Supplementary Table 1. Search strategy (up to January 10th 2015).

MEDLINE - Result: 253 studies

- 'clinical trial'** OR 'controlled trial'** OR 'randomized controlled trial'* OR 'randomised controlled trial'** OR 'randomized'** OR 'randomised'**
 OR 'trial'** OR 'controlled clinical trial'**
- 2. 'metabolic syndrome'** OR 'metabolic syndrome x'*
- 3. 1 and 2
- 4. 'resisted training'** OR 'resistance training'* OR 'resisted exercise'** OR 'resistance exercise'** OR 'strength training'** OR 'strength exercise'**
- 5. 3 and 4
- 6. Animal
- 7. 5 not 6

EMBASE – Result: 113 studies

- 1. 'clinical trial'** OR 'controlled trial'** OR 'randomized controlled trial'* OR 'randomised controlled trial'** OR 'randomized'** OR 'randomised'** OR 'trial'** OR 'controlled clinical trial'**
- 2. 'metabolic syndrome'** OR 'metabolic syndrome x'*
- 3. 1 and 2
- 4. 'resisted training'** OR 'resistance training'* OR 'resisted exercise'** OR 'resistance exercise'** OR 'strength training'** OR 'strength exercise'**
- 5. 3 and 4
- 6. Animal
- 7. 5 not 6

THE COCHRANE LIBRARY – Result: 46 studies

1. 'clinical trial'** OR 'controlled trial'** OR 'randomized controlled trial'* OR 'randomised controlled trial'** OR 'randomized'** OR 'randomised'**

OR 'trial'** OR 'controlled clinical trial'**

1. 'metabolic syndrome'** OR 'metabolic syndrome x'*

2. 1 and 2

- 3. 'resisted training'** OR 'resistance training'* OR 'resisted exercise'** OR 'resistance exercise'** OR 'strength training'** OR 'strength exercise'**
- 4. 3 and 4
- 5. Animal
- 6. 5 not 6

Filter: "trials".

SPORTDiscus – Result: 15 studies

- 1. 'clinical trial'** OR 'controlled trial'** OR 'randomized controlled trial'* OR 'randomised controlled trial'** OR 'randomized'** OR 'randomised'** OR 'trial'** OR 'controlled clinical trial'**
- 2. 'metabolic syndrome'** OR 'metabolic syndrome x'*
- 3. 1 and 2
- 4. 'resisted training'** OR 'resistance training'* OR 'resisted exercise'** OR 'resistance exercise'** OR 'strength training'** OR 'strength exercise'**
- 5. 3 and 4
- 6. Animal
- 7. 5 not 6

PEDro – Result: 45 studies

Abstract & Title: metabolic syndrome

Therapy: strength training

Method: clinical trial

*Medical Subject Headings (MeSH); **Keywords.

Supplementary Table 2.	Characteristics of included studies.
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Study name	Patient characteristics and sample size	Participants (inclusion/exclusion criteria)	Study length	Interventions	Metabolic syndrome risk factors assessed and time points	
Castaneda	n = 62	Inclusion: Confirmation of diabetes	3 days p/ week for 16 wks.	RT: 5 exercises;	Triglycerides	
2002[24]	RT group = 31	diagnosis by fasting plasma glucose ≥		intensity: 60-80% 1RM;	HDL-C	
	Control group = 31	7.0 mmol/l or use of diabetic medications.		dose: 9 S/MG/W;	Fasting plasma glucose SBP	
	% Female = 64.5%			Training sessions: Supervised.	DBP	
	RT group = 67.7%	Exclusion: myocardial infarction			WC	
	Control group = 61%	(within past 6 months) and any		Control group: continued		
		unstable chronic condition, including		usual medical care, received	Baseline	
	Age (Mean ± SD)	dementia, alcoholism, dialysis, retinal		Spanish translated diabetes	16 wks	
	RT group = 66 ± 2	hemorrhage or detachment, or		recommendations for self-		
	Control group = 66 ± 1	current participation in resistance		management, and were not		
		training.		given dietary counseling.		
Dunstan	n = 29	Inclusion: Overweight and sedentary;	3 days p/ week for 6 months.	RT: 9 exercises;	Triglycerides	
2002[26]	RT group = 16	had established but not optimally		intensity: 50-85% 1RM;	HDL-C	
	Control group = 13	controlled type 2 diabetes, were not		dose: 9 S/MG/W;	Fasting Plasma Glucose	
		taking insulin and were nonsmokers.			SBP	
	%Female = 44.8%			Training sessions: Supervised.	DBP	
	RT group = 37,5%	Exclusion: history or physical findings			WC	
	Control group = 53,8%	suggestive of ischemic heart disease,		Control group: offered static		
		systemic diseases, uncontrolled		stretching exercises.	Baseline	
	Age (Mean ± SD)	hypertension and advanced diabetic			3 months	
	RT group = 67.6 ± 5.2	neuropathy or retinopathy.			6 months	
	Control group = 66.9 ± 5.3					
Kukkonen-	N = 68	Inclusion: Age 35-50 years, BMI range	3 days p/ week for 6 months.	All groups performed a 2-	Triglycerides	
Harjula	RT group = 26	of 30-40 kg/m ² and waist		month very-low-energy diet	HDL-C	
2005[27]	Aerobic group = 20	circumference over 100 cm.		before training programs.	Fasting Plasma Glucose	

Control group = 22	SBP
Exclusion: Regular medication; plenty RT: 6 exercises;	DBP
All males of physical activity; smokers; resting intensity: 60-80% 1RM;	WC
blood pressure > 160/105 mmHg; dose: 9 S/MG/W.	
Age (Mean ± SD) fasting serum cholesterol >8 mmol L ¹ ;	Baseline
All participants = 42.6 ± triglycerides >4 mmol L ¹ ; blood AT: 60-70% VO ₂ max;	2 months
4.6 glucose >6.7 mmol L ¹ . dose: 135 min/week.	8 months
	31 months
Training sessions: 1 week	ly
training session w	as
supervised.	
Control group: advised not	to
increase physical activity.	
Sigal n = 251 Inclusion: type 2 diabetes for more 3 days p/ week for 6 months. Before randomization,	all Triglycerides
2007[28] RT group = 64 than 6 months and a baseline participants entered a 4-we	ek HDL-C
Aerobic group = 60 hemoglobin A1c value of 6.6% to run-in phase to asse	ss SBP
Combined group = 64 9.9%. adherence.	DBP
Control group = 63	WC
Exclusion: current insulin therapy; RT: 7 exercises;	
% Female = 36.2% participation in exercise 2 or more intensity: 7-9 RM;	Baseline
RT group = 37% times weekly or in any resistance dose: 6-9 S/MG/W.	3 months
Aerobic group = 35% training during the previous 6 months;	6 months
Combined group = 37% changes during the previous 2 months AT: 60-75% VO ₂ max;	
Control group = 35% in oral hypoglycemic, dose: 45-135 min/week.	
antihypertensive, or lipid- lowering	
Age (Mean ± SD) agents or body weight; serum Combined group: RT + AT.	
RT group = 54.7 ± 7.5 creatinine level of 200 mmol/L or	
Aerobic group = 53.9 ± greater; proteinuria greater than 1 Training sessions: Individu	al
6.6 g/d; blood pressure greater than exercise supervision w	as

	Combined group = 53.5 ± 7.3 Control group = 54.8 ± 7.2	160/95 mmHg; restrictions in physical activity because of disease; presence of other medical conditions that made participation inadvisable.		provided weekly for the first 4 weeks after randomization and biweekly thereafter.	
				Control group: asked to revert to pre-study activity levels.	
Stensvold 2010[8]	n = 43 RT group = 11 Aerobic group = 11 Combined group = 10	Inclusion: Patients with metabolic syndrome according to International Diabetes Federation.		RT: 7 exercises; intensity: 60-80% 1RM; dose: 9 S/MG/W.	Triglycerides HDL-C Fasting Plasma Glucose SBP
	Control group = 11 % Female = 39.5%	Exclusion: unstable angina pectoris, uncompensated heart failure, myocardial infarction during the past 4		AT: 70-95% HR _{peak} ; dose: 129 min/week.	DBP WC
	Age (Mean ± SD) RT group = 50.9 ± 7.6	wk, complex ventricular arrhythmias, and kidney failure.		Combined group: RT (1x p/wk) + AT (2x p/wk).	Baseline 12 wks
	Aerobic group = 49.9 ±10.1			Training sessions: Supervised.	
	Combined group = 52.9 ± 10.4 Control group = 47.3 ± 10.2			Control group: was instructed not to change their dietary patterns or physical activity levels during the study period.	
Saremi 2011[10]	n= 21 RT group = 11 Control group = 10	Inclusion: Males with the metabolic syndrome (based International Diabetes Federation); Low physical	Sessions 3 days p/ week for 12 weeks.	RT: intensity: 30-85% 1RM; dose: 6-9 SMG/W.	Triglycerides HDL-C Fasting Plasma Glucose
	All male	activity level (less than 30 minutes of physical activity per day); Aged		Training sessions: Supervised.	WC
	Age (Mean ± SD) All participants = 45,25 ±	between 20-60.		Control group: not participate in any regular exercise.	12 wks

	4,3	Exclusion: Cardiovascular disease;			
		Musculoskeletal problems; Receiving			
Venojarvi	n = 144	any other treatments. Inclusion: age 40-65 years; BMI	Sessions 2x n/ week for 12	RT: 50-85% 5RM;	Triglycerides
2013[11]	n = 144 RT group = 49	between 25.1 and 34.9 kg/m ² ; and	•	dose: 125 min/wk.	HDL-C
2012[11]	Aerobic group = 49	fasting plasma glucose between 5.6			Fasting Plasma Glucose
	Control group = 48	and 6.9 mmol/L.		AT: 55-75% of Heart Rate	-
	Control group - 47			reserve;	DBP
	All males	Exclusion: earlier detection of IGT and		dose: 103 min/wk.	WC
	All Illales	engagement in prescribed diet or			WC
	Age (Mean ± SD)	exercise programs, engagements in		Training sessions: Supervised.	Baseline
	RT group = 54 ± 6.1	regular and physically very rigorous		Hamming Sessions, Supervised.	12 wks
	Aerobic group = 54 ± 0.1	activities and usage of medication		Control group: not participate	12 WKS
	Control group = 54 ± 7.2	affecting glucose balance.		in any regular exercise.	
Earnest	n = 262	Inclusion: type 2 diabetes; sedentary	Sessions 3 days p/ week for 9	RT: 7 exercises;	Triglycerides
2014[7]	RT group = 73	(not participating in RT and Aerobic	, , , ,	intensity: 10-12 RM;	HDL-C
201.[.]	Aerobic group = 72	exercise.	montais.	dose: 6-9 S/MG/W.	Fasting Plasma Glucose
	Combined group = 76			0000.000,000,000	SBP
	Control group = 41	Exclusion: history of stroke, advanced		AT: 65% VO _{2peak} ;	DBP
		neuropathy or retinopathy, or other		dose: 150 min/wk.	WC
	%Female = 62.2%	serious medical condition			-
	RT group = 59%	contraindicated for exercise or that		Combined group: 2x p/wk of	Baseline
	Aerobic group = 62%	may prevent adherence to the study		RT and 3-5x p/wk of aerobic	
	Combined group = 64%	protocol.		training.	
	Control group = 68%	•		-	
	-			Training sessions: Supervised.	
	Age (Mean ± SD)			-	
	RT group = 57 ± 9			Control group: Offered weekly	
	Aerobic group = 54 ± 9			stretching and relaxation	
	Combined group = 55 ± 8			classes.	
	Control group = 59 ±8				

RT: Resistance Training; AT: Aerobic Training; RM: Repetition Maximum; CHD: Coronary Heart Disease; BMI: Body Mass Index; HDL-C: High Density Lipoprotein Cholesterol; SBP: Systolic Blood Pressure; DBP: Diastolic Blood Pressure; WC: Waist Circumference; IGT: Impaired Glucose Tolerance; S/MG/W: Sets for each muscle group per week; HR: Heart rate.

Study	Eligibility criteria specified	Random allocation	Concealed allocation	Groups similar at baseline	Participan t blinding	Therapist blinding	Assessor blinding	Adequate follow-up	Intention to treat analysis	Between group compariso ns	Point estimates and variability	Total (0-10)
Castaneda 2002	No	Yes	No	No	No	No	Yes	Yes	Yes	Yes	Yes	6
Dunstan 2002	Yes	Yes	No	Yes	No	No	No	No	No	Yes	Yes	4
Kukkonen- Harjula 2005	Yes	Yes	No	Yes	No	No	No	Yes	No	Yes	Yes	5
Sigal 2007	Yes	Yes	Yes	Yes	No	No	Yes	Yes	Yes	Yes	Yes	8
Stensvold 2010	Yes	Yes	Yes	Yes	No	No	No	Yes	No	Yes	Yes	6
Saremi 2011	Yes	Yes	No	Yes	No	No	No	No	No	Yes	Yes	4
Venojarvi 2013	Yes	Yes	No	Yes	No	No	No	No	No	Yes	Yes	4
Earnest 2014	No	Yes	No	Yes	No	No	No	No	No	Yes	Yes	4

	(Quality Assessmen	t	Patie	nt, n	Effect		
Outcomes	Risk of Bias ‡			RT Group	Control Group	MD† (95% CI)	- Quality	
Fasting Plasma Glucose Seven studies	Serious limitation (-1)	No serious inconsistency	Serious imprecision (-1)	202	168	0.04 (-0.12, 0.20)	Low	
HDL-Cholesterol Eight studies	Serious limitation (-1)	No serious inconsistency	No serious imprecision	266	231	-0.00 (-0.05, 0.04)	Moderate	
Triglycerides Eight studies	Serious limitation (-1)	No serious inconsistency	No serious imprecision	266	231	0.03 (-0.14, 0.20)	Moderate	
Diastolic Blood Pressure Seven studies	Serious limitation (-1)	No serious inconsistency	No serious imprecision	255	221	1.39 (-0.19, 2.98)	Moderate	
Systolic Blood Pressure Seven studies	Serious limitation (-1)	No serious inconsistency	No serious imprecision	255	221	4.08 (1.33, 6.82)	Moderate	
Waist Circumference Eight studies	Serious limitation (-1)	No serious inconsistency	No serious imprecision	266	231	1.09 (-0.12, 2.30)	Moderate	

Supplementary Table 4. Quality summary of outcome assessment (GRADE).

[†] Mean Difference (MD) of the resistance training group compared with the control group.

[‡] More than 25% of participants from studies with low methodological quality (Physiotherapy Evidence Database (PEDro) score <7 points).

§ Substantial I² (>75%).

¶Fewer than 400 participants for each outcome.