III-Data extraction form for randomised controlled trials

Reviewer:	Date:		Study num	ber:
GENERAL STUDY INFORM	IATION			
First author (e.g. Smith F):				
Correspondence to:			□not	provided
Title:				
Journal:				
Year of publication:	Vol.: N	um.:	Pages:	
Country:	La	anguage:		
Sources of support:				
STUDY DESIGN				
☐ Randomised Controlled Trial				
□ Cluster Randomised Controll	ed Trial			
Details:				
Setting: □Unicenter □Multio	enter (□National /	□Internationa	l) □ De	tail not provided
-	•		,	·
Recruitment period (months)	: 🗆	Detail not pro	vided	
. , ,		·		
STUDY POPULATION AND	PARTICIPANTS	6		
Study population description	:			
Inclusion criteria:				
Exclusion criteria:				
	Flow of par	ticipants		
		Gro	ups	
Invited to participate and/	or corooned for	Interventio	n / Control	Reasons/Details
Invited to participate and/o	or screened for			
Declined to participate				
Excluded				
Randomized				
Dropouts				
Completed				
Analysed				

Baseline characteristics					
	Total:	Intervention	Control		
	(n=)	(n=)	(n=)	Between group difference	
				(statistically significant)	
Age				□ Yes □ No	
Gender				□ Yes □ No	
Weight				□ Yes □ No	
Height				□ Yes □ No	
ВМІ				□ Yes □ No	
				□ Yes □ No	

INTERVENTION

Intervention general description and objectives:

Duration (weeks/months):

Intervention characteristics					
	Daration				Attendance / compliance rate
Intervention:					
Control:					

DATA COLLECTION

Data collection procedures: PRIMARY OUTCOME			
Variables	Responsible (who collected data)	Method (scale, instrument,etc.)	Details
Physical performance			

	Data collection procedures: SECONDARY OUTCOMES				
Variables	Responsible (who collected data)	Method (scale, instrument, etc.)	Details		
Other					
Subjective perception					
Technical & Tactical					
Physiological					

RESULTS

Drop-outs			
Group	Num. (%)	Description/Reasons	
Intervention			
Control			

Results					
	In	tervention (n=)			
Variable (and timing)	Baseline	Post-competition	P-value	Effect size	

Results			
Control (n=)		

Variable and timing	Bas	eline	Post-competition	P-val	ue Effect size
ADVERSE	EVENTS				
7.571.131		A diverse ou	ento collection and re-	novtina	
Registering	adverse		ents collection and re	oorang	
events		□ Yes	□ No		
Results		Total:	Intervention gr	oup:	Control group:
Other desci	riptions:		L		
CONCLUS	IONS				
			Conclusions		
Primary out	come:				
			COMMENTS		
(Add genera	l comments	if relevant)			

METHODS

Methodological details			
		Description / details	
Eligibility criteria specified	□ Yes □ No		
Power calculation	□ Yes □ No		
Method of randomization	□ Adequate / computer generated□ Inadequate□ Not reported		
Allocation concealment	☐ Adequate ☐ Doubtful ☐ Inadequate ☐ Not reported		
Participants: ☐ Yes ☐ No Blinding Coach/es: ☐ Yes ☐ No Outcome assessor/s: ☐ Yes ☐ No ☐ Not reported			
Handling of withdrawals description	□ Yes □ No		
Pre-published study protocol	□ Yes □ No		

APPENDIX 2:

RISK OF BIAS ASSESSMENT FORM FOR RANDOMISED CONTROLLED TRIALS

Reviewer:

First author (year): () Assessment data	: Study number:
Bias domain	Author's judgment (low, unclear, high)	Support for judgment
Random sequence generation (selection bias)	•	· · · ·
Allocation concealment (selection bias)		
Blinding of participants and researchers (performance bias)		
Blinding of outcome assessment (detection bias)		
Incomplete outcome data (attrition bias)		
Selective reporting (reporting bias)		
Other bias		