harris EJ. The natural history and pathophysiology of flexible flatfoot. *Clinic in Podiatric Medicine & Surgery* 2010;**27**:1-23.

Henry 2008

Hendry G, Gardner-Medwin J, Watt GF, Woodburn J. A survey of foot problems in juvenile idiopathic arthritis. *Musculoskeletal Care* 2008;**6**:221-32.

Higgins 2017

Higgins JPT, Altman DG, Sterne JAC, editor(s). Chapter 8: Assessing risk of bias in included studies. In: Higgins JPT, Churchill R, Chandler J, Cumpston MS, editor(s), *Cochrane Handbook for Systematic Reviews of Interventions* Version 5.2.0 (updated June 2017). Cochrane, 2017. Available from www.training.cochrane.org/handbook.

Higgins 2020b

Higgins JPT, Li T, Deeks JJ, editor(s). Chapter 6: Choosing effect measures and computing estimates of effect. In: Higgins JPT, Thomas J, Chandler J, Cumpston M, Li T, Page MJ, et al, editor(s). *Cochrane Handbook for Systematic Reviews of Interventions* Version 6.1 (updated September 2020). Cochrane, 2020. Available from training.cochrane.org/handbook/archive/v6.1.

Kerr 2015

Kerr CM, Stebbins J, Theologis T, Zavatsky B. Static postural differences between neutral and flat feet in children with and without symptoms. *Clinical Biomechanics* 2015;**30**:314-7.

Kim 2017

Kim HY, Shin HS, Ko JH, Cha YH, Ahn JH, Hwang JY. Gait analysis of symptomatic flatfoot in children: an observational study. *Clinics in Orthopedic Surgery* 2017;**9**:363-73.

Klaue 1997

Klaue K. Planovalgus and cavovarus deformity of the hind foot. *Journal of Bone and Joint Surgery. British Volume* 1997;**79**(6):892-5. [DOI: [10.1302/0301-620x.79b6.7195](https://doi.org/10.1302/0301-620x.79b6.7195)]

Klepper 2011

Klepper SE. Measures of pediatric function. *Arthritis Care & Research* 2011;**63**(S11):S371-82. [DOI: [10.1002/acr.20635](https://doi.org/10.1002/acr.20635)]

Kutlu 2018

Kutlu M, Dogan O. Test-retest reliability and validity of three different agility tests for various team sports in young male athletes. *Central European Journal of Sport Sciences and Medicine* 2018;**22**(2):33-8. [DOI: [10.18276/cej.2018.2-04](https://doi.org/10.18276/cej.2018.2-04)]

Landorf 2008

Landorf KB, Radford JA. Minimal important difference: values for the Foot Health Status questionnaire, foot function index and visual analogue scale. *The Journal of Foot and Ankle Surgery* 2008;**18**(1):15-9. [DOI: [10.1016/j.foot.2007.06.006](https://doi.org/10.1016/j.foot.2007.06.006)]

Lefebvre 2008

Lefebvre C, Manheimer E, Glanville J. Chapter 6: Searching for studies. In: Higgins JPT, Green S, editor(s). *Cochrane Handbook for Systematic Reviews of Interventions* Version 5.0.1 (updated

September 2008). The Cochrane Collaboration, 2008. Available from training.cochrane.org/handbook/archive/v5.0.1.

Martinez-Nova 2018

Martínez-Nova A, Gijón-Noguerón G, Alfageme-García P, Montes-Alguacil J, Evans AM. Foot posture development in children aged 5 to 11 years: a three-year prospective study. *Gait & Posture* 2018;**62**:280–4. [DOI: [10.1016/j.gaitpost.2018.03.032](https://doi.org/10.1016/j.gaitpost.2018.03.032)]

Mereday 1972

Mereday C, Dolan CM, Lusskin R. Evaluation of the University of California Biomechanics Laboratory shoe insert in 'flexible' pes planus. *Clinical Orthopedic Related Research* 1972;**82**:45e58.

Moher 2015

Moher D, Shamseer L, Clarke M, Ghersi D, Liberati A, Petticrew M, et al. Preferred reporting items for systematic review and meta-analysis protocols (PRISMA-P) 2015 statement. *Systematic Reviews* 2015;**4**(1):1-9. [DOI: [10.1186/2046-4053-4-1](https://doi.org/10.1186/2046-4053-4-1)]

Montalvo 2021

Montalvo S, Gonzalez MP, Dietze-Hermosa MS, Eggleston JD, Dorgo S. Common vertical jump and reactive strength index measuring devices: a validity and reliability analysis. *Journal of Strength & Conditioning Research* 2021;**35**(5):1234–43.

Moraleda 2011

Moraleda L, Mubarak SJ. Flexible flatfoot: differences in the relative alignment of each segment of the foot between symptomatic and asymptomatic patients. *Journal of Pediatric Orthopedics* 2011;**31**:421–8.

Moynihan 2012

Moynihan R, Doust J, Henry D. Preventing overdiagnosis: how to stop harming the healthy. *BMJ* 2012;**28**:344. [DOI: [10.1136/bmj.e3502](https://doi.org/10.1136/bmj.e3502)]

Moynihan 2014

Moynihan R, Henry D, Moons KG. Using evidence to combat overdiagnosis and overtreatment: evaluating treatments, tests, and disease definitions in the time of too much. *PLoS Medicine* 2014;**11**(7):e1001655.

Moynihan 2017

Pathirana T, Clark J, Moynihan R. Mapping the drivers of overdiagnosis to potential solutions. *BMJ* 2017;**16**:358. [DOI: [10.1136/bmj.j3879](https://doi.org/10.1136/bmj.j3879)]

Page 2020

Page MJ, Higgins JPT, Sterne JAC. Chapter 13: Assessing risk of bias due to missing results in a synthesis. In: Higgins JPT, Thomas J, Chandler J, Cumpston M, Li T, Page MJ, et al, editor(s). *Cochrane Handbook for Systematic Reviews of Interventions* Version 6.1 (updated September 2020). Cochrane, 2020. Available from training.cochrane.org/handbook/archive/v6.1.

Pfeiffer 2006

Pfeiffer M, Kotz R, Ledl T, Hauser G, Sluga M. Prevalence of flatfoot in preschool-aged children. *Pediatrics* 2006;**118**:634–9.

Rigoldi 2012

Rigoldi C, Galli M, Cimolin V, Camerota F, Celletti C, Tenore N, et al. Gait strategy in patients with Ehlers-Danlos syndrome hypermobility type and Down syndrome. *Research in Developmental Disabilities* 2012;**33**(5):1437–42. [DOI: [10.1016/j.ridd.2012.03.016](https://doi.org/10.1016/j.ridd.2012.03.016)]

Schünemann 2020a

Schünemann HJ, Higgins JPT, Vist GE, Glasziou P, Akl EA, Skoetz N, et al. Chapter 14: Completing 'Summary of findings' tables and grading the certainty of the evidence. In: Higgins JPT, Thomas J, Chandler J, Cumpston M, Li T, Page MJ, et al, editor(s). *Cochrane Handbook for Systematic Reviews of Interventions* Version 6.1 (updated September 2020). Cochrane, 2020. Available from training.cochrane.org/handbook/archive/v6.1.

Schunemann 2020b

Schünemann HJ, Vist GE, Higgins JPT, Santesso N, Deeks JJ, Glasziou P, et al. Chapter 15: Interpreting results and drawing conclusions. In: Higgins JPT, Thomas J, Chandler J, Cumpston M, Li T, Page MJ, et al, editor(s). *Cochrane Handbook for Systematic Reviews of Interventions* Version 6.1 (updated September 2020). Cochrane, 2020. Available from training.cochrane.org/handbook/archive/v6.1.

Staheli 1987

Staheli LT, Chew DE, Corbett M. The longitudinal arch: a survey of 802 feet in normal children and adults. *Journal of Bone and Joint Surgery. American Volume* 1987;**69**:426–8.

Varni 1996

Varni JW, Waldron SA, Gragg RA, Rapoff MA, Bernstein BH, Lindsley CB, et al. Development of the Waldron/Varni pediatric pain coping inventory. *Pain* 1996;**67**(1):141–50.

Varni 2002

Varni JW, Seid M, Knight TS, Szer JS. The PedsQL in pediatric rheumatology: reliability, validity, and responsiveness of the Pediatric Quality of Life Inventory, generic core scales and rheumatology module. *Arthritis and Rheumatism* 2002;**46**:714–25.

Weber 2014

Weber A, Martin K. Efficacy of orthoses for children with hypotonia. *Pediatric Physical Therapy* 2014;**26**:38–47.

Wegener 2011

Wegener C, Hunt AE, Vanwanseele B, Burns J, Smith RM. Effect of children's shoes on gait: a systematic review and meta-analysis. *Journal of Foot and Ankle Research* 2011;**4**(1):3. [DOI: [10.1186/1757-1146-4-3](https://doi.org/10.1186/1757-1146-4-3)]

Wilson 1968

Wilson JMG, Jungner G, World Health Organization. *Principles and Practice of Screening for Disease*; Public Health Papers No. 34. France: World Health Organization, 1968.

Yan 2013

Yan G, Yang Z, Lu M, Zhang J, Zhu Z, Guo Y. Relationship between symptoms and weight-bearing radiographic parameters of

idiopathic flexible flatfoot in children. *Chinese Medical Journal* 2013;**126**(11):2029-33.

Yoosefinejad 2014

Yoosefinejad AK, Ghalamghash R. The evaluation and prevalence of foot problems among Iranian students using 'alfoots' company scanner. *Health Science Journal* 2014;**8**(3):393-9.

Systematic Reviews 2010, Issue 7. Art. No: CD006311. [DOI: [10.1002/14651858.CD006311.pub2](https://doi.org/10.1002/14651858.CD006311.pub2)]

Rome 2010

Rome K, Evans A, Ashford R. Non-surgical interventions for paediatric pes planus. *Cochrane Database of Systematic Reviews* 2010, Issue 7. Art. No: CD006311. [DOI: [10.1002/14651858.CD006311.pub2](https://doi.org/10.1002/14651858.CD006311.pub2)]

References to other published versions of this review

Evans 2010

Evans AM, Rome K, Carroll M, Hawke F. Foot orthoses for treating paediatric flat feet. *Cochrane Database of*

* Indicates the major publication for the study

CHARACTERISTICS OF STUDIES

Characteristics of included studies [ordered by study ID]

Abd-Elmonem 2021

Study characteristics

Methods	Randomised controlled two-arm parallel trial
Participants	Initially recruited N = 134, excluded = 62 Study included: 72 children with flat feet, 36 in each group Age: 7 to 12 years old male and female Gender: 31 M, 35 F (M/F - 15/18 in intervention group; 16/17 in sham group) Setting: out-patient Physical Therapy Clinic of Faculty of Physical Therapy, Cairo University, Cairo, Egypt Inclusion criteria: (i) diagnosed (by an orthopedist) with asymptomatic flexible flatfoot, (ii) navicular drop more than 9 mm, (iii) grade III flatfoot Staheli Arch Index (midfoot width exceeding forefoot width) Exclusion criteria: (i) congenital deformities of the lower extremities (e.g. genu valgum, femoral anteversion), (ii) scar or osseous anomalies Baseline characteristics: age, mean (SD): 9.5 (1.02) in intervention group; 9.45 (0.76) in sham group
Interventions	Group 1: neuromuscular electrical stimulation and corrective exercise - 4 months (3 sessions/week) Group 2: sham neuromuscular electrical stimulation and corrective exercise - 4 months (3 sessions/week)
Outcomes	Assessments of Staheli's arch index (through foot print), navicular drop (through navicular drop test), and radiographic indexes (through anterior-posterior and medio-lateral X-ray) of both feet were performed before and after the intervention programmes
Source of funding	This research received no specific grant from any funding agency in the public, commercial, or not-for-profit sectors
Notes	Trial registration: NCT04410926

Risk of bias

Bias	Authors' judgement	Support for judgement
------	--------------------	-----------------------

Abd-Elmonem 2021 (Continued)

Random sequence generation (selection bias)	Low risk	"The online Graph Pad software was used to allocate study participants".
Allocation concealment (selection bias)	Low risk	"The randomisation was carried out by an independent person, who was unaware of the study protocol and not otherwise in control of the study".
Blinding of participants and personnel (performance bias) All outcomes	Unclear risk	"All children/legal guardians, radiologist and researchers responsible for evaluations were blinded to participants' allocation". However it is unclear whether the blinding was successful, as the trialists did not discuss the possibility that participants might have guessed their intervention based on whether they felt any discomfort related to the intervention or not.
Blinding of outcome assessment (detection bias) - self-reported outcomes (e.g., pain, function)	Low risk	There were no self-reported outcomes
Blinding of outcome assessment (detection bias) - objective outcomes	Low risk	Outcome assessors were blinded.
Incomplete outcome data (attrition bias) All outcomes	Low risk	3/36 (discontinued intervention = 3) in the intervention, and 3/36 in control group (discontinued intervention = 2; femoral fracture = 1) withdrew from the study and not included in the analysis
Selective reporting (reporting bias)	Low risk	trial registered, results for all pre-specified outcomes reported
Other bias	Low risk	none apparent

Aboutorabi 2013
Study characteristics

Methods	Randomised controlled three-arm parallel trial
Participants	<p>Initially recruited N = 62, 12 excluded</p> <p>Study included: 30 children with flat feet, 20 healthy controls</p> <p>Age: average 7.76 (1.4) years</p> <p>Gender: 30 M, 20 F (flat foot – 18 M, 12 F; healthy – 12 M, 8 F)</p> <p>Location - Hehran, Iran</p> <p>Setting: Iran University of Medical Sciences, Dept Orthopedics and Prosthetics, Faculty of Rehabilitation Sciences</p> <p>Inclusion criteria: flat foot and healthy children; flatfoot FPI > +6</p> <p>Exclusion criteria: contracture of soft tissue, genu varum or valgum, structural congenital deformity, lower extremity fracture, neurological problems, foot or ankle surgery in past 6 months, leg length discrepancy</p>

Aboutorabi 2013 (Continued)

Baseline characteristics: trial group/flat feet N = 30, 18 M:12 F, control group/non-flat feet N = 20, 12 M:8 F

Interventions	<ol style="list-style-type: none"> 1. functional foot orthoses and regular shoe 2. medical shoes 3. barefoot
Outcomes	<p>Outcomes were measured as immediate effects at baseline only.</p> <p>Gait parameters for the 3 interventions, i.e.</p> <ol style="list-style-type: none"> 1. step length 2. step width 3. walking velocity 4. symmetry 5. centre of pressure (CoP) displacement
Source of funding	none
Notes	<p>- FPI (Foot Posture Index) of healthy group is not defined, versus the flatfoot group FPI +6 to +12. Hence, basis for comparison from baseline unclear</p> <p>- Immediate effects only, hence, we did not include this trial in further analyses</p>

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	<p>Insufficient information about the sequence generation process to permit judgement of low risk or high risk.</p> <p>‘A randomised method was used for the sampling from the flat footed children who were referred for a new pair of shoes.’</p> <p>‘The test order was randomised for each child to minimize learning effect.’</p>
Allocation concealment (selection bias)	Unclear risk	Insufficient information to permit judgement of low risk or high risk
Blinding of participants and personnel (performance bias) All outcomes	High risk	<p>Each child participates in barefoot, regular shoe, and medical shoe trials. No blinding was performed.</p> <p>Research staff prepared each child for barefoot, regular shoe, and medical shoe trials. No blinding was performed.</p>
Blinding of outcome assessment (detection bias) - self-reported outcomes (e.g., pain, function)	High risk	N/A
Blinding of outcome assessment (detection bias) - objective outcomes	Unclear risk	Outcomes included assessment of gait parameters. Both the participant and the researcher were considered to be outcome assessors as both could influence the result. The participants were not blinded. No mention was made of blinding of researchers involved in outcome assessment.

Aboutorabi 2013 (Continued)

Incomplete outcome data (attrition bias) All outcomes	Unclear risk	Insufficient information to permit judgement of low risk or hHigh risk.
Selective reporting (reporting bias)	Unclear risk	No reference to study protocol. Insufficient information to permit judgement of low risk or high risk.
Other bias	Unclear risk	5 min rest was allowed between testing conditions for each child. It was not clear if this was sufficient to eliminate a carry-over effect. The random order of testing minimised this effect.

Ahn 2017
Study characteristics

Methods	Randomised controlled two-arm parallel trial
Participants	40 children with flexible flat feet Age: average 10 (4.5) years Gender: 24 M, 16 F Location: Daejeon, Korea Setting: Department of Pediatric Rehabilitation at Chung Nam National University Hospital Inclusion criteria: flexible flatfoot defined as > 4° valgus RCSP, one abnormal x-ray finding i.e. > 30° AP talo-calcaneal angle, > 45° lateral talo-calcaneal angles, > 4° lateral talo-metatarsal angles, < 10° calcaneal pitch Exclusion criteria: rigid flatfoot, of any cause Baseline characteristics: matched except for age - RFO 10.14 (4.99) years, TCFO 9.59 (4.24) years
Interventions	1. rigid foot orthoses (N = 20) 2. talus control foot orthoses (N = 20)
Outcomes	- RCSP - x-rays, 4 angles: AP talo-calcaneal angle, lateral talo-calcaneal angles, lateral talo-metatarsal angles, calcaneal pitch Both measures were assessed at baseline and after 12 months
Source of funding	none reported
Notes	no mention of pain, or gait parameters

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Insufficient information about the sequence generation process to permit judgement of low risk or high risk.

Ahn 2017 (Continued)

'Forty children were randomly prescribed a foot orthosis: 20 a RFO and 20 a TCFO'

Allocation concealment (selection bias)	Unclear risk	Insufficient information to permit judgement of low risk or high risk.
Blinding of participants and personnel (performance bias) All outcomes	High risk	Participants were not blinded Personnel were not blinded
Blinding of outcome assessment (detection bias) - self-reported outcomes (e.g., pain, function)	High risk	N/A
Blinding of outcome assessment (detection bias) - objective outcomes	Unclear risk	It was not clear if the person who measured resting calcaneal stance position or radiographs was blinded.
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	The number of children who completed the study, or for whom outcome data were presented was not reported.
Selective reporting (reporting bias)	Unclear risk	No reference to study protocol. Insufficient information to permit judgement of low risk or high risk.
Other bias	Unclear risk	Funding source for orthoses was not stated.

Asgaonkar 2012
Study characteristics

Methods	Randomised controlled two-arm parallel trial
Participants	80 children with flat feet, aged 5 to 15 years, in Mumbai, India 1. treatment group N = 45 2. age-sex matched control group N = 35 Age: 5 to 15 years Gender: not stated, but groups 'age and sex matched' (p 91) Location: primary and secondary schools Setting: Mumbai, India Inclusion criteria: flat feet - from footprint measures (Bleck arch index > 1.15) Exclusion criteria: fixed foot deformity, surgeries for foot deformities, pain injury that had required non-weight bearing, systemic problems Baseline characteristics: matched, but uneven group sizes
Interventions	1. valgus insole, rubber material of average thickness 4 cm (N = 45)

Asgaonkar 2012 (Continued)

2. no treatment (N = 35)

Outcomes	Outcome measures at baseline and 12 months 1. Pain - VAS 2. PCI (physical cost index) - average HR and speed for 100 m walk 3. Gait parameters - step length, stride length, cadence, walking velocity
Source of funding	none reported
Notes	- 20/80 dropouts in 12-month trial, most from treatment group - pain used VAS – some children were only 5 years old - valgus insole vague; unsure re: footwear used in either group (presume 'usual', but not stated)

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Insufficient information about the sequence generation process to permit judgement of low risk or high risk. "Out of the total 80 students, age, and sex, matched two groups were made at random; control group consisting of 35 students and experimental group consisting of 45 students."
Allocation concealment (selection bias)	Unclear risk	Insufficient information to permit judgement of low risk or high risk.
Blinding of participants and personnel (performance bias) All outcomes	High risk	No blinding, and the outcome was likely to be influenced by lack of blinding.
Blinding of outcome assessment (detection bias) - self-reported outcomes (e.g., pain, function)	High risk	No mention was made of blinding of researchers involved in outcome assessment.
Blinding of outcome assessment (detection bias) - objective outcomes	High risk	No mention was made of blinding of researchers involved in outcome assessment.
Incomplete outcome data (attrition bias) All outcomes	High risk	Imbalance in number of dropouts per group. n = 45 in the experimental group, 9 participants withdrew from the study and an additional 6 participants left the school. n = 35 in the control group, it appears 5 participants left the school and none withdrew from the study. Reasons for dropouts from the experimental group were not provided, but are likely related to the intervention.
Selective reporting (reporting bias)	Unclear risk	No reference to study protocol. Insufficient information to permit judgement of low risk or high risk.

Asgaonkar 2012 (Continued)

Other bias	Unclear risk	No funding source reported
------------	--------------	----------------------------

Bok 2016
Study characteristics

Methods	Randomised controlled four-arm parallel trial
Participants	21 children with symptomatic flexible flat feet Age: aged 8 to 13 years Gender: 8 M, 13 F Location: Jung-gu, Daejeon, South Korea Setting: Department of Rehabilitation Medicine, Chungnam National University Hospital Inclusion criteria: severe flat feet with pain, RCSP > 6° everted Exclusion criteria: (i) a fixed foot deformity (ii) reported previous intervention (e.g. orthoses or surgery) (iii) congenital and developmental foot disease (iv) neuromuscular or central nervous system disease Children with a history of overuse or traumatic injury to the lower limb in the past 6 months, bony surgery to the lower limb, or systemic endocrine, neurogenic, or musculoskeletal disorders were also excluded. Baseline characteristics: 8 M:13 F
Interventions	All children trialled 4 shoes with different degrees of orthotic (N = 21): 1. shoes 2. shoes and 0° orthotic 3. shoes and 15° orthotic 4. shoes and 30° orthotic
Outcomes	Plantar pressures - peak pressure, maximum force, contact area Reported shoe and orthotic condition, as immediate effects on plantar pressures
Source of funding	orthoses supplied by Korean company
Notes	- did not include any pain evaluation (yet all participants were symptomatic at baseline) - immediate effects only, hence, we did not include this trial in further analyses

Risk of bias

Bias	Authors' judgement	Support for judgement
------	--------------------	-----------------------

Bok 2016 (Continued)

Random sequence generation (selection bias)	Unclear risk	‘The 4 orthotic conditions were tested in a random order to minimize potential sequencing effects.’ Insufficient information about the sequence generation process to permit judgement of low risk or high risk.
Allocation concealment (selection bias)	Unclear risk	Insufficient information to permit judgement of low risk or high risk.
Blinding of participants and personnel (performance bias) All outcomes	Unclear risk	Each child participated in each orthotic condition. No blinding was performed. Research staff dispensed the orthoses to the children. It was unclear how involved the research staff were in preparing each child for each trial, for example, fitting the orthoses in the shoes.
Blinding of outcome assessment (detection bias) - self-reported outcomes (e.g., pain, function)	Unclear risk	Immediate effects only
Blinding of outcome assessment (detection bias) - objective outcomes	Unclear risk	Outcomes included assessment of gait parameters. Both the participant and the researcher were considered to be outcome assessors, as both could influence the result. The participants were not blinded. No mention was made of blinding of researchers involved in outcome assessment.
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	Insufficient reporting of attrition/exclusions to permit judgement of low risk or high risk.
Selective reporting (reporting bias)	Unclear risk	No reference to study protocol. Insufficient information to permit judgement of low risk or high risk.
Other bias	Unclear risk	It was not clear if participants were allowed an adjustment period for each orthotic design. The more aggressively posted orthoses may require a longer period to adjust to normalise gait.

Coda 2014
Study characteristics

Methods	Randomised controlled two-arm parallel trial, single blinded, multi-centre
Participants	60 children with JIA Ages - 11.17(3.51) control PFOs, 10.64(3.84) fitted PFOs Genders: control PFOs 6M/23F; fitted PFOs 9M/22F Setting: Paediatric Rheumatology hospital clinics Location: Scotland, UK Inclusion criteria: diagnosed with juvenile idiopathic arthritis (JIA; any subtype) according to International League of Associations for Rheumatology criteria, lower extremity joint involvement with disease onset ranging from 5 to 18 years, previous failure of orthotic management in which the child has not worn any foot orthoses for at least 3 months, ability to walk a minimum of 15 metres or more without assistive devices, at least 6 months after start of disease modifying antirheumatic drug therapy

Coda 2014 (Continued)

Exclusion criteria: inability to walk barefoot or shod, concomitant musculoskeletal disease, central or peripheral nerve disease and endocrine disorders, previous foot surgery, currently using foot orthoses, supply of foot orthoses is contraindicated

Baseline characteristics: CPFO group had higher VAS, 14 versus 6.5 in PFO group (NS); CPFO group had 65% oligoarthritis versus PFO 44.8%

Interventions	1. Fitted PFOs, customised in clinic (N = 31) 2. Control PFOs (N = 29) Both PFO types had same black covers to assist blinding
Outcomes	Baseline, 3 and 6 month outcomes - VAS - PedsQL
Source of funding	Queen Margaret University, Edinburgh - PhD scholarship
Notes	Foot posture was not defined Validated outcome measures used

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	"After obtaining informed consent, children were randomised in blocks of 10 each by an online computer random number generator (www.randomization.com)."
Allocation concealment (selection bias)	Unclear risk	No specific mention of allocation concealment. Insufficient information to permit judgement of high or low risk
Blinding of participants and personnel (performance bias) All outcomes	High risk	low - 'The control FOs was made with leather board (1 mm) without corrections. Both FOs had the same black-ethylene vinyl acetate (EVA) top cover (0.75 mm) to allow for blinding and monitoring the level of adherence to wearing the FOs.' high - study did not blind investigators to the intervention, and the outcome may be influenced by lack of blinding
Blinding of outcome assessment (detection bias) - self-reported outcomes (e.g., pain, function)	Low risk	Pain measured by participants HRQoL measured by participants and parents/carers Participants and their parents/carers are blinded
Blinding of outcome assessment (detection bias) - objective outcomes	Unclear risk	N/A
Incomplete outcome data (attrition bias) All outcomes	Low risk	'Overall, 179 out of a possible 180 assessments were completed (99.4%) and accounted for statistical analysis.' No reason for missing data given and it was not clear from which group and time point the data were missing.

Coda 2014 (Continued)

Selective reporting (reporting bias)	Unclear risk	No reference to study protocol. Insufficient information to permit judgement of low risk or high risk.
Other bias	Unclear risk	It is not clear how the orthoses in this study were funded.

Gould 1989

Study characteristics

Methods	Randomised controlled four-arm parallel trial
Participants	125 normal toddlers (beginner walkers) aged 11 to 14 months Age: 11 to 14 months Gender: not stated, except "boys and girls" Table 2 , p 243 – genders/groups at 5 years (27 M, 25 F) Location: University of Vermont, USA Setting: pediatrics dept Inclusion criteria: "clinically normal children... aged 11 to 14 months" Exclusion criteria: none stated Baseline characteristics: sex ratio at baseline not stated
Interventions	Group 1, N = 50 – straight last shoes Group 2, N = 25 – Group 1 shoes + long arch cookies Group 3, N = 15 – orthopaedic shoes with long counters, solid shanks, Thomas heels, 0.3 cm inside heel wedges Group 4, N = 25 – Group 3 shoes + thin longitudinal arch support
Outcomes	Baseline, 2,3, and 5 years - x-ray angles (AP talo-1st metatarsal, lateral talo-1st metatarsal, lateral talo-calcaneal) - biometry (pedoscope - arch appearance) - clinical examination - use of shoes, examination of femoral, tibial, knee configuration - "will be reported in detail subsequently"
Source of funding	1. Annual grant from Footwear Association 2. Shoes provided by Sabel shoes
Notes	Group 1, N = 50 acting as shod control group, across 4 years of the trial High attrition - 52/125 finished the study

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	'divided by lot into four difference footwear groups'

Gould 1989 (Continued)

		Insufficient information about the sequence generation process to permit judgement of low risk or high risk.
Allocation concealment (selection bias)	Unclear risk	Insufficient information to permit judgement of low risk or high risk
Blinding of participants and personnel (performance bias) All outcomes	High risk	Participants were not blinded Personnel were not blinded
Blinding of outcome assessment (detection bias) - self-reported outcomes (e.g., pain, function)	Unclear risk	N/A
Blinding of outcome assessment (detection bias) - objective outcomes	Unclear risk	'A determination was made as to whether the arch was slightly or moderately improved. The records were then consulted to determine what type of footwear was worn in each case.' It was not clear who made this determination, and if they had opportunity prior to looking at the records to know what footwear was worn in each case.
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	42% of children completed the trial. Attrition was uneven between groups. Reasons for attrition were not sufficiently reported to permit judgement of low risk or high risk.
Selective reporting (reporting bias)	Unclear risk	No reference to study protocol. Insufficient information to permit judgement of low risk or high risk.
Other bias	Unclear risk	Subjective and non-validated outcome assessments. 'A determination was made as to whether the arch was slightly or moderately improved'

Hsieh 2018
Study characteristics

Methods	Randomised controlled two-arm parallel trial
Participants	52 healthy children with symptomatic flexible flatfoot Age: all approximated 6 years - treatment group 6.9 (0.6) years, control group 6.2 (0.4) years Gender: (28 M, 24 F) Location: Shin Kong Wu Ho-Su Memorial Hospital, Taiwan Setting: Dept Physical Medicine and Rehabilitation Inclusion criteria: symptomatic flexible flatfoot (pain over the foot or calf, fatigue after prolonged walking, and gait disturbances) Exclusion criteria: history of foot injury or surgery, foot abnormalities affecting locomotion or foot mobility, or a confirmed diagnosis of developmental delays, such as developmental co-ordination disorder and neurological deficits

Hsieh 2018 (Continued)

	Baseline characteristics: treatment group had significantly lower scores for PODCI (transfer and mobility), PedsQL (physical, psychological, total health)
Interventions	Treatment group (N = 26): thermoplastic insoles, heat moulded to child's feet (CPFOs) Control group (N = 26): no treatment
Outcomes	Physical activity: timed 10-m walk, stair ascent, timed up and go, chair rise Physical function: parent reported PODCI (Chinese version) Psychometric: PODCI, PedsQL (parent report)
Source of funding	Multiple research grants from Shin Kong Wu Ho-Su Memorial Hospital, and Ministry of Science and Technology, Taiwan
Notes	Trial ran for 12 weeks Flatfoot assessment: ND test, FPI-6, lat and AP x-rays, Beighton score Self-selected usual footwear Insole use suggested to be 5 hours/day

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	The participants were randomly assigned to the treatment group (with customized insoles) or the control group (without customized insoles) according to computer-generated random numbers (Fig. 1).
Allocation concealment (selection bias)	Low risk	Allocation was initially concealed. A sealed envelope was opened for each consecutive participant to reveal the participant's group allocation when the participant was recruited to the study. One physician enrolled all participants, and another investigator generated the allocation sequence and assigned the participants to their groups.
Blinding of participants and personnel (performance bias) All outcomes	High risk	Control group had no orthotic intervention
Blinding of outcome assessment (detection bias) - self-reported outcomes (e.g., pain, function)	High risk	Due to nature of intervention
Blinding of outcome assessment (detection bias) - objective outcomes	High risk	High risk for primary outcome of pain as participants were not blinded. Secondary outcomes that were assessed by a blinded assessor had low risk of bias.
Incomplete outcome data (attrition bias) All outcomes	Low risk	Numbers and reasons provided for missing outcome data. Number was small and reason provided unlikely to be due to the intervention.
Selective reporting (reporting bias)	High risk	ClinicalTrials.gov (NCT02414087) The published study included more outcomes than are listed in the trial registry.

Hsieh 2018 (Continued)

Other bias	Low risk	none apparent
------------	----------	---------------

Jafarnezhadgero 2018
Study characteristics

Methods	Randomised controlled two-arm parallel trial
Participants	<p>30 healthy boys, with flexible flat feet</p> <p>Age: 8 to 12 years</p> <p>Gender: all boys</p> <p>Location: not clear - boys 'recruited from orthopaedic specialists in local community' (p 4)</p> <p>Setting: not clear, ethics approval and trial registry within Iran (p 4)</p> <p>Inclusion criteria: boys who volunteered, flat feet assessed using navicular drop, arch height index, RCSP</p> <p>Exclusion criteria: history of bone fractures, signs of functional lower limb instability, ligament injury, reconstruction of ligaments, neuromuscular dysfunction, dysfunction of lower limb muscles, leg length differences larger than 1 cm, and a history of lower extremity trauma or surgery.</p> <p>Baseline characteristics: all were right foot dominant, no significant baseline differences between groups for examined variables</p>
Interventions	<p>Treatment group (N = 15): medial arch support foot orthoses (custom made medial arch support foot orthoses), CFOs</p> <p>Control group (N = 15): flat 2-mm thick insoles (sham)</p>
Outcomes	Gait kinematics and ground reaction forces (kinetic) - joint vectors, moments, and ground reaction forces
Source of funding	Deutsche Forschungsgemeinschaft (DFG), and Open Access Publishing Fund of University of Potsdam, Germany.
Notes	All participants issued with same footwear (New Balance 759)

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Says only 'The block randomization method was used to allocate study participants into experimental groups.'
Allocation concealment (selection bias)	Unclear risk	<p>Says only; 'Another naïve examiner controlled the allocation of each participant and was responsible for delivering the treatment to both groups.'</p> <p>Not enough information to determine risk.</p>
Blinding of participants and personnel (performance bias) All outcomes	Low risk	Sham orthoses used

Jafarnezhadgero 2018 (Continued)

Blinding of outcome assessment (detection bias) - self-reported outcomes (e.g., pain, function)	High risk	Due to nature of intervention
Blinding of outcome assessment (detection bias) - objective outcomes	Low risk	Examiners blinded for all outcomes
Incomplete outcome data (attrition bias) All outcomes	Low risk	No missing data
Selective reporting (reporting bias)	Unclear risk	Iranian Registry of Clinical Trials (IRCT2017082235517N1 ; URL: www.irct.ir/user/trial/26811/view) - unable to access the online trial registry.
Other bias	Low risk	none apparent

Kanatli 2016
Study characteristics

Methods	Randomised controlled two-arm parallel trial
Participants	<p>45 children, gender (33 boys, 12 girls) - with moderate flexible flatfoot</p> <p>Age: mean age 39.5 months (17 to 72 months)</p> <p>Location: Ankara, Turkey</p> <p>Setting: Gazi University School of Medicine</p> <p>Inclusion criteria: moderate flexible flatfoot (Volpe classification - Jack's test, standing heel position)</p> <p>Exclusion criteria: rigid flatfoot, neuromuscular disorders, genetic disease associated with collagen abnormalities</p> <p>Baseline characteristics: sample sex ratio 33 M: 12 F, not stated/group</p>
Interventions	<p>1. orthopaedic shoes, N = 21</p> <p>2. control group, N = 24</p>
Outcomes	<p>Baseline and at follow up, i.e. 34.6 (10.9) months (range 24 to 57 months)</p> <p>- Joint laxity (Wynne-Davis method)</p> <p>- Arch Index (Staheli method)</p> <p>- x-ray angles (lateral and AP talo-1st metatarsal, lateral talo-horizontal, lateral calcaneal pitch, lateral and AP talo-calcaneal)</p>
Source of funding	not stated
Notes	Orthopaedic shoes had internal 'orthosis' and Thomas heels

Risk of bias
Foot orthoses for treating paediatric flat feet (Review)

Kanatli 2016 (Continued)

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	High risk	Systematic, non-random approach. 'Patients were randomised and separated in two groups. Randomization was made by weekly basis. Patients that fulfilled the inclusion criteria during even number weeks consisted group 1 and in odd number weeks consisted group 2.'
Allocation concealment (selection bias)	High risk	Inexplicably unconcealed procedure: allocated depending on week odd or even week at enrolment.
Blinding of participants and personnel (performance bias) All outcomes	High risk	Children were not blinded. Children received orthopaedic shoes or no intervention. Personnel fitted the orthopaedic shoes or no intervention.
Blinding of outcome assessment (detection bias) - self-reported outcomes (e.g., pain, function)	Unclear risk	N/A
Blinding of outcome assessment (detection bias) - objective outcomes	Unclear risk	It is not clear who assessed the outcomes and if this person was aware of group allocation.
Incomplete outcome data (attrition bias) All outcomes	Low risk	All enrolled children completed the study.
Selective reporting (reporting bias)	High risk	Arch index scores were not reported for any time point.
Other bias	Unclear risk	It was not clear how the orthopaedic shoes in this study were funded.

Khamooshi 2017
Study characteristics

Methods	Randomised controlled three-arm parallel trial
Participants	60/130 girls with flat feet Age: 9 to13 years Gender: all girls Location: Tehran, Iran Setting: Schools of Khalil Abad county Inclusion criteria: female, good general health, flexible flat feet – assessed using tip toe test, navicular collapse Exclusion criteria: lower extremity surgery, fracture, orthopaedic problems Baseline characteristics: all girls, matched re age, BMI
Interventions	1. control group, N = 20

Khamooshi 2017 (Continued)

2. stretches, strengthening; i.e. Achilles tendon stretching and strengthening, exercises effective on the interior longitudinal foot arch, N = 20

3. stretches, strengthening, core exercises; i.e. exercises related to the muscles, Achilles tendon stretching and strengthening, exercises effective on the interior longitudinal foot arch and activities for core stability, N = 20

Baseline, and after exercise programme: groups 2 and 3 performed the exercises for 8 weeks, three times a week, in the form of three turns with 20 repetitions.

Outcomes	Pre/post exercise programme (8 weeks apart) - Staheli footprint arch index: the narrowest section of the arch (A) was divided by the broadest section of the foot (B) based on Staheli's formula ($AI = A/B$) - Navicular collapse test: seated with knee flexed 90 degrees, and subtalar neutral position. The navicular height from ground measured (mm). This is repeated with participant standing, and the difference recorded as the navicular collapse rate.
Source of funding	Not stated
Notes	Only Staheli index reported, and yet Navicular collapse is stated to improve with exercises (both groups), versus control (p 154)

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Insufficient information about the sequence generation process to permit judgement of low risk or high risk. 'In the present study, 60 female students with flat feet disorders were selected, who were stochastically assigned to three 20-individual groups'
Allocation concealment (selection bias)	Unclear risk	Insufficient information to permit judgement of low risk or high risk.
Blinding of participants and personnel (performance bias) All outcomes	Unclear risk	Participants were not blinded Personnel were not blinded
Blinding of outcome assessment (detection bias) - self-reported outcomes (e.g., pain, function)	Unclear risk	N/A
Blinding of outcome assessment (detection bias) - objective outcomes	Unclear risk	It was not clear if the person who measured Staheli arch index was aware of group allocation.
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	The number of children who completed the study, or for whom outcome data were presented was not reported.
Selective reporting (reporting bias)	Unclear risk	Data for 'navicular bone collapse' (also referred to as Navi Loss within the paper) was not reported.
Other bias	Unclear risk	The training programmes were inadequately described.

Morrison 2013

Study characteristics

Methods	Randomised controlled two-arm parallel trial
Participants	<p>22 children were approached to participate in the study. All were male, white, British, and FPI > +4</p> <p>20 children consented to participate, then 5 failed to attend. 1 participant withdrew</p> <p>14 participants (median age 7.5 years) completed the study</p> <p>Age: aged 6 to 11 years</p> <p>Gender; all white British boys</p> <p>Location: London, UK</p> <p>Setting: children's physical rehabilitation programme (Children's Therapy Service at Medway Community Healthcare)</p> <p>Inclusion criteria: diagnosis DCD, flat feet defined by FPI > +4</p> <p>Exclusion criteria: medical complications likely to affect gait – neuromuscular, orthopaedic conditions</p> <p>Baseline characteristics: all boys, stated that groups were similar for age, foot posture, and hypermobility (but no demographic data)</p>
Interventions	<p>- CFOs (N = 9)</p> <p>- no CFOs (N = 5)</p> <p>Both groups completed a 7-week rehabilitation programme</p>
Outcomes	<p>- Spatio-temporal gait (Gait-rite system)</p> <p>- Six-minute walk test</p> <p>No significant differences from baseline to 7 week completion.</p>
Source of funding	Canonbury Healthcare funding
Notes	<p>High attrition - 14/21 (68%) finished the trial</p> <p>Small sample</p> <p>Specific cohort</p> <p>Preliminary trend inferred from NS results</p> <p>No true control group – quasi RCT</p>

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	All participants entering the study were quasi-randomised into one of the two groups..? Method of randomisation unclear
Allocation concealment (selection bias)	Low risk	'...using a sealed envelope technique. Each envelope was opened after the child consented to taking part in the programme'

Morrison 2013 (Continued)

Blinding of participants and personnel (performance bias) All outcomes	High risk	Participants aware of foot orthoses being supplied at the start of end of rehabilitation program. Personnel supplied the orthoses at the start or end of the rehabilitation programme.
Blinding of outcome assessment (detection bias) - self-reported outcomes (e.g., pain, function)	Unclear risk	N/A
Blinding of outcome assessment (detection bias) - objective outcomes	High risk	The investigator was blind to the treatment group during the initial data collection. It was not clear if investigators were blind at the time of outcome assessment at follow-up. The gait parameters reported are influenced by the participants who were not blinded.
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	Reasons not provided for attrition.
Selective reporting (reporting bias)	Unclear risk	No reference to study protocol. Insufficient information to permit judgement of low risk or high risk
Other bias	Low risk	Source of funding declared and independence of funder from research design, conduct and reporting assured.

Powell 2005
Study characteristics

Methods	Randomised controlled three-arm parallel trial
Participants	48 children with JIA, and persistent lower limb pain were enrolled (sample size calculation not stated) Age: 5 to 19 years (mean 12 years, 7 months, SD 3.7) Gender: 30 F, 10 M Setting; outpatients, Southern Californian children's hospitals, Location: San Diego, USA Inclusion criteria: diagnosed juvenile chronic arthritis; over 5 years old, active foot disease, history of foot pain over 1 month, able to walk 50 metres, stable medications Exclusion criteria: previous use of shoe inserts, joint injections in last 6 months, osseous anomaly Baseline characteristics: fewer males in CFO group (13.3% versus 33.3% in PFO, 30.8% in shoes); more polyarthritis in CFO group (73.3% versus 50% in PFOs, and 30.8% in shoes); CFO group had more pain on VAS (5.23) versus PFO (VAS 3.5), versus shoe (VAS 4.7)
Interventions	1: custom-made semi-rigid foot orthoses (CFO) 2: prefabricated foot orthoses (PFO) 3: new athletic footwear with a medial longitudinal arch and shock-absorbing sole
Outcomes	Outcomes measured at baseline and 3 months

Foot orthoses for treating paediatric flat feet (Review)

Powell 2005 (Continued)

1. Paediatric Pain questionnaire: 0 to 10-point VAS scale (0 = no pain)
2. Timed walking speed
3. Foot Function index: 3 subscales - activity limitation, foot pain, disability, 0 to 100-point scale (0 = best function)
4. Physical functioning subscale of PedsQL, 0 to 100-point scale (0 = worst function)

Source of funding	Foot orthoses supplied by Langer Biomechanics, Neoprene insoles supplied by Spenco Medical
Notes	Pain reported in the trial using two instruments (Paediatric pain questionnaire and foot pain subscale of the Foot Function index) For this review, we extracted data using the Paediatric pain questionnaire for the analysis of pain

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Method of random sequence generation unclear. 'Once accepted into the study, each subject was randomly placed...' Insufficient information about the sequence generation process to permit judgement of low risk or high risk.
Allocation concealment (selection bias)	Low risk	'Sealed envelope containing a predetermined numbered placement card into one of the 3 intervention groups'
Blinding of participants and personnel (performance bias) All outcomes	High risk	No blinding and the outcome may be influenced by lack of blinding Physical therapist 1 who administered the interventions was not blinded Physical therapist 2 who performed baseline and follow-up measures was blinded
Blinding of outcome assessment (detection bias) - self-reported outcomes (e.g., pain, function)	High risk	Pain, level of disability, and activity limitation measured by participants
Blinding of outcome assessment (detection bias) - objective outcomes	High risk	Speed of ambulation influenced by participant and measured by personnel The outcome measurement may be influenced by lack of blinding
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	Withdrawals few and similar between groups No child withdrew from the study because of discomfort or lack of efficacy. Children who withdrew were not different from children who completed the study in terms of parental education level, family income, race/ethnicity, or child's age, gender, or type of arthritis.'
Selective reporting (reporting bias)	Unclear risk	No reference to study protocol. Insufficient information to permit judgement of low risk or high risk.
Other bias	Unclear risk	Orthoses and shoe inserts reported to be donated by medical products companies. No assurance was made of their independence from the research design, conduct, or report.

Solanki 2020
Study characteristics

Methods	Randomised controlled two-arm parallel trial
Participants	<p>166 children screened, with 70 meeting the criteria, 44 randomised into two arms (22 each) and 44 provided final outcome data</p> <p>Age: children were in 8th and 9th standard in the school (probably 13 and 14 year olds)</p> <p>Gender: boys and girls, no other details provided</p> <p>Location: India</p> <p>Setting: schools in Bardoli, Gujarat, India</p> <p>Inclusion criteria: children with flat feet assessed using 'too many toes' sign, calcaneal angle, and navicular height</p> <p>Exclusion criteria: previous trauma or fracture of lower limb, history of previous surgeries of the lower limb during last 3 months, hypersensitive skin, and any allergy to tape</p> <p>Baseline characteristics: not reported</p>
Interventions	<p>Group A: anti-pronation taping + conventional treatment for 4 weeks</p> <p>Group B: sham taping + conventional treatment for 4 weeks</p> <p>A common (4 week) intervention program was executed for both groups as conventional therapy, which included strengthening exercise, stretching, & faradic foot bath that was given for 30 minutes per day</p>
Outcomes	star excursion balance test (SEBT), vertical jump height (VJH), and Illinois agility test (IAT) measured at 1, 2, 3, and 4 weeks
Source of funding	not described
Notes	We used 4 weeks of data for the SEBT scores, VJH, and IAT, as the intervention was given for 4 weeks.

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	"By use of lottery method, 44 children were randomly selected". We presumed this was drawing of lots.
Allocation concealment (selection bias)	Low risk	Allocation was concealed using opaque envelopes.
Blinding of participants and personnel (performance bias) All outcomes	Low risk	It appeared that from the participants' point of view, sham taping was not distinguishable from the anti-pronation taping.
Blinding of outcome assessment (detection bias) - self-reported outcomes (e.g., pain, function)	Low risk	no self-reported outcomes reported

Solanki 2020 (Continued)

Blinding of outcome assessment (detection bias) - objective outcomes	Low risk	measurement of the outcomes were unlikely to be affected by knowledge of the intervention
Incomplete outcome data (attrition bias) All outcomes	Low risk	no withdrawals
Selective reporting (reporting bias)	Unclear risk	trial was not registered; baseline characteristics not reported
Other bias	Low risk	none apparent

Wenger 1989
Study characteristics

Methods	Randomised controlled four-arm parallel trial
Participants	<p>131 children recruited, with 129 meeting the criteria, 98 provided final outcome data</p> <p>Age: 1 to 6 years; mean age at baseline 29.3 (13.6) months (N = 98)</p> <p>Gender: N = 129, not reported; final N = 98 - male 60, female 38</p> <p>Location: USA, Dallas</p> <p>Setting: Dept Orthopedic surgery, Children's Hospital, San Diego</p> <p>Inclusion criteria: 1 to 6 years old and flexible flat feet (observed to have valgus heel and poor arch formation in stance, further assessed by tip toe test, where varus shift of heel used to confirm flexible flat foot)</p> <p>Exclusion criteria: neurological condition (cerebral palsy or muscular disease), excessive laxity (Down or Ehlers-Danlos syndrome), or previous treatment with foot orthoses or corrective footwear</p> <p>Baseline characteristics: (CFO group had more younger children (12 to 32 months), fewer in 32 to 52 month age range, fewer at 52 to 72 months age. Shoe group had lowest joint laxity, heel cup group had highest laxity, sex ratio - not reported.</p>
Interventions	<p>Gp 1 - controls</p> <p>Gp 2 - corrective orthopaedic shoes</p> <p>Gp 3 - Helfet heel cups</p> <p>Gp 4 - custom moulded plastic inserts (UCBL)</p>
Outcomes	<p>- x-rays (lateral talo-horizontal angle > 35°, lateral talo-1st metatarsal > 10°, AP talo-calcaneal - too difficult to interpret)</p> <p>- photos (all x-rays and photos recorded every 6 months)</p> <p>- foot progression angle of gait (intoe/straight/out-toe), heel cord - assessed by passive ankle dorsiflexion, laxity (rated on Wynne-Davies 1 to 5 scale; low laxity = 0 or 1, high laxity = 2 to 5)</p> <p>- examined every 3 months by flatfoot team (nurse, Orthop Surg, Pedorthist) in unspecified manner. Shoe fit checked every 3 months by pedorthist, and shoes replaced whenever required for correct fitting.</p>

Wenger 1989 (Continued)

Commenced N = 129, completed N = 98 (lost n = 31, over 3 years of trial)

Group 1, 31 controls; 21 completed, lost 10

Group 2, 32 shoes; 28 completed, lost 4

Group 3, 35 Helfet; 27 completed, lost 8

Group 4, 31 UCBL; 22 completed, lost 9

Source of funding	shoes were provided
Notes	All participants wore specified footwear (some variation between groups) Minimum 3 years treatment and follow-up Attrition 98/129 (76%), and at completion, boys = 61/98 (61%)

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	The patients were assigned to treatment groups by a nurse who picked numbers randomly. It is not clear how this was done.
Allocation concealment (selection bias)	Unclear risk	Insufficient information to permit judgement of low risk or high risk.
Blinding of participants and personnel (performance bias) All outcomes	High risk	Participants were aware of intervention group. Personnel were aware of intervention group.
Blinding of outcome assessment (detection bias) - self-reported outcomes (e.g., pain, function)	Low risk	'At the time of three-year follow-up, clinical, radiographic, and photographic analyses were completed with the examiners in ignorance of the child's treatment group.'
Blinding of outcome assessment (detection bias) - objective outcomes	Unclear risk	31 participants were excluded from analysis as they were lost to follow up, mostly due to non-compliance; the trialists did not report which treatment these exclusions received, and this may have biased the results of the study
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	98/129 completed the trial; 31/129 participants: 'did not return for all of the follow-up visits, did not wear the shoes regularly, or moved from the state and were dropped from the study'. Insufficient reporting of attrition/exclusions to permit judgement of low risk or high risk
Selective reporting (reporting bias)	Unclear risk	No reference to study protocol. Insufficient information to permit judgement of low risk or high risk
Other bias	Unclear risk	'All shoes were provided by the Prescription Footwear Association'. No assurance was made of their independence from the research design, conduct, or report. However, since the report published negative findings, it suggests independence.

Whitford 2007

Study characteristics

Methods	Randomised clinical three-arm parallel trial
Participants	<p>672 children responded to advertisements</p> <p>Recruited 178 with flexible flat feet</p> <p>Age: 7 to 11 years</p> <p>Gender: male 53.4%, female 46.6%</p> <p>Inclusion: bilateral flexible flat feet, as indicated by RCSP > 5° everted, Navicular drop 10 mm or greater</p> <p>Exclusion: unilateral flexible excessive foot pronation, history of lower limb surgery, any serious medical pathology, such as cancer, known neuromuscular motor co-ordination condition, prior or recent foot orthoses use, intellectual or learning difficulty, or any chromosomal abnormality</p> <p>Location: Adelaide, South Australia</p> <p>Setting: outpatients</p> <p>Baseline characteristics: lower laxity score in CFO group 17.0 versus PFO 26.2, and shoes 30.6</p>
Interventions	<p>Gp 1: CFOs</p> <p>Gp 2: PFOs</p> <p>Gp 3: no orthoses (control)</p>
Outcomes	<p>Assessment at baseline, 3 months, 12 months</p> <p>Outcome included in the trial</p> <ol style="list-style-type: none"> 1. motor proficiency 2. pain - proportion with pain in subgroup that had pain (post hoc analysis) 3. exercise efficiency 4. self-perception profile <p>Outcome included in this review is proportion without pain at 12 months</p>
Source of funding	not stated (NHMRC)
Notes	<p>Most children did not have pain at baseline, pain analysed as subgroup</p> <p>Secondary outcomes measured included ligamentous laxity, tight calf muscles, and body mass index</p>

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Randomization was undertaken using computer generated lists of equally sized groups.
Allocation concealment (selection bias)	Unclear risk	Insufficient information to permit judgement of low risk or high risk.

Whitford 2007 *(Continued)*

Blinding of participants and personnel (performance bias) All outcomes	High risk	'... participants, and parents were not blinded.' 'Prescribing podiatrists... were not blinded'.
Blinding of outcome assessment (detection bias) - self-reported outcomes (e.g., pain, function)	High risk	Motor proficiency, pain, Self Perception Profile - outcomes are subjective or influenced by participants, and participants were not blinded
Blinding of outcome assessment (detection bias) - objective outcomes	Low risk	exercise efficiency - 'As in-shoe orthoses are concealed within the child's shoes when worn, raters of outcome measures were blinded to the orthotic status of the child.'
Incomplete outcome data (attrition bias) All outcomes	Low risk	Attrition rate and reasons similar between groups
Selective reporting (reporting bias)	Unclear risk	No reference to study protocol. The measurement of pain was done using VAS as a continuous measure, but reported only as a dichotomous outcome.
Other bias	Unclear risk	Funding source not stated

Characteristics of excluded studies *[ordered by study ID]*

Study	Reason for exclusion
Benedetti 2011	Not a randomised controlled trial
Ford 2017	Not a randomised controlled trial
Hill 2020	Review - not an RCT
Hurd 2010	Not a randomised controlled trial
MacKenzie 2012	Review - not a randomised controlled trial
Mosca 2010	Not a randomised controlled trial
Okamura 2020	Participants were university students, not children
Perhamre 2011	Study not addressing flat feet
Perhamre 2012	Study not addressing flat feet
Pothrat 2013	Not a randomised controlled trial
Riccio 2009	Not a randomised controlled trial
Uden 2017	Not a randomised controlled trial
Yung 2011	Participants were not children

Characteristics of studies awaiting classification *[ordered by study ID]*
Pandey 2013

Methods	Randomised controlled four-arm parallel trial
Participants	150 symptomatic flat foot, 50 controls (age > 8 years)
Interventions	<ol style="list-style-type: none"> 1. Foot exercises 2. Thomas heel ± arch support 3. Rose Schwartz insole 4. Foot exercise and both footwear modifications
Outcomes	<ul style="list-style-type: none"> • Foot Print Index • Valgus index • Pain • Gait (wear on medial versus lateral shoes)
Notes	<p>No variance, no age definition, no term of intervention, no randomisation details</p> <p>AE contacted Pandey via email x 3, searched internet for alternative address; unable to contact for additional data</p>

Sinha 2013

Methods	Randomised controlled two-arm parallel trial
Participants	<p>101 recruited – children with symptomatic flat feet</p> <p>81 children finished trial</p> <p>20 lost or non-compliant</p>
Interventions	<ol style="list-style-type: none"> 1. Foot orthosis N = 55 2. Control N = 26
Outcomes	<ul style="list-style-type: none"> • x-ray angles • AOFAS pain scores
Notes	<ul style="list-style-type: none"> • Check randomisation method "81 children with bilateral symptomatic flatfoot were randomised into orthosis (N = 55) and control (N = 26) groups" • Will need mean and SD for each group for American Orthopaedic Foot and Ankle Society (AOFAS) scores for pain • Age 3 to 17: will need data for children 3 to 15 only <p>Unable to contact author – 3 emails; searched for alternative addresses – no contact made</p>

Characteristics of ongoing studies *[ordered by study ID]*
ACTRN12616001082493

Study name	Effectiveness of foot orthoses in children with juvenile idiopathic arthritis: a randomised control trial
------------	-----------------------------------------------------------------------------------------------------------

ACTRN12616001082493 (Continued)

Methods	Randomised controlled two-arm trial - 12 months
Participants	<p>Children with juvenile idiopathic arthritis (JIA)</p> <ol style="list-style-type: none"> 1. Diagnosed with JIA according to ILAR (International League of Associations for Rheumatology) criteria 2. Age 5 to 18 years old 3. Active lower-limb joint arthritis involvement 4. No previous use of foot orthoses, or previous failure of foot orthotic management, where the patient has not worn any foot orthoses (FO) for a period of at least 3 months 5. Ability to walk a minimum of 15 meters without assistive devices 6. If disease-modifying antirheumatic drugs (DMARD), biological therapy, or both are used, not having started these drug therapies within 6 months of enrolling in the trial
Interventions	Customised prefabricated foot orthoses (PFO), sham insole (control)
Outcomes	pain, disability, quality of life
Starting date	not stated
Contact information	www.who.int/trialsearch/Trial2.aspx?TrialID=ACTRN12616001082493
Notes	

CTRI/2018/07/014989

Study name	Effect of adhesive taping and exercises on outward rotation of foot in affected children
Methods	Randomised controlled two-arm cross-over trial - 4 weeks
Participants	<p>Children with pronated feet and neurodevelopmental disorders:</p> <ol style="list-style-type: none"> 1) Diagnosed cases of non-progressive neurodevelopmental disorders by paediatrician 2) Foot Posture Index score more than 6 3) Navicular Drop Test more than 10 mm 4) Both male and female 5) Age 6 to 11 years 6) Patient should be able to stand without support or with minimum hand support 7) Flexible pes planus
Interventions	<p>Intervention: Group A: kinesio taping for 5 days for 2 weeks, and then group A will be switched to Group B therapy after 2 weeks of window period</p> <p>Control: Group B: exercises for 5 days for 2 weeks, and then group B will be switched to Group A after 2 weeks of window period</p>
Outcomes	1. Foot Posture Index/time point

CTRI/2018/07/014989 (Continued)

2. Navicular Drop Test/time point

Starting date	not stated
Contact information	www.who.int/trialsearch/Trial2.aspx?TrialID=CTRI/2018/07/014989
Notes	

CTRI/2019/08/020925

Study name	Effects of barefoot walking on the flat foot in school going children: a randomised control trial
Methods	Randomised control trial
Participants	N = 38, age between 6 and 14 years
Interventions	<p>Intervention 1: Barefoot walking and exercises: children have to be barefoot while doing exercises, as follows:</p> <ol style="list-style-type: none"> 1. Towel gathering exercise for 15 minutes 2. Heel cord stretching (holding for 30 seconds and then relax for 30 seconds; repeat once) 3. Toe spread (5 seconds, than 2 second relax) 4. Posterior tibialis exercises (3 sets, 10 repetitions) <p>Intervention 2: Bare foot exercises and Exercises: Bare foot walking and exercises.
Exercises are,</p> <ol style="list-style-type: none"> A. Towel gathering exercise for 15 minutes B. Heel cord stretching (holding for 30 seconds and then relax for 30 seconds and then repeat for 1times) C. Toe spread [5 sec than 2 sec relax] D. Posterior Tibialis exercises [3sets, 10reps] <p>Control Intervention1: Not applicable</p>
Outcomes	The Oxford Ankle Foot Questionnaire for Children (OxAFQ-C) time point: 8 weeks
Starting date	27 August 2019, not recruiting
Contact information	Sharath HV, chippala1979@gmail.com. www.ctri.nic.in/Clinicaltrials/pmaindet2.php?trialid=36115
Notes	Nitte institute of physiotherapy, Nitte University, India

IRCT2017082235517N1

Study name	Effects of long-term use of arch support foot orthoses on lower limb kinematics and kinetics during walking in children with flexible flat foot
Methods	Randomised control trial

IRCT2017082235517N1 (Continued)

Participants	N = 30 Inclusion criteria: age range between 9 and 12 years; flexible flat feet; male Exclusion criteria: history of lower limb injury; history of surgery
Interventions	Intervention 1: intervention group: 4 months use of arch support foot orthoses Intervention 2: control group: long-term use of placebo orthoses; rehabilitation; placebo; Intervention group: 4 months use of arch support foot orthoses; Control group: long term use of placebo orthoses
Outcomes	Walking kinematics Time point: before and after 4 months Method of measurement: vicon cameras; walking kinetics. Timepoint: before and after 4 months. Method of measurement: Force platform.
Starting date	4 September 2017
Contact information	Amir Ali Jafarnejadgero, amiralijafarnejad@gmail.com; en.irct.ir/trial/26811
Notes	Mohaghegh Ardabili University, Iran (Islamic Republic of)

ISRCTN14602568

Study name	Orthotics for treatment of symptomatic flat feet in children - the OSTRICH study
Methods	Interventional randomised controlled trial including a qualitative study within a trial (treatment)
Participants	N = 1055 Inclusion criteria: 1. Aged between 6 and 14 years, inclusive 2. Have one or both symptomatic pes planus* 3. The child, parent, or legal guardian is able to speak, write and understand English 4. The parent or legal guardian is able to give informed consent *Symptomatic pes planus is described as the manifestation of foot and lower limb symptoms, secondary to altered foot alignment (reduced medial longitudinal arch, everted rearfoot, and abducted forefoot). The diagnosis will be made pragmatically, by treating clinicians, in line with current practice. Eligibility for the birthday card SWAT All participants recruited into the host trial will be eligible to take part in this SWAT.
Interventions	Participants will be allocated to one of three groups: 1. A package consisting of an exercise programme and advice covering topics, such as typical arch development in children, coping strategies, and footwear advice 2. The exercise and advice package plus a pair of prefabricated, off-the-shelf orthoses (i.e. insoles that are mass produced to a generic shape, but can be adapted by a clinician)

ISRCTN14602568 (Continued)

3. The exercise and advice package plus a pair of custom-made foot orthoses, where the shape of the insole is made for a specific person, based on a 3D impression or scan of the patient's foot

The participant will be informed which group they have been allocated to at their first trial appointment. Participants will be asked to wear their orthoses every day in their shoes, and to do their exercises. Participants in group 1 and 2 will, on the whole, need one clinic appointment, but those in group 3, may need a total of 2 or 3 appointments. Blinding of participants to the treatment allocation is not possible.

Measures will be collected at baseline, weeks 1 to 12, and at three, six, and 12 months after a participant is enrolled into the study. The measures are all self-reported by either the participant or their parent or legal guardian.

Birthday card study: In the birthday card SWAT, the researchers will evaluate whether sending a participant a birthday card increases the number of questionnaires they return to the study team. Participants will be allocated to one of three groups; birthday card, birthday card informed by nudge theory to encourage completion of questionnaires, or no birthday card.

Outcomes	Physical domain subscale of the Oxford Ankle Foot Questionnaire for Children (OxAFQ-C); Time point(s): over the 12-month follow-up period
Starting date	18 May 2020
Contact information	David Torgerson, PhD; Sarah Cockayne, MSc; sarah.cockayne@york.ac.uk www.isrctn.com/ISRCTN14602568
Notes	University of York, UK

ISRCTN49672274

Study name	Foot disease in juvenile idiopathic arthritis: foot care trial
Methods	Randomised clinical two-arm trial - 12 months
Participants	Children with JIA: <ol style="list-style-type: none"> juvenile idiopathic arthritis (JIA) diagnosed according to the International League of Associations for Rheumatology (ILAR) 2004 criteria Lower limb arthritis of two or more large joints (hips, knees, ankles, and subtalar joints) Widespread polyarthritis involving large and small joints
Interventions	Usual clinical care (standard podiatry) versus individualised care package (foot orthoses (FO), shoes, physiotherapy, podiatry)
Outcomes	Juvenile Arthritis Foot Disability Index (JAFI) Function HRQoL (health-related quality of life)
Starting date	not stated
Contact information	/www.who.int/trialsearch/Trial2.aspx?TrialID=ISRCTN49672274
Notes	

Foot orthoses for treating paediatric flat feet (Review)

KCT0001717

Study name	The effects of talus control foot orthosis in children with flexible flatfoot
Methods	Randomised control trial
Participants	N = 40 Inclusion criteria: diagnosed with flexible flatfoot; visited the Department of Pediatric Rehabilitation over 6 years Exclusion criteria: rigid flatfoot caused by hereditary foot disease or neuromuscular disease, fixed foot deformity, or previous surgery history
Interventions	Device: plantar orthoses
Outcomes	X-ray measurements of the angles: talus-calcaneus, talus-navicular, and talus-first metatarsal angle
Starting date	7 December 2015
Contact information	So-Young Ahn
Notes	Chungnam National University Hospital, Republic of Korea

NCT02414087

Study name	Therapeutic effects of customized insoles on children with flatfoot
Methods	Randomised controlled trial - two arms - 12 weeks
Participants	52 children with flat feet
Interventions	1. customised full-length insoles 2. control
Outcomes	physical function balance quality of life
Starting date	not stated
Contact information	clinicaltrials.gov/show/NCT02414087
Notes	Ru-Lan Hsieh, Shin Kong Wu Ho-Su Memorial Hospital, Taipei Medical University

NCT02633566

Study name	Clinical trial of the effect of functional orthoses in children with flat feet
Methods	randomised, double-blind, parallel design clinical trial - 1 year

Foot orthoses for treating paediatric flat feet (Review)

NCT02633566 (Continued)

Participants	61 children, aged 3 and 4 years old
Interventions	1. Intervention group - functional plantar orthotics 2. Control group - placebo-type plantar orthotics
Outcomes	X-rays: talus-calcaneus, tibionavicular, and talus-first metatarsal angles were compared pre- and post-treatment
Starting date	not stated
Contact information	clinicaltrials.gov/show/NCT02633566
Notes	Raquel Lopez Fresno, PhD Student (Podiatrist)

NCT03151538

Study name	Effects on pes planus exercise training mixed with play on pre-school children
Methods	not stated
Participants	Pre-school children with pes planus (flat feet)
Interventions	exercise training mixed with play
Outcomes	effects on pes planus and femoral anteversion angle
Starting date	not stated
Contact information	clinicaltrials.gov/show/NCT03151538
Notes	Burcu Talu, Assistant Professor, Inonu University

NCT04104555

Study name	Orthotics for treatment of symptomatic flat feet in children - the OSTRICH study
Methods	
Participants	<ul style="list-style-type: none"> - Have pes planus secondary to any systematic condition or syndrome**, malignancy - Have a history of foot or ankle surgery, or both - Require an ankle-foot orthoses, or other lower limb device, or have received treatment previously for their flat feet - This does not exclude children with hypermobility spectrum disorder (HSD), where the manifestation is non-syndromic and isolated (L-HSD), peripheral (P-HSD), or generalised hypermobility (G-HSD)(14). <p>OSTRICH pen and signposting to multimedia trial information SWAT inclusion criteria: any potential participant identified as eligible to be sent trial information will be eligible for the study</p> <p>OSTRICH birthday card SWAT inclusion criteria: all participants recruited into the host trial will be eligible to take part in this SWAT. Exclusion criteria: any participant who has withdrawn from the main OSTRICH study, or is not due a follow-up questionnaire will be excluded.</p>
Interventions	Other: Birthday card;

Foot orthoses for treating paediatric flat feet (Review)

NCT04104555 (Continued)

	Other: Birthday card plus nudge;
	Device: Prefabricated orthoses;
	Device: Custom orthoses;
	Other: Pen;Other: Signposting to multimedia;
	Other: Pen and signposting to multimedia;
	Other: Usual supportive care - exercises and footwear advice (the comparator)
Outcomes	Physical domain subscale of the Oxford Ankle Foot Questionnaire for Children (OxAFQ-C) over the 12-month follow-up period
Starting date	19 September 2019
Contact information	David Torgerson, PhD; Sarah Cockayne, MSc; sarah.cockayne@york.ac.uk clinicaltrials.gov/ct2/show/NCT04104555
Notes	University of York, UK

NCT04410926

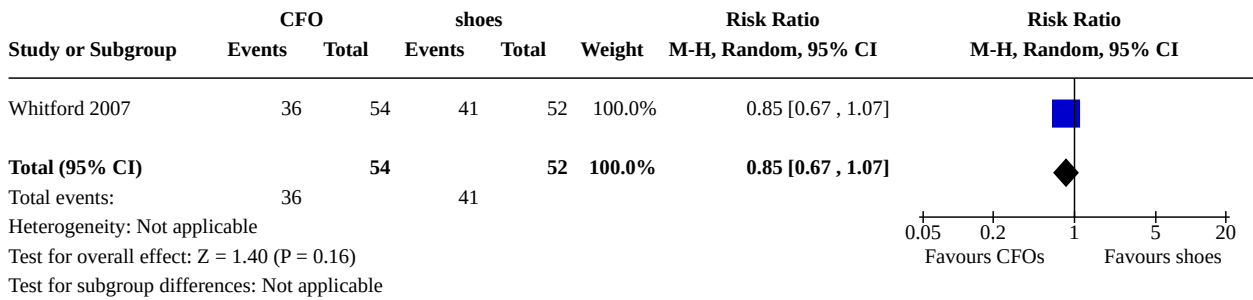
Study name	Corrective exercises with neuromuscular electrical stimulation In children with flexible flat feet
Methods	prospective double-masked randomised controlled trial
Participants	70 school children were recruited, 67 were randomised
Interventions	1. intervention group (corrective exercise and neuromuscular electrical stimulation (NMES)) 2. control group (corrective exercise and placebo NMES)
Outcomes	not stated
Starting date	1 September 2018 to 31 December 2019
Contact information	clinicaltrials.gov/show/NCT04410926
Notes	Ethics Statement - study was approved by the Institutional Review Board of the Faculty of Physical Therapy, Cairo University, Egypt (no. P.T.REC/012/0016370)

DATA AND ANALYSES
Comparison 1. Custom foot orthoses (CFOs) versus shoes for asymptomatic flat feet

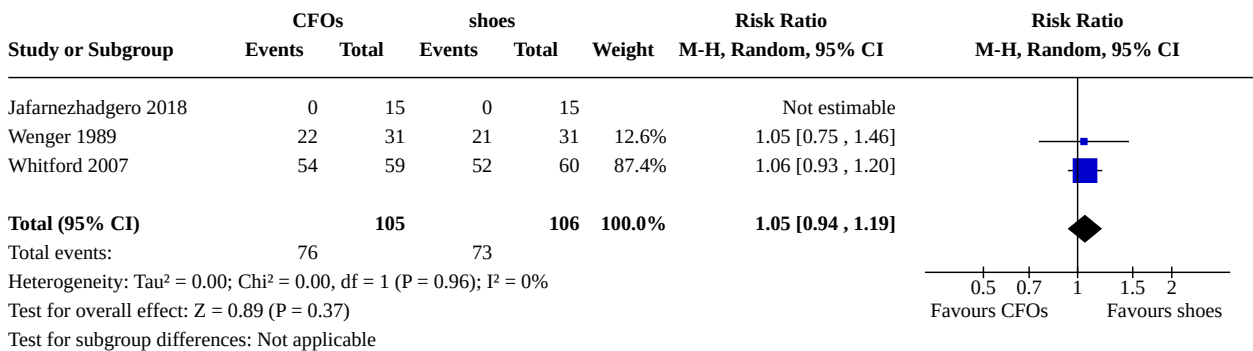
Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1.1 Proportion without pain	1	106	Risk Ratio (M-H, Random, 95% CI)	0.85 [0.67, 1.07]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1.2 Withdrawal due to adverse events	3	211	Risk Ratio (M-H, Random, 95% CI)	1.05 [0.94, 1.19]

Analysis 1.1. Comparison 1: Custom foot orthoses (CFOs) versus shoes for asymptomatic flat feet, Outcome 1: Proportion without pain



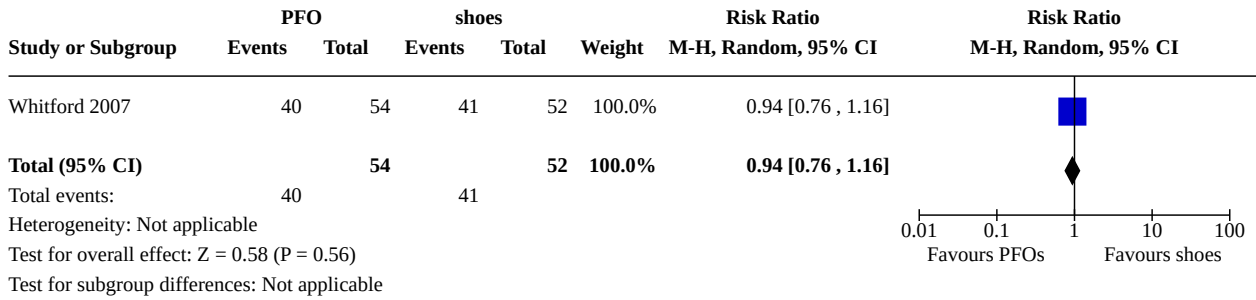
Analysis 1.2. Comparison 1: Custom foot orthoses (CFOs) versus shoes for asymptomatic flat feet, Outcome 2: Withdrawal due to adverse events



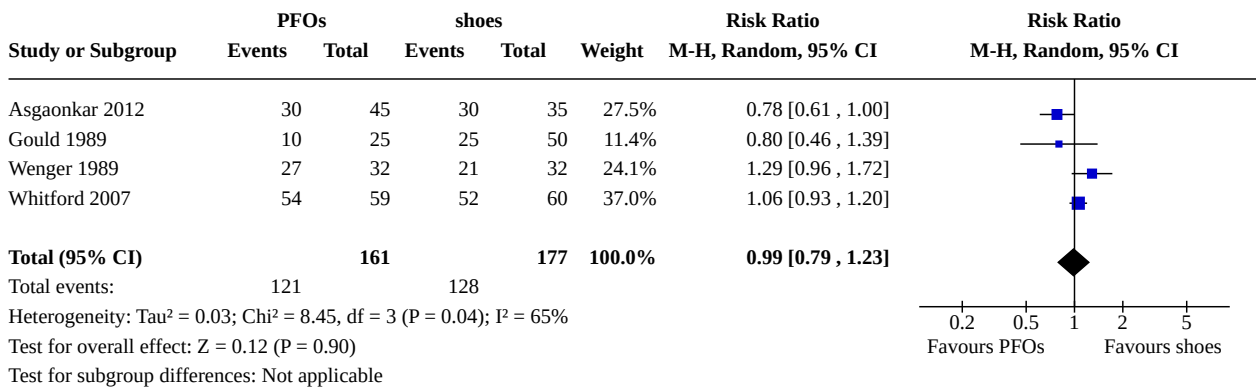
Comparison 2. Prefabricated foot orthoses (PFOs) versus shoes in asymptomatic flat feet

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
2.1 Proportion without pain	1	106	Risk Ratio (M-H, Random, 95% CI)	0.94 [0.76, 1.16]
2.2 Withdrawal due to adverse events	4	338	Risk Ratio (M-H, Random, 95% CI)	0.99 [0.79, 1.23]

Analysis 2.1. Comparison 2: Prefabricated foot orthoses (PFOs) versus shoes in asymptomatic flat feet, Outcome 1: Proportion without pain



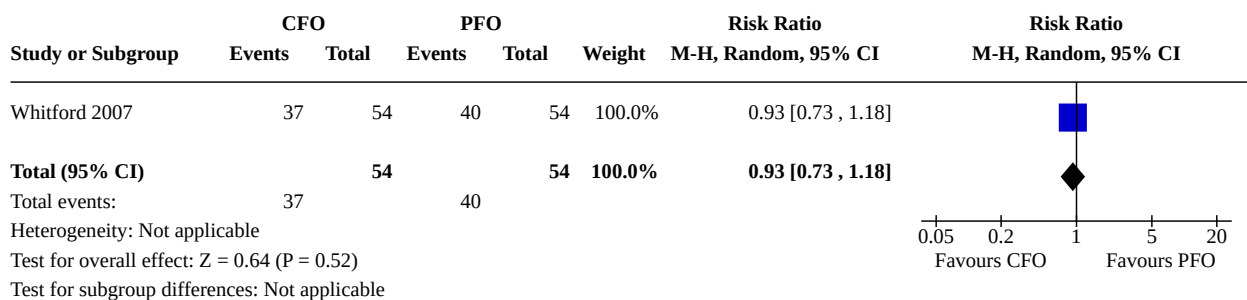
Analysis 2.2. Comparison 2: Prefabricated foot orthoses (PFOs) versus shoes in asymptomatic flat feet, Outcome 2: Withdrawal due to adverse events



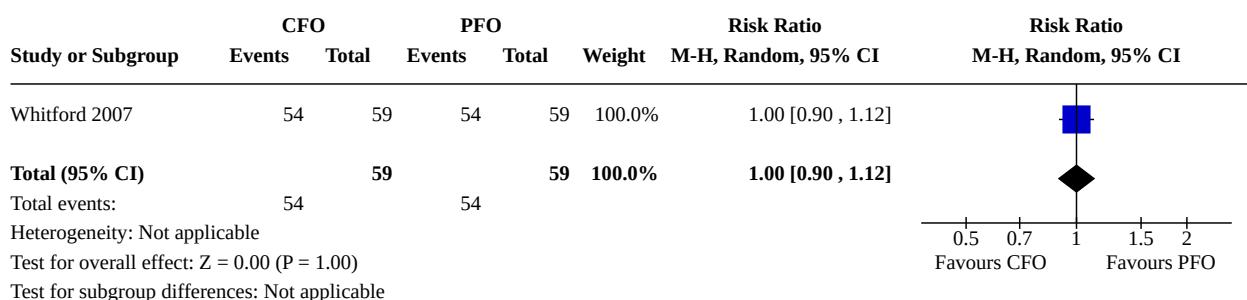
Comparison 3. CFOs versus PFOs in asymptomatic flat feet

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
3.1 Pain	1	108	Risk Ratio (M-H, Random, 95% CI)	0.92 [0.73, 1.18]
3.2 Withdrawal due to adverse events	1	118	Risk Ratio (M-H, Random, 95% CI)	1.00 [0.90, 1.12]

Analysis 3.1. Comparison 3: CFOs versus PFOs in asymptomatic flat feet, Outcome 1: Pain



Analysis 3.2. Comparison 3: CFOs versus PFOs in asymptomatic flat feet, Outcome 2: Withdrawal due to adverse events





Comparison 4. CFOs versus shoes in juvenile idiopathic arthritis (JIA)

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
4.1 Pain	1	28	Mean Difference (IV, Random, 95% CI)	-1.50 [-2.78, -0.22]
4.2 Function	1		Mean Difference (IV, Random, 95% CI)	Subtotals only
4.2.1 foot pain - FFI	1	28	Mean Difference (IV, Random, 95% CI)	-19.19 [-35.50, -2.88]
4.2.2 activity limitation - FFI	1	28	Mean Difference (IV, Random, 95% CI)	-19.38 [-35.54, -3.22]
4.2.3 disability - FFI	1	28	Mean Difference (IV, Random, 95% CI)	-18.55 [-34.42, -2.68]
4.3 Quality of life	1		Mean Difference (IV, Random, 95% CI)	Subtotals only
4.3.1 PedsQL physical - child-rated	1	25	Mean Difference (IV, Random, 95% CI)	12.10 [-1.60, 25.80]
4.3.2 Peds QL physical - parent-rated	1	26	Mean Difference (IV, Random, 95% CI)	9.01 [-4.08, 22.10]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
4.4 Treatment success (gait parameters)	1	28	Mean Difference (IV, Random, 95% CI)	-1.33 [-2.77, 0.11]
4.5 Withdrawal due to adverse events	1	28	Risk Ratio (M-H, Random, 95% CI)	0.58 [0.11, 2.94]







Analysis 4.1. Comparison 4: CFOs versus shoes in juvenile idiopathic arthritis (JIA), Outcome 1: Pain

Study or Subgroup	CFOs			Shoes			Weight	Mean Difference IV, Random, 95% CI	Mean Difference IV, Random, 95% CI
	Mean	SD	Total	Mean	SD	Total			
Powell 2005	1.32	1.3	15	2.82	2.01	13	100.0%	-1.50 [-2.78, -0.22]	
Total (95% CI)			15			13	100.0%	-1.50 [-2.78, -0.22]	

Heterogeneity: Not applicable
Test for overall effect: Z = 2.31 (P = 0.02)
Test for subgroup differences: Not applicable

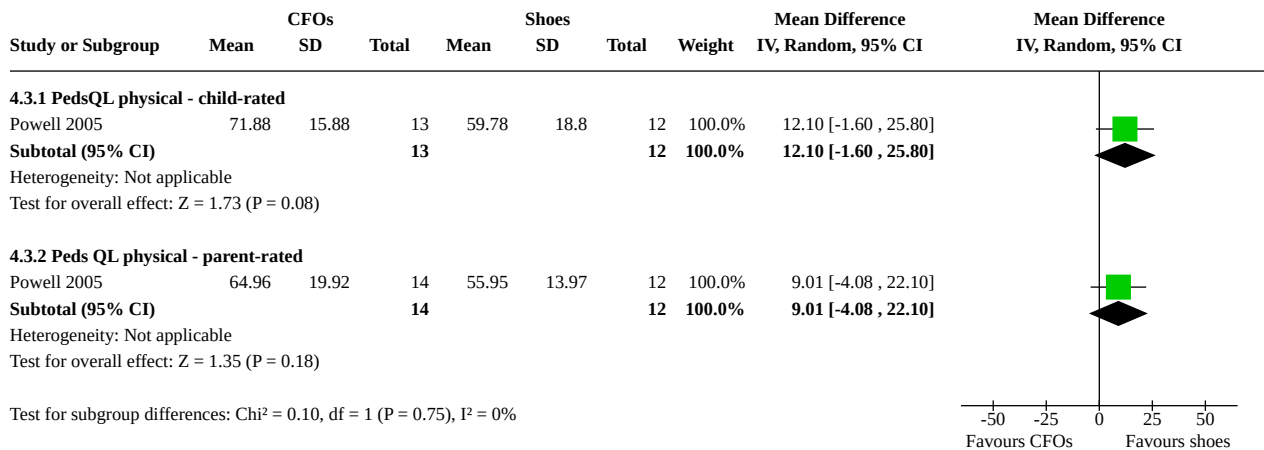
-4 -2 0 2 4
Favours CFO Favours shoes

Analysis 4.2. Comparison 4: CFOs versus shoes in juvenile idiopathic arthritis (JIA), Outcome 2: Function

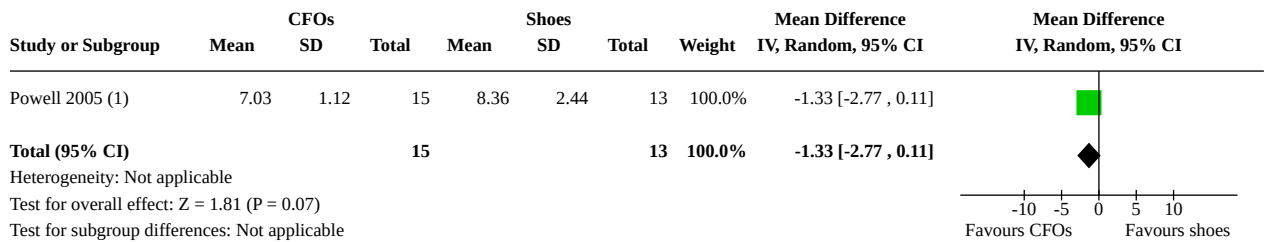
Study or Subgroup	CFOs			shoes			Weight	Mean Difference IV, Random, 95% CI	Mean Difference IV, Random, 95% CI
	Mean	SD	Total	Mean	SD	Total			
4.2.1 foot pain - FFI									
Powell 2005	18.35	17.05	15	37.54	25.47	13	100.0%	-19.19 [-35.50, -2.88]	
Subtotal (95% CI)			15			13	100.0%	-19.19 [-35.50, -2.88]	
Heterogeneity: Not applicable Test for overall effect: Z = 2.31 (P = 0.02)									
4.2.2 activity limitation - FFI									
Powell 2005	8.54	11.06	15	27.92	27.89	13	100.0%	-19.38 [-35.54, -3.22]	
Subtotal (95% CI)			15			13	100.0%	-19.38 [-35.54, -3.22]	
Heterogeneity: Not applicable Test for overall effect: Z = 2.35 (P = 0.02)									
4.2.3 disability - FFI									
Powell 2005	15.6	13.51	15	34.15	26.35	13	100.0%	-18.55 [-34.42, -2.68]	
Subtotal (95% CI)			15			13	100.0%	-18.55 [-34.42, -2.68]	
Heterogeneity: Not applicable Test for overall effect: Z = 2.29 (P = 0.02)									
Test for subgroup differences: Chi ² = 0.01, df = 2 (P = 1.00), I ² = 0%									

-100 -50 0 50 100
Favours CFOs Favours shoes

Analysis 4.3. Comparison 4: CFOs versus shoes in juvenile idiopathic arthritis (JIA), Outcome 3: Quality of life



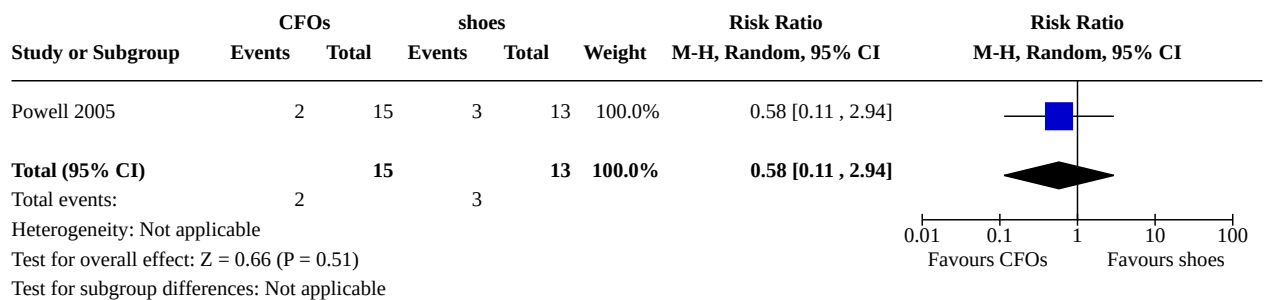
Analysis 4.4. Comparison 4: CFOs versus shoes in juvenile idiopathic arthritis (JIA), Outcome 4: Treatment success (gait parameters)



Footnotes

(1) 50FWT (seconds)

Analysis 4.5. Comparison 4: CFOs versus shoes in juvenile idiopathic arthritis (JIA), Outcome 5: Withdrawal due to adverse events

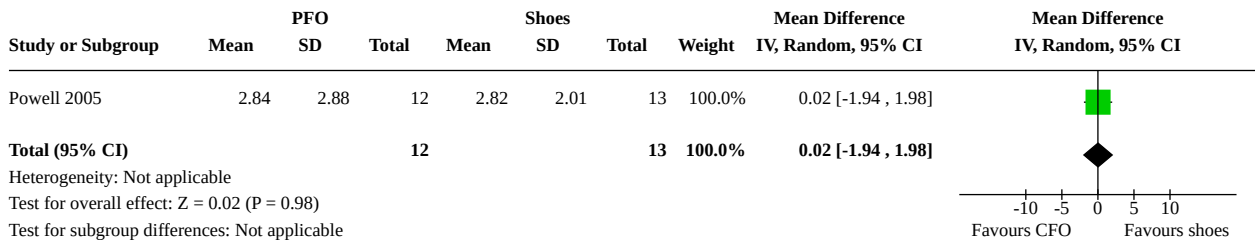


Comparison 5. PFOs versus shoes in JIA

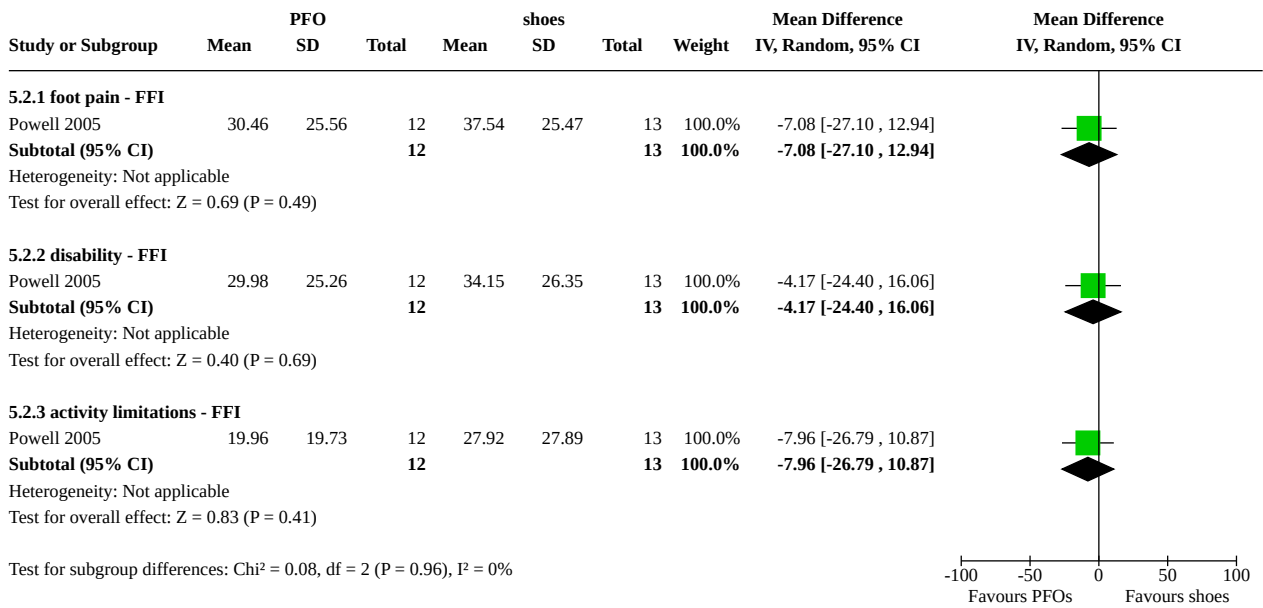
Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
5.1 Pain	1	25	Mean Difference (IV, Random, 95% CI)	0.02 [-1.94, 1.98]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
5.2 Function	1		Mean Difference (IV, Random, 95% CI)	Subtotals only
5.2.1 foot pain - FFI	1	25	Mean Difference (IV, Random, 95% CI)	-7.08 [-27.10, 12.94]
5.2.2 disability - FFI	1	25	Mean Difference (IV, Random, 95% CI)	-4.17 [-24.40, 16.06]
5.2.3 activity limitations - FFI	1	25	Mean Difference (IV, Random, 95% CI)	-7.96 [-26.79, 10.87]
5.3 Quality of life	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
5.3.1 PedsQL physical - child-rated	1	22	Mean Difference (IV, Fixed, 95% CI)	-3.84 [-19.01, 11.33]
5.3.2 PedsQL physical - parent-rated	1	22	Mean Difference (IV, Fixed, 95% CI)	-0.64 [-13.22, 11.94]
5.4 Treatment success (Timed walking)	1	25	Mean Difference (IV, Random, 95% CI)	-0.38 [-1.90, 1.14]
5.5 Withdrawal due to adverse events	1	25	Risk Ratio (M-H, Random, 95% CI)	0.72 [0.14, 3.61]

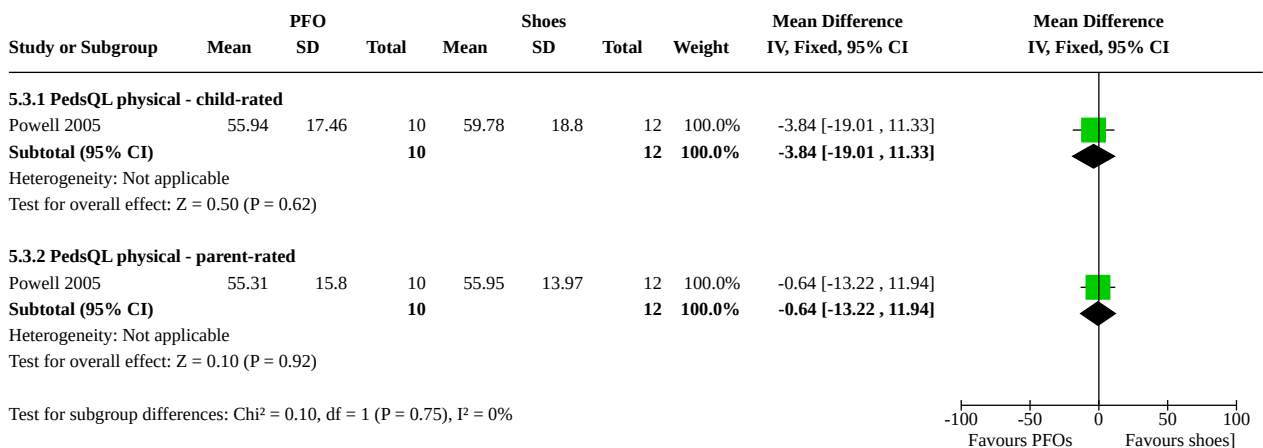
Analysis 5.1. Comparison 5: PFOs versus shoes in JIA, Outcome 1: Pain



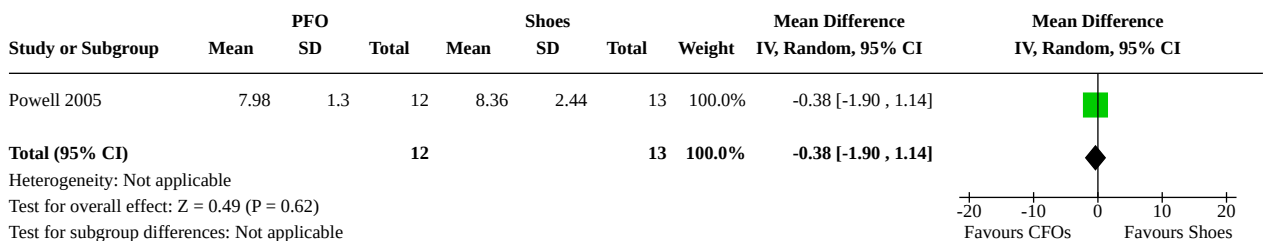
Analysis 5.2. Comparison 5: PFOs versus shoes in JIA, Outcome 2: Function



Analysis 5.3. Comparison 5: PFOs versus shoes in JIA, Outcome 3: Quality of life



Analysis 5.4. Comparison 5: PFOs versus shoes in JIA, Outcome 4: Treatment success (Timed walking)



Analysis 5.5. Comparison 5: PFOs versus shoes in JIA, Outcome 5: Withdrawal due to adverse events

Study or Subgroup	PFO		shoes		Weight	Risk Ratio		Risk Ratio	
	Events	Total	Events	Total		M-H, Random, 95% CI	M-H, Random, 95% CI		
Powell 2005	2	12	3	13	100.0%	0.72 [0.14 , 3.61]			
Total (95% CI)		12		13	100.0%	0.72 [0.14 , 3.61]			
Total events:	2		3						
Heterogeneity: Not applicable									
Test for overall effect: Z = 0.40 (P = 0.69)									
Test for subgroup differences: Not applicable									

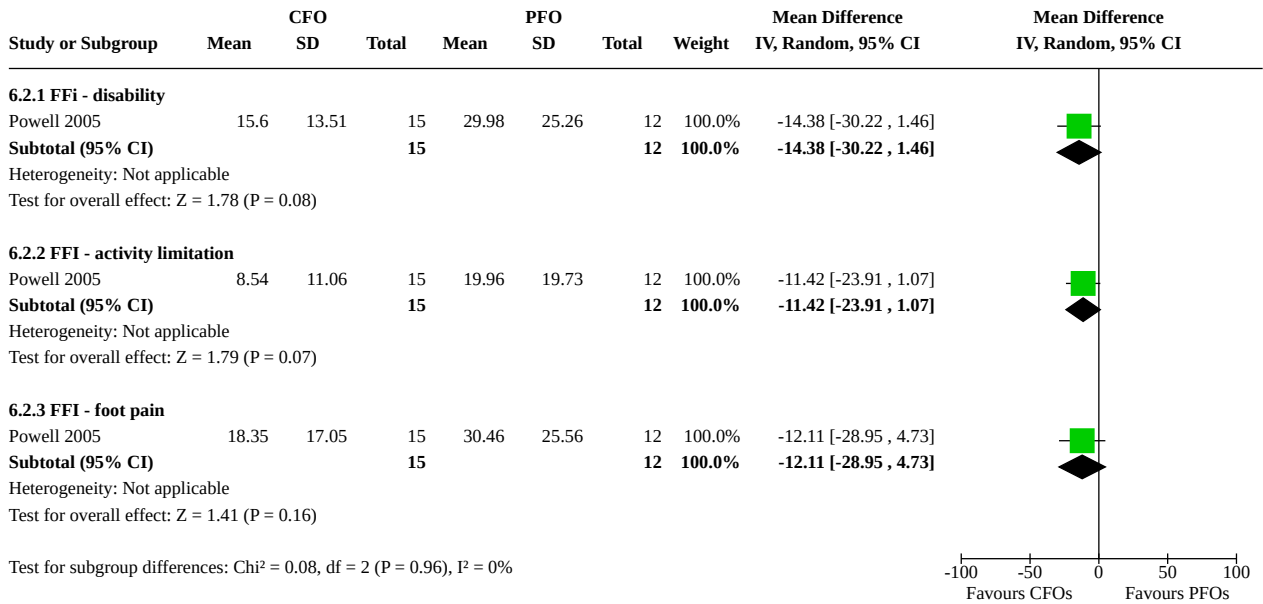
Comparison 6. CFOs versus PFOs in JIA

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
6.1 Pain	2	87	Mean Difference (IV, Random, 95% CI)	-1.48 [-3.23, 0.26]
6.2 Function	1		Mean Difference (IV, Random, 95% CI)	Subtotals only
6.2.1 FFI - disability	1	27	Mean Difference (IV, Random, 95% CI)	-14.38 [-30.22, 1.46]
6.2.2 FFI - activity limitation	1	27	Mean Difference (IV, Random, 95% CI)	-11.42 [-23.91, 1.07]
6.2.3 FFI - foot pain	1	27	Mean Difference (IV, Random, 95% CI)	-12.11 [-28.95, 4.73]
6.3 Quality of life	2		Mean Difference (IV, Random, 95% CI)	Subtotals only
6.3.1 PedsQL - child-rated	2	83	Mean Difference (IV, Random, 95% CI)	8.64 [-3.90, 21.18]
6.3.2 PedsQL - parent-rated	2	84	Mean Difference (IV, Random, 95% CI)	2.94 [-11.00, 16.88]
6.4 Treatment success (timed walking)	1	27	Mean Difference (IV, Random, 95% CI)	-0.95 [-1.88, -0.02]
6.5 Withdrawal due to adverse events	2	87	Risk Ratio (M-H, Random, 95% CI)	0.80 [0.13, 4.87]

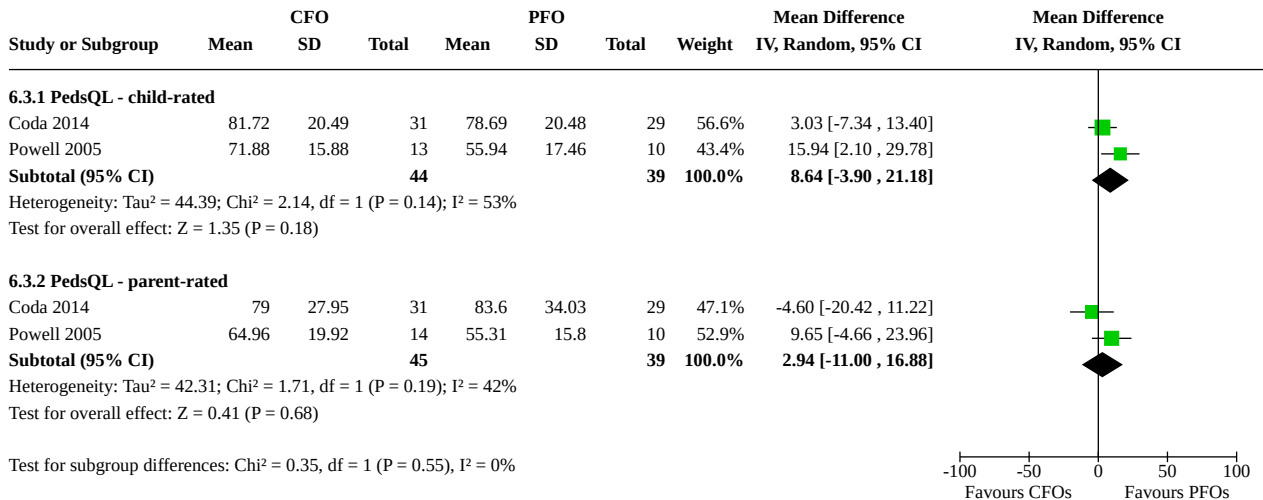
Analysis 6.1. Comparison 6: CFOs versus PFOs in JIA, Outcome 1: Pain

Study or Subgroup	CFO		PFO		Total	Weight	Mean Difference		Mean Difference	
	Mean	SD	Mean	SD			IV, Random, 95% CI	IV, Random, 95% CI		
Coda 2014	5	24	31	3.6	34	29	1.4%	1.40 [-13.58 , 16.38]		
Powell 2005	1.32	1.3	15	2.84	2.88	12	98.6%	-1.52 [-3.28 , 0.24]		
Total (95% CI)			46			41	100.0%	-1.48 [-3.23 , 0.26]		
Heterogeneity: Tau ² = 0.00; Chi ² = 0.14, df = 1 (P = 0.70); I ² = 0%										
Test for overall effect: Z = 1.66 (P = 0.10)										
Test for subgroup differences: Not applicable										

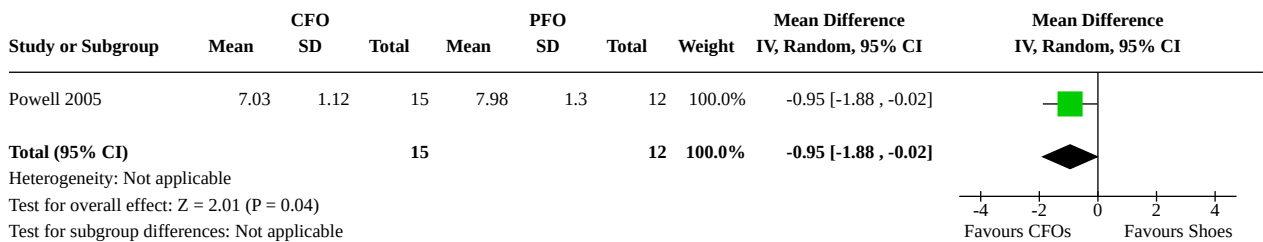
Analysis 6.2. Comparison 6: CFOs versus PFOs in JIA, Outcome 2: Function



Analysis 6.3. Comparison 6: CFOs versus PFOs in JIA, Outcome 3: Quality of life



Analysis 6.4. Comparison 6: CFOs versus PFOs in JIA, Outcome 4: Treatment success (timed walking)



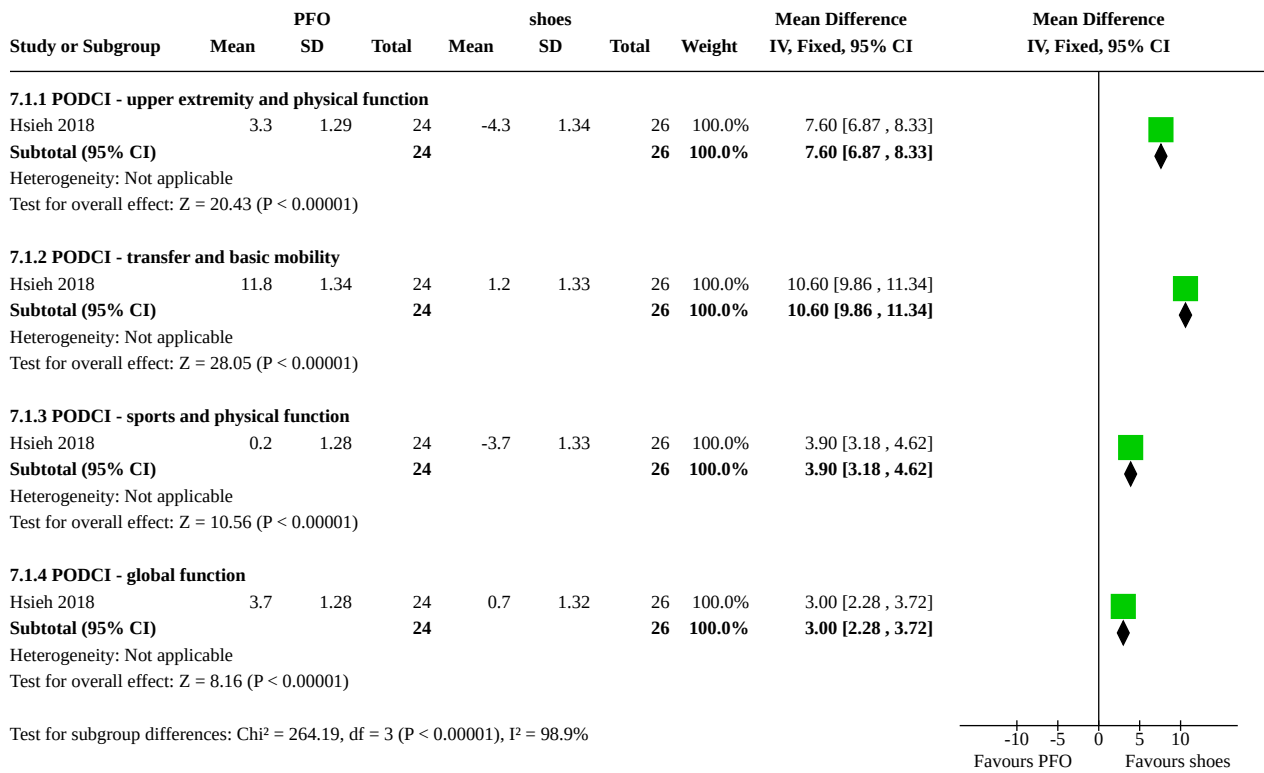
Analysis 6.5. Comparison 6: CFOs versus PFOs in JIA, Outcome 5: Withdrawal due to adverse events

Study or Subgroup	CFO		PFO		Weight	Risk Ratio	Risk Ratio
	Events	Total	Events	Total		M-H, Random, 95% CI	M-H, Random, 95% CI
Coda 2014	0	31	0	29		Not estimable	
Powell 2005	2	15	2	12	100.0%	0.80 [0.13, 4.87]	
Total (95% CI)		46		41	100.0%	0.80 [0.13, 4.87]	
Total events:	2		2				
Heterogeneity: Not applicable							
Test for overall effect: Z = 0.24 (P = 0.81)							
Test for subgroup differences: Not applicable							

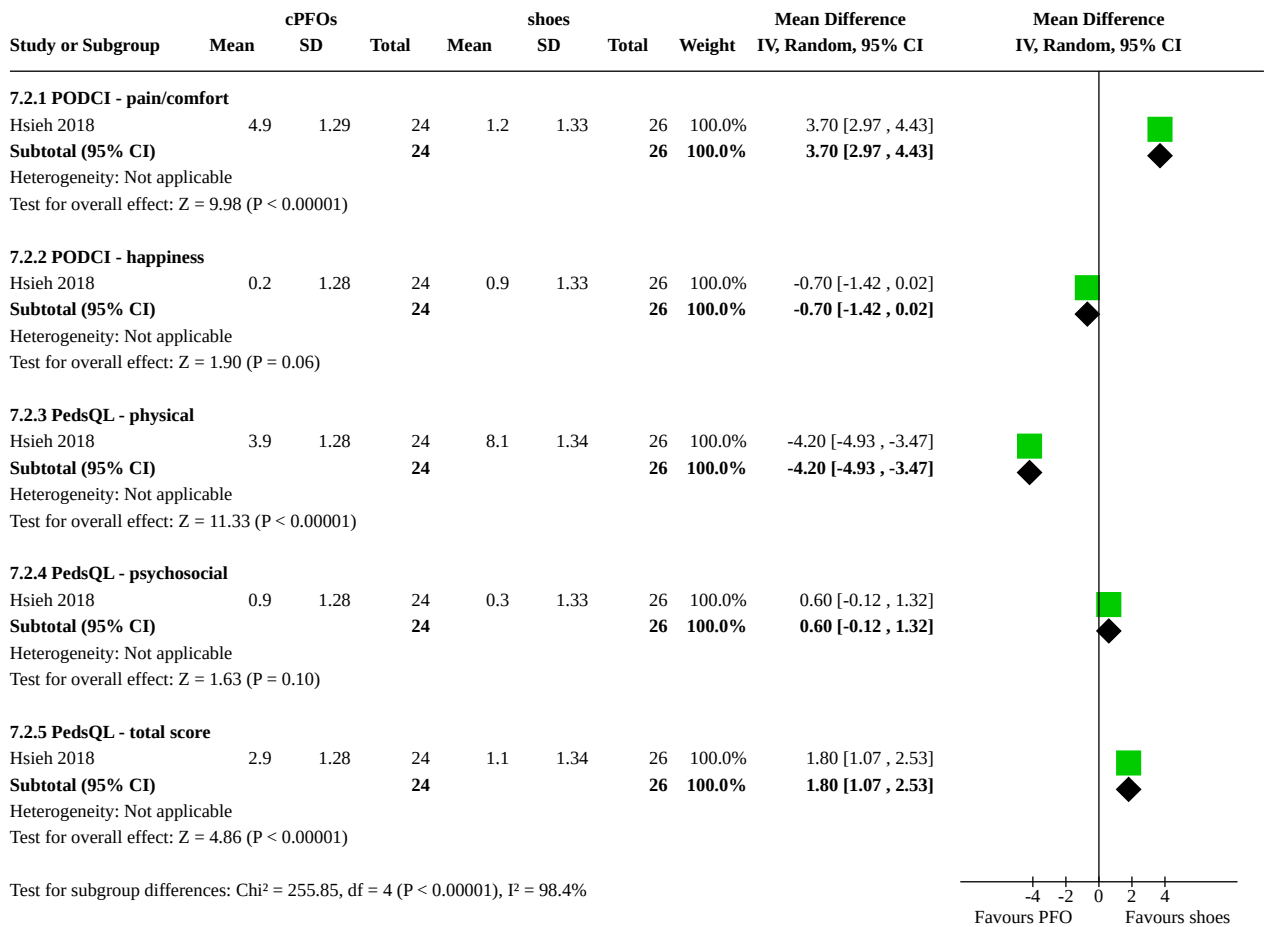
Comparison 7. PFOs versus shoes in symptomatic flat feet

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
7.1 Function	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
7.1.1 PODCI - upper extremity and physical function	1	50	Mean Difference (IV, Fixed, 95% CI)	7.60 [6.87, 8.33]
7.1.2 PODCI - transfer and basic mobility	1	50	Mean Difference (IV, Fixed, 95% CI)	10.60 [9.86, 11.34]
7.1.3 PODCI - sports and physical function	1	50	Mean Difference (IV, Fixed, 95% CI)	3.90 [3.18, 4.62]
7.1.4 PODCI - global function	1	50	Mean Difference (IV, Fixed, 95% CI)	3.00 [2.28, 3.72]
7.2 Quality of life	1		Mean Difference (IV, Random, 95% CI)	Subtotals only
7.2.1 PODCI - pain/comfort	1	50	Mean Difference (IV, Random, 95% CI)	3.70 [2.97, 4.43]
7.2.2 PODCI - happiness	1	50	Mean Difference (IV, Random, 95% CI)	-0.70 [-1.42, 0.02]
7.2.3 PedsQL - physical	1	50	Mean Difference (IV, Random, 95% CI)	-4.20 [-4.93, -3.47]
7.2.4 PedsQL - psychosocial	1	50	Mean Difference (IV, Random, 95% CI)	0.60 [-0.12, 1.32]
7.2.5 PedsQL - total score	1	50	Mean Difference (IV, Random, 95% CI)	1.80 [1.07, 2.53]

Analysis 7.1. Comparison 7: PFOs versus shoes in symptomatic flat feet, Outcome 1: Function



Analysis 7.2. Comparison 7: PFOs versus shoes in symptomatic flat feet, Outcome 2: Quality of life



Comparison 8. CFOs versus shoes in DCD flat feet

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
8.1 Gait parameters	1		Other data	No numeric data
8.1.1 six minute walk test (6MWT; m)	1		Other data	No numeric data
8.1.2 cadence (steps/min.)	1		Other data	No numeric data
8.1.3 double support (%)	1		Other data	No numeric data
8.1.4 stride (m)	1		Other data	No numeric data

Analysis 8.1. Comparison 8: CFOs versus shoes in DCD flat feet, Outcome 1: Gait parameters

Gait parameters

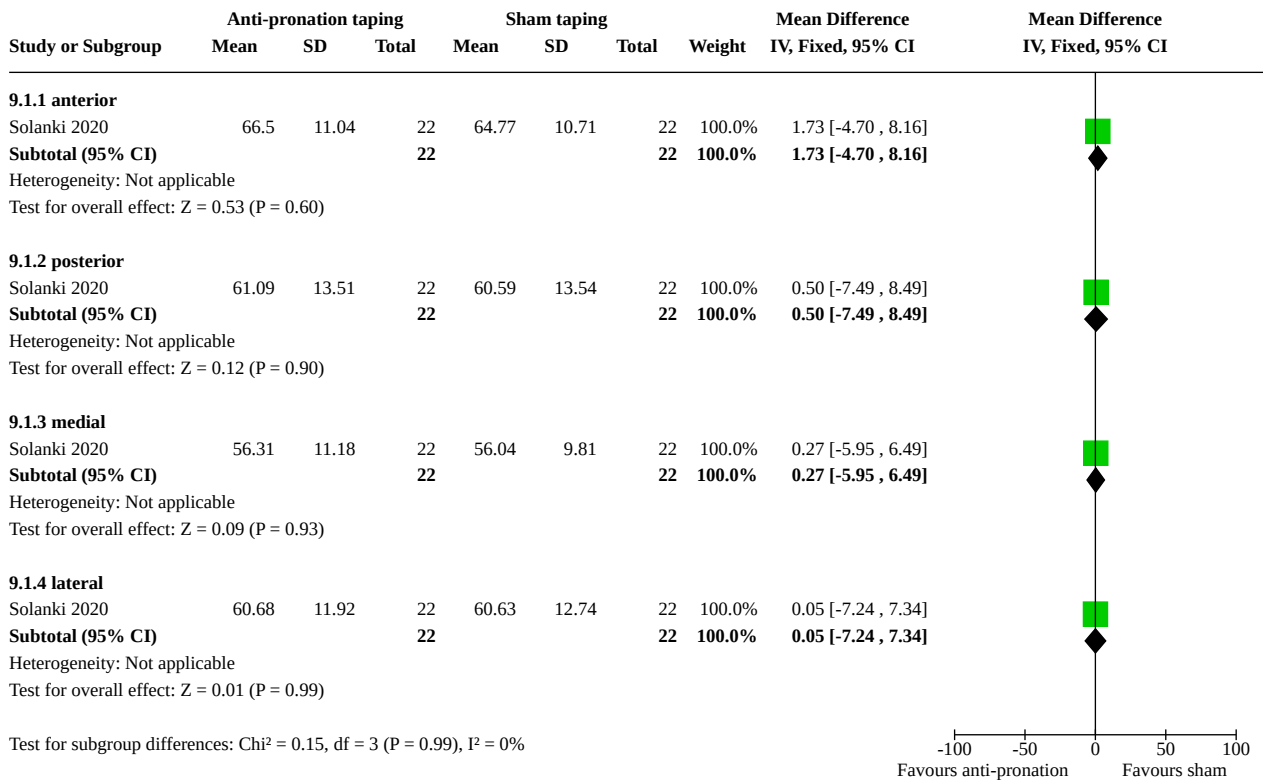
Study	Median	IQR	significance	group (N)
six minute walk test (6MWT; m)				
Morrison 2013	351	312.5 to 433.25	P = 0.43	CFO (9)

	390	375.5 to 437		shoes (5)
cadence (steps/min.)				
Morrison 2013	114.7	108.6 to 124.9	P = 0.019	CFO (9)
	131.9	123.8 to 145		shoes (5)
double support (%)				
Morrison 2013	25	21.7 to 26.9	P = 0.042	CFO (9)
	21.5	19.3 to 22.3		shoes (5)
stride (m)				
Morrison 2013	107.5	102.2 to 122.3	P = 0.23	CFO (9)
	102	95.2 to 111.9		shoes (5)

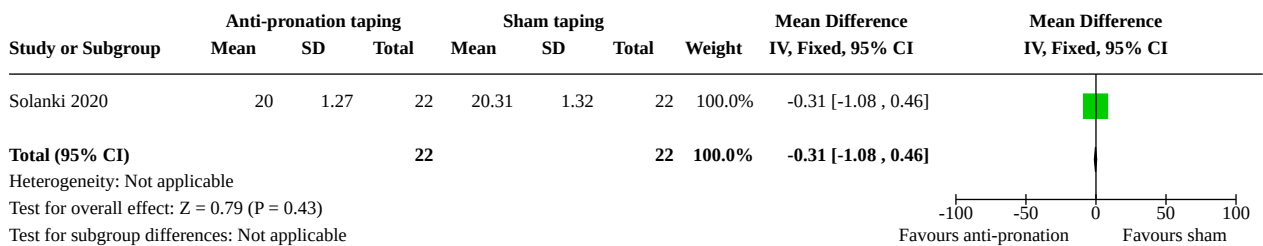
Comparison 9. Anti-pronation taping versus sham taping

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
9.1 Balance test (SEBT) at 4 weeks	1		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
9.1.1 anterior	1	44	Mean Difference (IV, Fixed, 95% CI)	1.73 [-4.70, 8.16]
9.1.2 posterior	1	44	Mean Difference (IV, Fixed, 95% CI)	0.50 [-7.49, 8.49]
9.1.3 medial	1	44	Mean Difference (IV, Fixed, 95% CI)	0.27 [-5.95, 6.49]
9.1.4 lateral	1	44	Mean Difference (IV, Fixed, 95% CI)	0.05 [-7.24, 7.34]
9.2 Agility test (Illinois Agility Test) at 4 weeks	1	44	Mean Difference (IV, Fixed, 95% CI)	-0.31 [-1.08, 0.46]
9.3 Vertical jump height (cm)	1	44	Mean Difference (IV, Fixed, 95% CI)	0.37 [-1.37, 2.11]

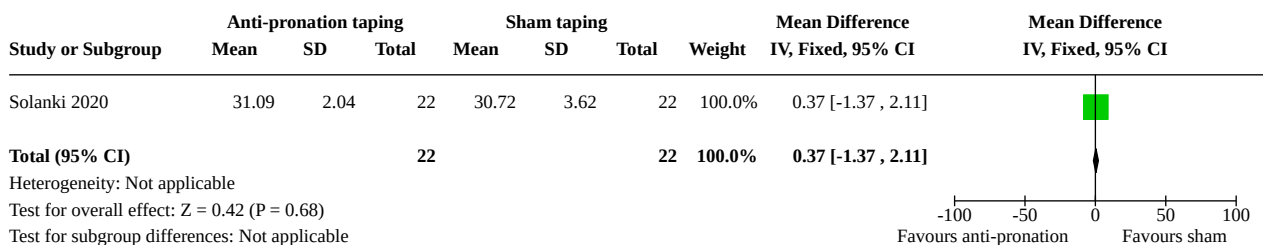
Analysis 9.1. Comparison 9: Anti-pronation taping versus sham taping, Outcome 1: Balance test (SEBT) at 4 weeks



Analysis 9.2. Comparison 9: Anti-pronation taping versus sham taping, Outcome 2: Agility test (Illinois Agility Test) at 4 weeks



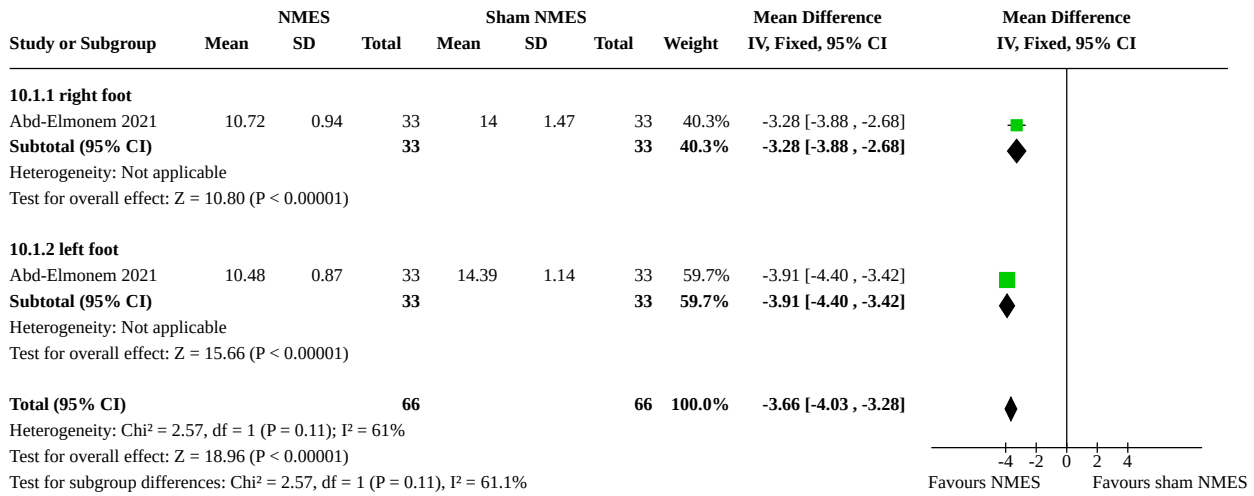
Analysis 9.3. Comparison 9: Anti-pronation taping versus sham taping, Outcome 3: Vertical jump height (cm)



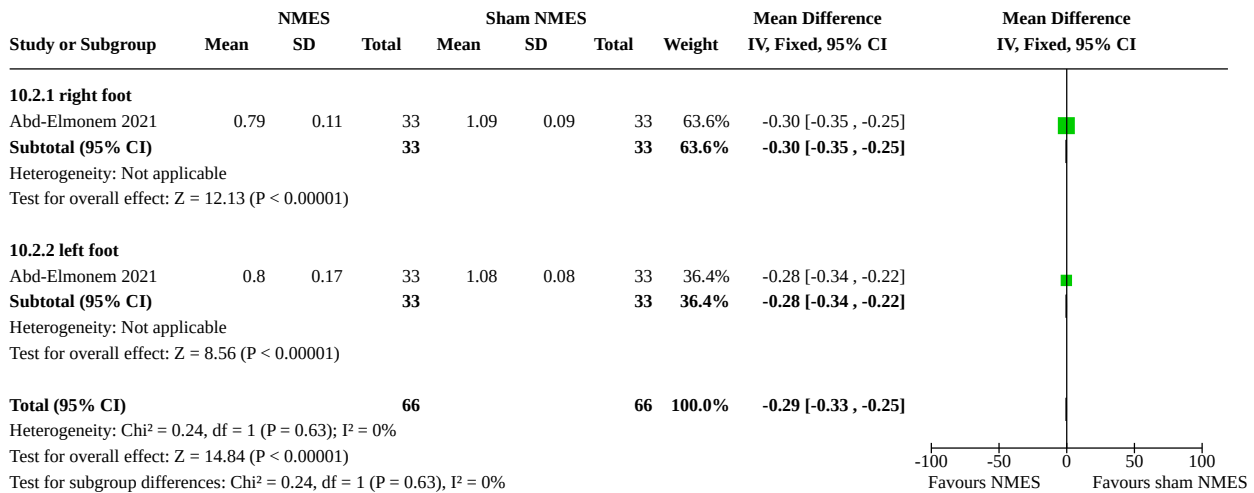
Comparison 10. Neuromuscular electrical stimulation (NMES) versus sham NMES

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
10.1 Navicular height (mm)	1	132	Mean Difference (IV, Fixed, 95% CI)	-3.66 [-4.03, -3.28]
10.1.1 right foot	1	66	Mean Difference (IV, Fixed, 95% CI)	-3.28 [-3.88, -2.68]
10.1.2 left foot	1	66	Mean Difference (IV, Fixed, 95% CI)	-3.91 [-4.40, -3.42]
10.2 Staheli's arch index (mm)	1	132	Mean Difference (IV, Fixed, 95% CI)	-0.29 [-0.33, -0.25]
10.2.1 right foot	1	66	Mean Difference (IV, Fixed, 95% CI)	-0.30 [-0.35, -0.25]
10.2.2 left foot	1	66	Mean Difference (IV, Fixed, 95% CI)	-0.28 [-0.34, -0.22]
10.3 Calcaneal inclination angle (degrees)	1	132	Mean Difference (IV, Fixed, 95% CI)	-2.64 [-3.99, -1.29]
10.3.1 right foot	1	66	Mean Difference (IV, Fixed, 95% CI)	-2.87 [-5.06, -0.68]
10.3.2 left foot	1	66	Mean Difference (IV, Fixed, 95% CI)	-2.50 [-4.22, -0.78]
10.4 Talus second metatarsal angle (degrees)	1	132	Mean Difference (IV, Fixed, 95% CI)	-2.53 [-3.04, -2.02]
10.4.1 right foot	1	66	Mean Difference (IV, Fixed, 95% CI)	-2.45 [-3.25, -1.65]
10.4.2 left foot	1	66	Mean Difference (IV, Fixed, 95% CI)	-2.58 [-3.24, -1.92]
10.5 Talo-navicular coverage angle	1	132	Mean Difference (IV, Fixed, 95% CI)	-1.32 [-1.87, -0.77]
10.5.1 right foot	1	66	Mean Difference (IV, Fixed, 95% CI)	-1.91 [-2.82, -1.00]
10.5.2 left foot	1	66	Mean Difference (IV, Fixed, 95% CI)	-0.98 [-1.67, -0.29]

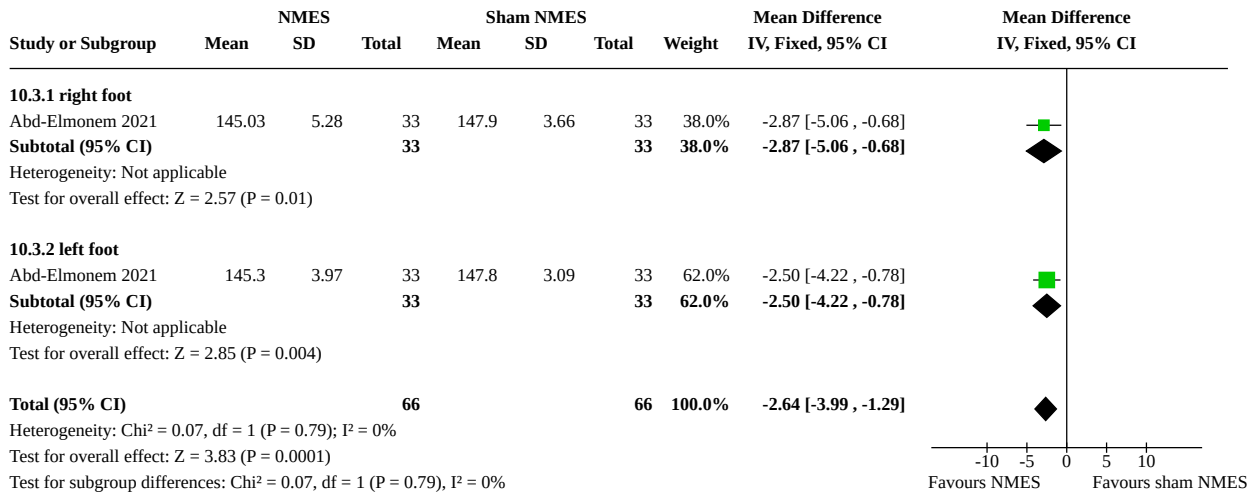
Analysis 10.1. Comparison 10: Neuromuscular electrical stimulation (NMES) versus sham NMES, Outcome 1: Navicular height (mm)



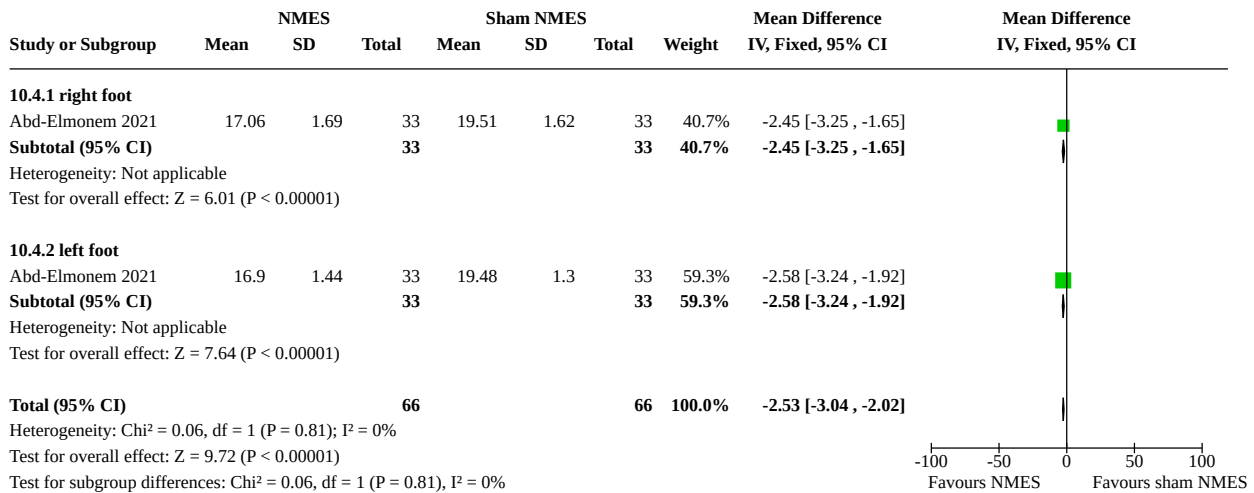
Analysis 10.2. Comparison 10: Neuromuscular electrical stimulation (NMES) versus sham NMES, Outcome 2: Staheli's arch index (mm)



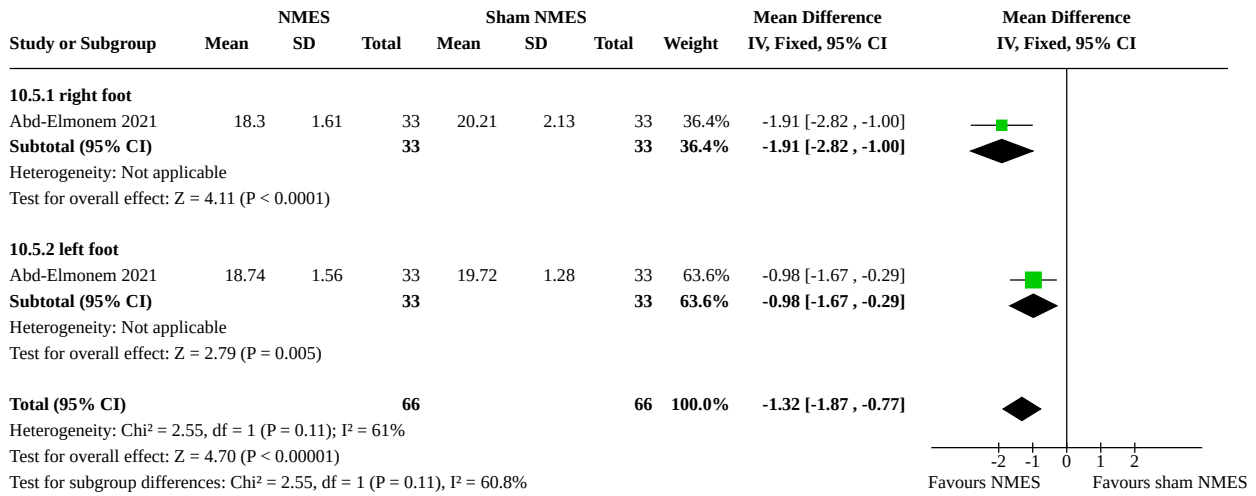
Analysis 10.3. Comparison 10: Neuromuscular electrical stimulation (NMES) versus sham NMES, Outcome 3: Calcaneal inclination angle (degrees)



Analysis 10.4. Comparison 10: Neuromuscular electrical stimulation (NMES) versus sham NMES, Outcome 4: Talus second metatarsal angle (degrees)



Analysis 10.5. Comparison 10: Neuromuscular electrical stimulation (NMES) versus sham NMES, Outcome 5: Talo-navicular coverage angle



ADDITIONAL TABLES
Table 1. Study characteristics of the 16 included trials

Study Country	Follow-up time	Baseline sample size	Age (SD)	Final sample size (% of baseline)	Intervention	Outcome measures
Flexible asymptomatic flat feet in healthy children (9 studies)						
Wenger 1989 USA	3 years	131	1 to 6 years	98 (75%)	Shoe: N = 28 Heel cup: N = 27 UCBL: N = 22 Control: N = 21	X-ray Clinical photos
Gould 1989 USA	5 years	125	11 to 14 months	52 (42%)	SL shoe: N = 25 SL shoe/ cookie: N = 10 Ortho shoes: N = 7 Ortho/mla: N = 10	X-ray Pedotopography Clinical assessment
Whitford 2007 Australia	1 year	178	7 to 11 years	160 (90%)	CFO: N = 59 FO: N = 59 Control: N = 60	Pain SPPC Motor skills VO ² max
Asgaonkar 2012 India	1 year	80	5 to 15 years	60 (75%)	Valgus insole: N = 30 Control: N = 30	Pain (VAS) Physical cost (HR) Gait (step parameters)
Kanatli 2016 Turkey	2 to 5 years	45	17 to 72 months (average 39.5 months)	45 (100%)	Orthotic shoes: N = 21 Control: N = 24	X-ray Laxity AI
Ahn 2017 Korea	1 year	40	10.14 years (4.99)	40 (100%)	TCFO: N = 20 RFO: N = 20	X-ray RCSP
Khamooshi 2017 Iran	8 weeks	60	9 to 13 years	60 (100%)	Foot exercises: N = 20 Foot/core exercises: N = 20 Control: N = 20	Pedoscope Staheli AI ND Tiptoe/mla
Ja-farnezhadgero 2018 Iran	4 months	30	8 to 12 years	30	CFO: N = 15 Sham insole: N = 15	Gait kinematic Kinetic parameters

Table 1. Study characteristics of the 16 included trials (Continued)

Solanki 2020 India	4 weeks	44	approximately 13 to 14 years	44	Conventional exercises + Faradic foot bath + rigid taping: N = 22 Conventional exercises + Faradic foot bath + sham tape: N = 22	SEBT VJH IAT
Abd-Elmonem 2021 Egypt	4 months	72	7 to 12 years	66	Corrective exercises + NMES: N = 36 Corrective exercises + sham NM ES: N = 36	Staheli AI ND x-ray
Children with juvenile idiopathic arthritis and foot pain (2 studies)						
Powell 2005 USA	3 months	48	5 to 19 years	40 (83%)	CFO: N = 15 Neoprene inserts: N = 12 Sports shoe: N = 13	Pain (VAS) PedsQL Timed walk FFI
Coda 2014 UK	0, 3, 6 months	60	10 to 11 years (3.5)	60 (100%)	CPFO: N = 31 PFO: N = 29	VAS PedsQL
Flexible flat feet in children with foot pain (1 study)						
Hsieh 2018 Taiwan	12 weeks	52	6 to 7 years	50	PFO: N = 24 Control: N = 26	Physical activity Function (PODCI) Psychometric (PODCI, HRQoL)
Flexible flat feet in children with foot pain (immediate effects only; 1 study)						
Bok 2016 South Korea	immediate	21	8 to 13 years (average 9.9 years)	21 (100%)	0° inverted CFO/15° inverted CFO/30° inverted CFO: N = not specified Shoes only (usual): N = not specified	Pedar - peak pressure, max. force, contact area
Flexible flat feet in children without foot pain (immediate effects only; 1 study)						
Aboutorabi 2013 Iran	immediate	50 (30 flat feet: 20 controls)	7.76 years (1.4)	50 (100%)	Shoes + CFO/ Medical shoes/Bare-foot: N = 30 Control (no flat feet): N = 20	Gait - Step - length, width, symmetry Velocity CoP

Table 1. Study characteristics of the 16 included trials (Continued)

Flexible flat feet in children with developmental co-ordination disorder (1 study)						
Morrison 2013 UK	7 weeks	22	6 to 11 years	14 (64%)	CFO: N = 9 Control: N = 5	6-minute walk Gait rite

Abbreviations:

ADRs: adverse reactions;

AI: arch index;

CFO: customised/bespoke foot orthoses;

CoP: centre of pressure;

FF: flat feet;

FO: foot orthoses;

HR: heart rate;

HRQoL: health-related quality of life

IAT: Illinois Agility test;

JIA: juvenile idiopathic arthritis;

ND: navicular drop;

NMES: neuromuscular electrical stimulation

NS: not significant;

PedsQL: Pediatric quality of life inventory;

PFO: prefabricated foot orthoses;

PODCI: Paediatric outcome data collection instrument

RCSP: resting calcaneal stance position;

RFO: rigid FO;

SEBT: start excursion balance test;

SL: straight last (shoe);

SPPC: self perception profile;

TCFO: talus control FO;

UCBL: University of California Biomechanics Laboratory heel cup orthosis;

Table 1. Study characteristics of the 16 included trials (Continued)

VAS: visual analogue score;

VJH: vertical jump height.

Prefabricated foot orthoses definition

A prefabricated foot orthosis is an in-shoe medical device that is not made from an individual scan, cast, or mould of the foot. This generic device is intended to alter the magnitudes and temporal patterns of the reaction acting on the plantar aspect of the foot and normalise foot and lower extremity function; decreasing abnormal loading forces on the structural components of the foot and lower extremity during weight-bearing and related activity.

Customised prefabricated foot orthoses definition

A modified version of a basic generic device, which is initially mass produced, and then specifically modified for the foot and gait requirements of an individual child. The modifications are usually added by the treating clinician, and may include: additional arch fill, varus or valgus wedges, and topcovers.

Custom foot orthoses definition

A bespoke foot orthosis is an individually customised in-shoe medical device that is made from an individual scan, cast, or mould of the foot. The design is prescribed by a qualified healthcare professional to alter the magnitudes and temporal patterns of the reaction forces acting on the plantar aspect of the foot, in order to allow more normal foot and lower extremity function, and to decrease pathologic loading forces on the structural components of the foot and lower extremity during weight-bearing and related activity.

Table 2. Outcome matrix per trial group comparison

Diagnosis	Pain	Function	HRQoL	Treatment success	Withdrawals	Adverse events	Serious adverse events
1. CFO versus shoes							
asymptomatic flat feet	Whitford 2007 – post hoc subgroup (% pain)	Whitford 2007 – VO ² max, motor skills	NR	NR	Wenger 1989; Whitford 2007	NR	NR
JIA	Powell 2005 – VAS	Powell 2005 – timed walk	Powell 2005 – FFI	Powell 2005	Powell 2005	Powell 2005 – none	NR
DCD	NR	Morrison 2013 – 6MWT	NR	Morrison 2013	Morrison 2013	NR	NR
2. PFO versus shoes							
asymptomatic flat feet	Whitford 2007 post hoc subgroup (% pain)	Whitford 2007 – VO ² max, motor skills	NR	Asgaonkar 2012; Gould 1989;	Asgaonkar 2012; Gould 1989; Wenger 1989; Whitford 2007	NR	NR
symptomatic	NR	Hsieh 2018	Hsieh 2018 – PODCI	Hsieh 2018	Hsieh 2018	NR	NR

Table 2. Outcome matrix per trial group comparison (Continued)

flat feet							
JIA	Powell 2005 – VAS	Powell 2005 – timed walk	Powell 2005 – FFI	Powell 2005	Powell 2005	Powell 2005 – none	NR
3. CFO versus PFO							
asymptomatic	Whitford 2007 – post hoc subgroup (% pain)	Whitford 2007 – VO ² max, motor skills	NR	NR	Wenger 1989; Whitford 2007	NR	NR
flat feet							
JIA	Coda 2014; Powell 2005 – VAS	Powell 2005 – timed walk	Coda 2014; Powell 2005 – PedsQL	Coda 2014; Powell 2005;	Coda 2014; Powell 2005;	Coda 2014 – NR Powell 2005 – none	NR

6MWT: 6-minute walk test; CFO: custom foot orthoses; DCD: developmental co-ordination disorder; HRQoL: health-related quality of life; JIA: juvenile idiopathic arthritis; HR: heart rate; NR: not reported; PedsQL: Pediatric quality of life inventory; PFO: prefabricated foot orthoses; PODCI: Paediatric outcome data collection instrument; FFI: Foot Function index; VAS: visual analogue scale

Table 3. Shoes used within the trials

Study ID	Control shoe	Comparator shoes
Asymptomatic flat feet		
Wenger 1989	usual shoes	corrective shoes, usual shoes + Helfet heel cups, usual shoes + UCBL CFO
Gould 1989	straight last shoes	- straight last shoes plus longitudinal arch cookies - orthopaedic shoes with long counters, solid shanks, Thomas heels, and 0.312 cm inside heel wedges - orthopaedic shoes with long counters, solid shanks, Thomas heels, and 0.312 cm inside heel wedges, with supplemental thin longitudinal arch support
Whitford 2007	usual shoes	none (PFO, CFO)
Asgaonkar 2012	usual shoes	none (valgus insole)
Kanatli 2016	usual shoes	corrective shoes, i.e. custom-made orthopaedic shoes that had 0.5 to 0.9cm longitudinal arch support and 3 to 4 mm heel wedges
Ahn 2017	usual shoes	none (2 CFO types)
Khamooshi 2017	usual shoes	none (foot, core exercises)
Jafarnezhadgero 2018	New Balance 759 (trainers)	New Balance 759 (trainers)
Solanki 2020	not stated	not stated
Abd-Elmonem 2021	not stated	not stated
Symptomatic flat feet		
Hsieh 2018	usual shoes (encouraged to wear at least 5 hours daily)	usual shoes (encouraged to wear at least 5 hours daily)
JIA		
Powell 2005	new supportive athletic shoes with a medial longitudinal arch support and shock absorbing soles (cross-training type shoes)	all children, regardless of intervention, received new athletic shoes at beginning of the study
Coda 2014	usual shoes	none (PFO, CPFO)
DCD		
Morrison 2013	usual shoes	none (CFO)
Immediate effects studies		
Bok 2016	usual shoes	none (3 inverted CFOs)

Table 3. Shoes used within the trials (Continued)

[Aboutorabi 2013](#) no shoes (bare feet) medical shoes, regular shoes (with FO)

CFO: customised foot orthoses, CPFO: customised prefabricated foot orthoses; DCD: developmental co-ordination disorder; FO: foot orthoses; JIA: juvenile idiopathic arthritis; PFO: prefabricated foot orthoses; UCBL: University of California Biomechanics Laboratory heel cup orthosis

Table 4. Prefabricated foot orthoses versus control on function and pain outcomes at 12 months

Outcome Measure	No of participants	Prefabricated orthoses	Controls	P value	Effect size
Physical cost (mean (SD))	Whitford 2007 = 160	45.10 (4.88)	44.95 (3.81)	P = 0.899	MD 0.15, 95% CI -1.51 to 1.81
- Whitford 2007 (VO ² max)		0.20 (0.06)	0.26 (0.12)	P = 0.0264	
- Asgaonkar 2012 (HR)		Asgaonkar 2012 = 60			
					MD -0.06, 95% CI -0.11 to -0.01
Pain (mean (SD))	60	0.64 (1.09)	4.33 (2.58)	P < 0.0001	MD -3.69, 95% CI -4.60 to -2.78
- Asgaonkar 2012 (VAS, mean (SD))					
Pain (numbers (%))	160	36/54 (67%)	41/52 (79%)	P = 0.56	RR 0.85, 95% CI 0.67 to 1.07
- Whitford 2007 (% without pain)					

[Asgaonkar 2012](#) reported improvement in both pain and physical cost of children treated with prefabricated orthoses at 12 months versus the control group

[Whitford 2007](#) found no difference between groups

CI: confidence interval; **MD:** mean difference; **RR:** risk ratio

Table 5. Shoes versus control on x-ray outcomes at 3 years

Outcome Measure	No of participants	Corrective shoes	Controls	P value	Effect size
Talo-horizontal x-ray change (mean (SD))	98	6.47 (0.59)	5.48 (0.71)	P > 0.4	-0.16 (-0.44 - 0.16)
- Wenger 1989	45	0.17	0.13	P = 0.19	
- Kanatli 2016					
Talo-1st metatarsal x-ray change (mean (SD))	98	6.80 (0.7)	5.78 (0.83)	P > 0.5	-0.50 (-1 - (-0.02))
- Wenger 1989	45	0.45	0.46	P = 0.72	
- Kanatli 2016					
Talocalcaneal (AP) x-ray change (mean (SD))	98	7.36 (0.78)	4.50 (0.91)	P > 0.5	-0.12 (-0.05 - 0.20)
- Wenger 1989	45	0.13	0.23	P = 0.09	
- Kanatli 2016					

Wenger showed positive correlation between all radiographic parameters between initial and changed angles over three years (P < 0.001). Both studies showed that the measured change in x-ray angles was the same between treatment (shoes) and control groups after three years.

APPENDICES

Appendix 1. CENTRAL, in the Cochrane Library search

Search Name: Cochrane A028-R limit date 2009 - 300720

Date Run: 02/09/2021 11:00:04

ID Search Hits

- #1 MeSH descriptor: [Flatfoot] explode all trees 60
- #2 flat next foot*:ti,ab 81
- #3 flatfoot:ti,ab 53
- #4 flat next feet:ti,ab 103
- #5 flatfeet:ti,ab 5
- #6 pes next planus:ti,ab 44
- #7 painful next foot:ti,ab 26
- #8 pes next planovalgus:ti,ab 4
- #9 "posterior tibial tendon dysfunction":ti,ab 9
- #10 subtalar:ti,ab 163
- #11 sub* next talar:ti,ab 4
- #12 calcane*:ti,ab 723
- #13 heel next bone*:ti,ab 35
- #14 medical arch*:ti,ab 1702
- #15 (#1 OR #2 OR #3 OR #4 OR #5 OR #6 OR #7 OR #8 OR #9 OR #10 OR #11 OR #12 OR #13 OR #14) 2715
- #16 MeSH descriptor: [Musculoskeletal Diseases] explode all trees 43202
- #17 MeSH descriptor: [Neuromuscular Diseases] explode all trees 10781
- #18 MeSH descriptor: [Nervous System Diseases] explode all trees 87486
- #19 ehlers-danlos:ti,ab 51
- #20 "down* syndrome":ti,ab 747
- #21 trisomy:ti,ab 215
- #22 mongolism:ti,ab 2
- #23 "inflammatory arthritis":ti,ab 439
- #24 juvenile near/3 arthritis:ti,ab 839
- #25 (#16 OR #17 OR #18 OR #19 OR #20 OR #21 OR #22 OR #23 OR #24) 124705
- #26 MeSH descriptor: [Diabetes Mellitus, Type 1] explode all trees 5764
- #27 (diabet* or IDDM):ti,ab 95002
- #28 MeSH descriptor: [Diabetes Insipidus] explode all trees 73

- #29 mellitus:ti,ab 28759
- #30 diabet* next insipidus:ti,ab 123
- #31 diabet* next mellitus:ti,ab 28693
- #32 (#26 OR #27) 95299
- #33 # 32 NOT (#26 OR #29 OR #31) 127788
- #34 (#32 OR #33) 217218
- #35 MeSH descriptor: [Joint Instability] explode all trees 777
- #36 ligament* next laxity:ti,ab 48
- #37 pronat*:ti,ab 521
- #38 malalignment:ti,ab 297
- #39 (#35 OR #36 OR #37 OR #38) 1626
- #40 (#25 OR #34 OR #39) 329079
- #41 (#15 AND #40) 875
- #42 child:ti,ab 122707
- #43 children:ti,ab 122705
- #44 childhood:ti,ab 15011
- #45 infant:ti,ab 16230
- #46 teenage:ti,ab 481
- #47 adolescen*:ti,ab 29258
- #48 paediatric:ti,ab 31799
- #49 pediatric:ti,ab 31807
- #50 (#42 OR #43 OR #44 OR #45 OR #46 OR #47 OR #48 OR #49) 165257
- #51 (#41 AND #50) with Publication Year from 2009 to 2020, with Cochrane Library publication date Between Jul 2009 and Jul 2020, in Trials 57
- #52 (#41 AND #50) with Publication Year from 2020 to 2021, with Cochrane Library publication date Between Jul 2020 and Sep 2021, in Trials 9

Appendix 2. MEDLINE search

Database: Ovid MEDLINE(R) <1946 to September 01, 2021>

Search Strategy:

1 exp flatfoot/ (2420)

2 flat foot\$.mp. (492)

3 flatfoot\$.mp. (2639)

4 flat feet.mp. (314)

5 flatfeet.mp. (140)

6 pes planus.mp. (444)

Foot orthoses for treating paediatric flat feet (Review)

Copyright © 2022 The Cochrane Collaboration. Published by John Wiley & Sons, Ltd.

- 7 painful foot.mp. (139)
- 8 pes planovalgus.mp. (178)
- 9 posterior tibial tendon dysfunction.mp. (407)
- 10 subtalar.mp. (3432)
- 11 (sub\$ adj talar).mp. (51)
- 12 calcane\$.mp. (12164)
- 13 heel bone\$.mp. (192)
- 14 medical arch\$.mp. (4)
- 15 or/1-14 (16654)
- 16 exp musculoskeletal diseases/ (1132028)
- 17 exp neuromuscular diseases/ (321208)
- 18 exp nervous system diseases/ (2639188)
- 19 ehlers danlos.mp. (3950)
- 20 down\$ syndrome.mp. (29398)
- 21 trisomy.mp. (20802)
- 22 mongolism.mp. (1563)
- 23 inflammatory arthritis.tw. (4517)
- 24 (juvenile adj3 arthritis).tw. (9551)
- 25 or/16-24 (3553109)
- 26 exp diabetes mellitus, type 1/ (79337)
- 27 (diabet\$ or IDDM).tw. (589388)
- 28 26 or 27 (596792)
- 29 exp diabetes insipidus/ (8046)
- 30 mellitus.tw. (176625)
- 31 29 not (26 or 30) (7373)
- 32 (diabet\$ adj (insipidus not mellitus)).tw. (7731)
- 33 31 or 32 (9648)
- 34 28 not 33 (588951)
- 35 joint instability.sh. (21632)
- 36 ligament\$ laxity.mp. (806)
- 37 pronat\$.mp. (6019)
- 38 malalignment.mp. (5292)
- 39 or/35-38 (32599)
- 40 or/25,34,39 (4079777)
- 41 15 and 40 (10302)

- 42 randomized controlled trial.pt. (542072)
- 43 controlled clinical trial.pt. (94342)
- 44 randomized.ab. (461416)
- 45 placebo.ab. (200637)
- 46 drug therapy.fs. (2367832)
- 47 randomly.ab. (309518)
- 48 trial.ab. (488467)
- 49 groups.ab. (1911898)
- 50 or/42-49 (4665240)
- 51 (animals not (humans and animals)).sh. (4847355)
- 52 50 not 51 (4000598)
- 53 41 and 52 (1452)
- 54 limit 53 to ("infant (1 to 23 months)" or "preschool child (2 to 5 years)" or "child (6 to 12 years)" or "adolescent (13 to 18 years)") (476)
- 55 child.mp. (2156988)
- 56 children.mp. (987895)
- 57 childhood.mp. (243050)
- 58 infant\$.mp. (1278871)
- 59 teenag\$.mp. (20079)
- 60 adolescen\$.mp. (2155197)
- 61 paediatric.mp. (56044)
- 62 pediatric.mp. (285434)
- 63 or/55-62 (4022120)
- 64 53 and 6 (70)
- 65 54 or 64 (512)
- 66 limit 65 to dt=20090701-20200728 (236)
- 67 limit 66 to dt=20200730-20210901 (16)

Appendix 3. Embase search strategy

Database: Embase Classic+Embase <1947 to 2021 September 01>

Search Strategy:

-
- 1 Flatfoot/ (4297)
- 2 flat foot\$.mp. (1085)
- 3 flatfoot\$.mp. (4756)
- 4 flat feet.mp. (638)
- 5 flatfeet.mp. (234)

- 6 pes planus.mp. (882)
- 7 painful foot.mp. (239)
- 8 pes planovalgus.mp. (390)
- 9 posterior tibial tendon dysfunction.mp. (604)
- 10 subtalar.mp. (5576)
- 11 (sub\$ adj talar).mp. (85)
- 12 calcane\$.mp. (18956)
- 13 heel bone\$.mp. (324)
- 14 medical arch\$.mp. (3)
- 15 or/1-14 (26512)
- 16 exp Musculoskeletal Disease/ (2653609)
- 17 exp Neuromuscular Disease/ (222845)
- 18 exp Neurologic Disease/ (4051261)
- 19 ehlers danlos.mp. (6829)
- 20 down\$ syndrome.mp. (43417)
- 21 trisomy.mp. (36587)
- 22 mongolism.mp. (2063)
- 23 inflammatory arthritis.tw. (10525)
- 24 (juvenile adj3 arthritis).tw. (18516)
- 25 or/16-24 (6050059)
- 26 Insulin Dependent Diabetes Mellitus/ (124599)
- 27 (diabet\$ or IDDM).tw. (1059361)
- 28 26 or 27 (1074667)
- 29 exp Diabetes Insipidus/ (17717)
- 30 mellitus.tw. (311351)
- 31 29 not (26 or 30) (16444)
- 32 (diabet\$ adj (insipidus not mellitus)).tw. (12278)
- 33 31 or 32 (17980)
- 34 28 not 33 (1061737)
- 35 joint instability.sh. (12223)
- 36 ligament\$ laxity.mp. (1257)
- 37 pronat\$.mp. (8728)
- 38 malalignment.mp. (6085)
- 39 or/35-38 (27594)
- 40 or/25,34,39 (6917639)

- 41 15 and 40 (21433)
- 42 random\$.ti,ab. (1712450)
- 43 factorial\$.ti,ab. (42419)
- 44 (crossover\$ or cross over\$ or cross-over\$).ti,ab. (115459)
- 45 placebo\$.ti,ab. (335897)
- 46 (doubl\$ adj blind\$).ti,ab. (228318)
- 47 (singl\$ adj blind\$).ti,ab. (27619)
- 48 assign\$.ti,ab. (434599)
- 49 allocat\$.ti,ab. (172371)
- 50 volunteer\$.ti,ab. (277903)
- 51 crossover procedure.sh. (68322)
- 52 double blind procedure.sh. (189697)
- 53 randomized controlled trial.sh. (676185)
- 54 single blind procedure.sh. (43642)
- 55 or/42-54 (2581294)
- 56 exp animal/ or nonhuman/ or exp animal experiment/ (31561864)
- 57 exp human/ (24025004)
- 58 56 and 57 (24025004)
- 59 56 not 58 (7536860)
- 60 55 not 59 (2270319)
- 61 41 and 60 (1208)
- 62 limit 61 to (infant or child or preschool child <1 to 6 years> or school child <7 to 12 years> or adolescent <13 to 17 years>) (173)
- 63 child.mp. (2719502)
- 64 children.mp. (1537907)
- 65 childhood.mp. (477947)
- 66 infant\$.mp. (1065321)
- 67 teenag\$.mp. (32051)
- 68 adolescen\$.mp. (1863177)
- 69 paediatric.mp. (112907)
- 70 pediatric.mp. (520089)
- 71 or/63-70 (4483036)
- 72 61 and 71 (188)
- 73 62 or 72 (188)
- 74 limit 73 to dd=20090701-20200728 (99)
- 75 limit 74 to dd=20200730-20210901 (5)

Appendix 4. CINAHL search strategy

S40 S26 and S39

S39 S27 or S28 or S29 or S30 or S31 or S32 or S33 or S34 or S35 or S36 or S37 or S38

S38 TI Allocat* random* or AB Allocat* random*
 S37 (MH "Quantitative Studies")

S36 (MH "Placebos")

S35 TI Placebo* or AB Placebo*

S34 TI Random* allocat* or AB Random* allocat*
 S33 (MH "Random Assignment")

S32 TI Randomi?ed control* trial* or AB Randomi?ed control* trial*

S31 AB singl* blind* or AB singl* mask* or AB doub* blind* or AB doubl* mask* or AB trebl* blind* or AB trebl* mask* or AB tripl* blind* or AB tripl* mask*

S30 TI singl* blind* or TI singl* mask* or TI doub* blind* or TI doubl* mask* or TI trebl* blind* or TI trebl* mask* or TI tripl* blind* or TI tripl* mask*

S29 TI clinical* trial* or AB clinical* trial*
 S28 PT clinical trial

S27 (MH "Clinical Trials+")

S26 S5 and S25

S25 S10 or S19 or S24

S24 S20 or S21 or S22 or S23

S23 TI malalignment or AB malalignment
 S22 TI pronat* or AB pronat*

S21 TI ligament* laxity or AB ligament* laxity
 S20 (MH "Joint Instability+")

S19 S13 not S18

S18 S16 or S17

S17 TI diabet* insipidus or AB diabet* insipidus not TI diabet* mellitus or AB diabet* mellitus

S14 not (S11 or S15)

S15 ti mellitus or ab mellitus

S14 (MH "Diabetes Insipidus")
 S13 S11 or S12

S12 TI diabet* or AB diabet* or TI IDDM or AB IDDM
 S11 (MH "Diabetes Mellitus, Insulin-Dependent")

S10 S6 or S7 or S8 or S9

S9 TI ehlers-danlos or AB ehlers-danlos or TI down* syndrome or AB down* syndrome or TI trisomy or AB trisomy or TI mongolism or AB mongolism or TI inflammatory arthritis or AB inflammatory arthritis or TI juvenile N3 arthritis or AB juvenile N3 arthritis

S8 (MH "Nervous System Diseases+") S7 (MH "Neuromuscular Diseases+")

S6 (MH "Musculoskeletal Diseases+") S5 S1 or S2 or S3 or S4

S4 TI subtalar or AB subtalar or TI sub* adj talar or AB sub* adj talar or TI calcane* or AB calcane* or TI heel bone* or AB heel bone* or TI medical arch* or AB medical arch* Search modes - Boolean/Phrase S3 TI pes planus or AB pes planus or TI painful foot or AB painful foot

or TI pes planovalgus or AB pes planovalgus or TI posterior tibial tendon dysfunction or AB posterior tibial tendon dysfunction Search S2
TI flat foot* or AB flat foot* or TI flatfoot* or AB flatfoot* or TI flat feet or AB flat feet or TI flatfeet or AB flatfeet

S1 (MH "Flatfoot")

Appendix 5. Dissertation abstracts search

(flatfoot) OR (flat foot*) OR (flatfoot*) OR (flat feet) OR (flatfeet) OR (pes planus) OR (painful foot) in Citation and Abstract

FEEDBACK

Feedback, June 2011

Summary

Date of Submission: 20 June 2011

Name: Alan Cooper BSc

Personal Description: Occupation UK Podiatrist

Feedback: When one of my daughters was born 32 years ago, the first thing I noticed (as soon as she was delivered) was her left foot laying flat against the tibia, and fibula - so much so that I observed a clear imprint of the outline of the bones. At the time, I was in my 2nd year at Podiatry college. The condition was undoubtedly T. calcaneovalgus, and in my opinion, severe. The foot was dorisflexed, abducted, which in my view as an adult it would have exhibited severe abnormal pronation, with an everted calcaneum, a flat arch with the whole of the mid-tarsal joint contacting the ground plantar grade. An orthopaedic surgeon was asked to look at her feet, and as he felt all the components of the foot were there, we were referred to a physiotherapist for treatment. The treatment was of course manipulation of the foot, with splinting for several months. The splint was removed six or seven times a day, and the foot manipulated. Today, looking at my daughter's feet, there is no difference between them, and both stand correctly in their neutral position. Interestingly, on our last appointment with the physiotherapist, she had asked a father to bring in his child with the same condition, who was born at the same time. The child had not received manipulation, and as a consequence, there was no real change in the foot's appearance from birth.

Over my 30 years of practice, I have seen many conditions of pes planus, and wondered if there was a relationship between this and untreated mild, moderate, or severe T. calcaneovalgus: does this go unseen, and untreated? If it was picked up by the podiatrist at birth, is it possible that the condition of pes planus in many folk could be eliminated? Just recently, a toddler was referred to me with bilateral flat feet: heels everted, feet abducted, with metatarsals, and metatarsal joints appearing to be either close, or contacting the ground. The father had almost a cavoid foot, and the mother's foot was quite normal. Bearing in mind the limited evidence for many treatments, I think I shall recommend small orthosis, together with some manipulation. When I have enquired of old, and older clients in the past, the treatment appeared to be just exercises, which seemed to have made no difference to the adult foot at all.

Submitter agrees with default conflict of interest statement: I certify that I have no affiliations with or involvement in any organisation or entity with a financial interest in the subject matter of my feedback.

Reply

27 June 2011

Dear Mr Cooper

Thank you for your response relating to the Cochrane Review. Dealing with children's feet is an issue. Dr Evans recently described a traffic light system for treating flat feet that has been strongly debated [1-3]. From a clinical viewpoint, identifying a foot problem at a very young age is still based upon clinical expertise and personal choice for management [4]. Unfortunately, this is the lowest form of evidence [5], and further work is needed in this area to ensure that clinicians are using an evidence-base approach to manage flat feet in children.

From the Cochrane Review, we only found reports on the use of foot orthoses for the management of paediatric flat feet [4]. The use of manipulation as a form of treatment was not reported in the literature. However, in a recent critical review [6], we found one article that evaluated a group of children who followed a rehabilitative programme versus a historical group of children who had been treated with insoles and orthopaedic footwear [7]. Over a two-year period, 300 children (mean age was 3.4 years; 184 male, 116 female) with bilateral flexible flatfoot (600 feet) were recruited and underwent a rehabilitative programme for a mean period of 2.8 years. The rehabilitative programme consisted of simple therapeutic exercises, which could be easily learned by both patients and their caregivers. These children were compared to a historical group of children (674 feet) who had been treated in a paediatric department for infantile flexible flatfoot with the use of foot orthoses. The results demonstrated that in the two groups (children treated with rehabilitation and children treated with foot orthoses), the rehabilitative approach seemed to be more effective. The authors suggested that rehabilitation has a marginal influence on the natural history of paediatric valgus flexible flatfoot, even though it plays a role in maintaining good flexibility of the flatfoot, thus limiting functional impairment.

In summary, the use of exercise is commonly reported, but the evidence is limited. Future studies need to further investigate the potential effect of rehabilitative exercises, and foot manipulation, and continue to investigate the efficacy of foot orthoses, and footwear modifications, and consider the effect of other contemporary clinical interventions.

References

1. Evans AM. The flat-footed child - to treat or not to treat: What is the clinician to do? *J Am Podiatr Med Ass* 2008; 98: 386-393.
2. D'Amico JC. The flat-footed child - to treat or not to treat: What is the clinician to do? *J Am Podiatr Med Ass* 2009; 99:267.
3. Bresnahan P. The flat-footed child - to treat or not to treat: What is the clinician to do? *J Am Podiatr Med Ass* 2009; 99: 178.
4. Rome K, Ashford RL, Evans AM. Non-surgical interventions for pediatric pes planus. *Cochrane Database Syst Rev* 2010; 7(CD006311):DOI:10.1002
5. Sackett DL, Straus SE, Richardson WS, Rosenberg W, Haynes RB. *Evidence-based Medicine. How to practice and teach EBM.* 2000. 2nd edition. Churchill Livingstone, UK.
6. Evans AM, Rome K. A review of the evidence for non-surgical interventions for flexible paediatric flat feet. *Eur J Phys Rehabil Med* 2011; 47: 1-2.
7. Riccio I, Gimigliano AE, Gimigliano R, Porpora G, Iolascon G. Rehabilitative treatment in flexible flatfoot: a perspective cohort study *Musculoskeletal Surg* 2009; 93:101-1

Contributors

Professor Keith Rome, Professor Podiatry, AUT University, New Zealand

Dr Angela Evans, Senior Research Fellow, University of South Australia, Australia

Feedback, May 2012

Summary

Date of Submission: 12 May 2012

Name: A. van Heukelum

Personal Description: Occupation General Practice Resident

Feedback: During our residency, all students are required to make an "critically appraised topic", based on a personal case during practice. The problem of painful flat feet or pes planus came to my attention, so I was very pleased to find this review on the subject, and would like to thank you for your clear, insightful review. Clearly, much more research needs to be done to make evidence. I have one question though, about the included article of Powell (2005). As I understood, the inclusion criterium, amongst others, was patients with pes planus. In the short description of the article by Powell (2005), it was stated that it included cases of pes planus with juvenile idiopathic arthritis, although in this article, this was not clear to me. Could you perhaps give more details about why the participants of this study would have pes planus?

Submitter has modified conflict of interest statement:

I certify that I have no affiliations with or involvement in any organization or entity with a financial interest in the subject matter of my feedback.

Reply

Pes planus (or a valgus/pronated rearfoot posture) is by far the most common presentation in juvenile idiopathic arthritis, but hindfoot varus posture can also occur, and is usually associated with significantly reduced joint range of motion. I have included three references that may assist you:

1. Ferrari J. A review of foot deformities seen in juvenile chronic arthritis. *The Foot*, 1998; 8: 193-196.2.
2. Mavidrou A et al. Conservative management of the hindfoot in juvenile chronic arthritis. *The Foot*, 1991; 1; 139-143.3.
3. Hendry G et al. A survey of foot problems in juvenile idiopathic arthritis. *Musculoskeletal Care*, 2008; 6: 221-232.

The Cochrane review team, at the time of reviewing the literature, made a decision to include the Powell (2005) study, based upon the current evidence of pes planus in this population.

One issue that has not been addressed, and is debated in the Cochrane Review and other articles we have published, is the definition of pes planus. Perhaps one day a consensus of opinion will address the problem.

Contributors

Professor Keith Rome, Professor Podiatry, AUT University, New Zealand

WHAT'S NEW

Date	Event	Description
26 January 2022	Amended	Minor typographical errors in the abstract and plain language summary corrected.
11 November 2021	New citation required but conclusions have not changed	A total of 19 studies are included in this version of the review, an addition of 16 new studies since the last version was published in 2010.
11 November 2021	New search has been performed	Review updated, and current until 01 September 2021

HISTORY

Protocol first published: Issue 1, 2007

Review first published: Issue 7, 2010

Date	Event	Description
25 September 2012	Feedback has been incorporated	New feedback
10 August 2011	Feedback has been incorporated	Feedback added

CONTRIBUTIONS OF AUTHORS

AE, KR, and FH conceived and designed the review. AE collected and inputted the data. AE, and MC conducted the GRADE analyses. AE, KR, and FH compiled the data, and AE drafted the manuscript. All authors read and approved the final manuscript.

DECLARATIONS OF INTEREST

KR, AE are authors of the 2010 systematic review addressing this topic.

AE declares authorship of Evidence Essentials (www.evidenceessentials.com; blog and monograph series), Board Directorship Australian Podiatry Association, Board Directorship AnglicareSA

MC: none known

KR: none known

FH: none known

SOURCES OF SUPPORT

Internal sources

- Auckland University of Technology, New Zealand
- University of Newcastle, NSW, Australia
- La Trobe University, Melbourne, Australia

External sources

- No sources of support provided

DIFFERENCES BETWEEN PROTOCOL AND REVIEW

Post hoc, we decided to include only FOs as the main comparisons, viz CFO versus shoes, PFO versus shoes, PFOs versus CFOs. Similarly, we had intended to include any paediatric population, and separate symptomatic versus asymptomatic participants. Post hoc, we decided to address two paediatric populations, namely asymptomatic, juvenile idiopathic arthritis (JIA), or other clinical concerns.

We did not describe risk of bias and summary of findings tables, or GRADE analyses in the original protocol, but we included them in this review update, in accordance with current Cochrane standards.

We extended both outcomes and comparisons, and better defined them in this review update, which includes 16 trials versus the 3 trials available in 2010 ([Rome 2010](#)). Further, the updated review is more structured regarding participants, intervention comparisons (three comparisons), and outcomes.

Intended comparisons were: any FOs versus sham; any FOs versus shoes; customised FOs (CFOs) versus prefabricated FOs (PFOs).

INDEX TERMS

Medical Subject Headings (MeSH)

*Flatfoot [therapy]; *Foot Orthoses; Pain; Pain Measurement; Quality of Life

MeSH check words

Child; Humans