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The effect of exercise interventions on resting metabolic rate: A systematic review and metaanalysis

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1	The effect of exercise interventions on resting metabolic rate: a systematic review and meta-
2	analysis.
3	
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1. ABSTRACT

37

38	The systematic review and meta-analysis evaluated the effect of aerobic, resistance and
39	combined exercise on RMR (kCal/day) and performed a methodological assessment of
40	indirect calorimetry protocols within the included studies. Subgroup analyses included
41	energy/diet restriction and body composition changes. Randomized control trials (RCTs),
42	quasi – RCTs and cohort trials featuring a physical activity intervention of any form and
43	duration excluding single exercise bouts were included. Participant exclusions included
44	medical conditions impacting upon RMR, the elderly (≥ 65 years of age) or pregnant,
45	lactating or post-menopausal women. The review was registered in the International
46	Prospective Register of Systematic Reviews (CRD 42017058503). 1669 articles were
47	identified; 22 were included in the qualitative analysis and 18 were meta-analysed. Exercise
48	interventions (aerobic and resistance exercise combined) did not increase resting metabolic
49	rate (mean difference (MD): 74.6 kcal/d [95% CI: -13.01, 161.33], $P = 0.10$). While there
50	was no effect of aerobic exercise on RMR (MD: 81.65 kcal/d [95% CI: -57.81, 221.10], P =
51	0.25), resistance exercise increased RMR compared to controls (MD: 96.17 kcal/d [95% CI:
52	45.17, 147.16], $P = 0.0002$). This systematic review effectively synthesises the effect of
53	exercise interventions on RMR in comparison to controls; despite heterogenous
54	methodologies and high risk of bias within included studies.
55	

- 56 Abstract Word Count 200 words
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58 **2. KEYWORDS**

59 Measurement, Metabolism, Nutrition, Physiology, Exercise.

60

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3. INTRODUCTION

63

Human energy expenditure has three primary components: activity energy expenditure, 64 resting metabolic rate (RMR) and dietary induced thermogenesis (DIT) [1]. The accurate 65 66 measurement and interpretation of RMR is beneficial as it is a principal contributor to daily energy expenditure. In practice, this is usually measured by Indirect Calorimetry, a method 67 that is 'indirect' as it measures airflow and the percentage of oxygen (O₂) and carbon dioxide 68 (CO₂) to generate the respiratory exchange ratio (RER) which is subsequently converted to 69 energy expended through known relationships [2, 3]. It is important for practitioners to 70 understand how behaviours and lifestyle can impact on components of energy expenditure, in 71 particular the effect of exercise on RMR is of interest as it has implications for health and 72 sports performance. Despite this, there is a lack of agreement in the literature regarding the 73 74 potential for exercise to modulate RMR in humans.

75

Previous studies have reported increases, decreases or no change in RMR as a result of 76 77 chronic adaptations to endurance or resistance exercise programs [4-9]. These differences may be attributable to a range of factors. For example, changes in body composition directly 78 impact RMR due to the relative energy contribution of different body tissues; fat-free mass is 79 known to explain 25 - 70% of the variance in RMR and therefore gains and/or losses in 80 skeletal muscle due to resistance or aerobic exercise can impact on RMR [10, 11]. As well, 81 82 changes in dietary intake and/or energy expenditure with an exercise program will impact RMR and its interpretation [12]. In addition to these primary factors, other physiological and 83 genetic factors contribute as exercise has the ability to impact thyroid status, protein turnover, 84 85 circulating leptin [13], thermogenesis [14], β-adrenergic stimulation [15] and mitochondrial activity in the liver [16]. While understanding these factors is important for the interpretation 86 of changes in RMR, equivocal changes in RMR as a response to exercise have also been 87

88 attributed to sample size, differences in methodology - particularly the timing and technique

89 of measurement - and the intensity and duration of exercise programs [17].

90

While Indirect Calorimetry is widely accepted as a valid and reliable method of determining 91 RMR, high precision in the estimate of RMR is achieved when best-practice methodologies 92 are employed [18, 19]. In short, several aspects of measurement must be standardised 93 including familiarisation and/or acclimatisation with the measurement and the ventilated 94 95 hood, test conditions, stimulant intake, food intake and physical activity prior to measurement, physiological state (e.g. illness, medications, altitude) and the method of 96 measurement and analysis [18, 19]. The method has been used successfully in the general 97 population and is regularly reported in studies examining the effects of exercise on whole 98 body metabolism [20, 21]. However, it is currently unclear whether publications that report 99 changes in RMR adhere to, and report, best practice protocols. 100

101

This systematic review synthesised evidence from experimental intervention studies that 102 assessed the effect of exercise programs including resistance exercise or endurance/aerobic 103 104 exercise on RMR to assess the primary research question 'what is the effect of aerobic, resistance and combined exercise training modalities on RMR (kCal/day) measured by 105 106 indirect calorimetry in comparison to a control group?. In addition, secondary aims for this systematic review included 1) performing subgroup analyses assessing the impact of 107 energy/diet restriction, changes in body weight and body composition on changes in RMR 108 and 2) providing an overview of the methodologies reported in the included studies 109 measurement of RMR and how these align with best practice guidelines. It is hypothesised 110 that regular or prolonged exercise would have a measurable effect on RMR in accord with 111 changes in body composition. 112

113 4. MATERIALS AND METHODS

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1	1	Δ
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115	This systematic review was conducted in line with the guidelines of the Preferred Reporting
116	Items for Systematic Reviews and Meta-Analysis: The PRISMA statement [22], and the
117	guidelines of the Cochrane Handbook for Systematic Reviews and Interventions [23]. The
118	methods including the eligibility criteria, search strategy, extraction process and analysis
119	were pre-specified and documented in a protocol that was published in the International
120	Prospective Register of Systematic Reviews (CRD42017058503) available at
121	https://www.crd.york.ac.uk/PROSPERO/display_record.php?RecordID=58503.
122	
123 124	4.1. Literature search
125	A literature search was performed in the electronic databases MEDLINE, EMBASE,
126	CENTRAL and SPORTSDISCUS (from inception to July 22, 2018), using a combination of
127	subject headings, free text terms and synonyms relevant to this review, in consultation with a
128	systematic review search librarian (Supplemental Table 1). There was no date or language
129	restriction in the search strategy non-English studies were translated and assessed against
130	inclusion criteria. A multi-step search approach was taken to retrieve relevant studies through
131	additional hand-searching. Two review authors (DS and JK) screened articles in a blinded,
132	standardized manner, with disagreements in judgement resolved by consensus or a third
133	reviewer (KMcKS).
134	
135 136	4.2. Study selection
137	Search results were merged into reference management software Endnote (X8; Thomson
138	Reuters) and de-duplicated prior to screening. Studies were included if they met all of the
139	following criteria: 1) randomized controlled trial (RCT), cluster RCT, quasi-RCT,
140	prospective cohort and retrospective cohort trials; 2) inclusion of adult participants (≥18 years

of age); 3) intervention involving exercise and physical activity training; 4) inclusion of nonexercising control group as a comparator; 5) assessed resting metabolic rate (RMR) at the
beginning and end of intervention using indirect calorimetry.

144

Studies involving populations with conditions impacting upon RMR - including medical 145 conditions such as sepsis and thyroid conditions the elderly (≥65 years of age), or pregnant, 146 147 lactating, or post-menopausal women were excluded. Studies involving the use of medications or known stimulants known to elevate RMR were also excluded [18, 19]. 148 Eligible interventions included physical activity or training of any form (e.g. aerobic exercise, 149 resistance training or concurrent training) of any duration, although studies involving a single 150 (acute) exercise bout were excluded. Studies involving multifactorial interventions involving 151 physical activity and dietary change were included if the dietary change delivered as the 152 intervention also served as the non-exercising comparator. 153

154

The primary outcome was between-group differences in either RMR, resting energy
expenditure or basal metabolic rate at the end of intervention, as well as changes from
baseline. Studies were included only if they reported on the primary study outcome, as either
between-group differences or changes from baseline.

159

160 **4.3. Data extraction and management**

161

162 Three reviewers (DS, JK and KMcKS) independently extracted the data from eligible studies,

and one reviewer (KMcKS) determined the final extraction when there were differences or

164 omissions. Data extracted included: study design (duration, location, details of 'run-in'

165 periods); participant characteristics, intervention details (type of physical activity, intensity,

166 duration and compliance); and other information including indirect calorimetry methodology

used, body composition assessment method and change in body composition analysis.

169	For all pre-specified primary, secondary and exploratory outcome data, the mean, standard
170	deviation (SD), standard error (SE) or 95% confidence intervals (CI) that were reported at
171	end of intervention were extracted for analysis. Where studies involved multiple intervention
172	groups involving different types of physical activity, data was extracted for each intervention
173	for separate analysis. Where multiple intervention arms reported the same type of activity (for
174	example two different aerobic activities) results were combined and compared against the
175	control in one analysis.
176	
177	Risk of bias was independently assessed by two reviewers (DS and JK) using Cochrane
178	methodology [24] which assesses five domains of potential bias with each domain rated
179	either low, unclear or high risk of bias. Disagreements in risk of bias between the two
180	independent reviewers were resolved through discussion.
181	
182 183	4.4. Statistical analysis
182 183 184	4.4. Statistical analysis The overall treatment effect of physical activity on primary and secondary outcomes was
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182 183 184 185 186	4.4. Statistical analysis The overall treatment effect of physical activity on primary and secondary outcomes was calculated using the difference between either the end of intervention values or change scores for the intervention and comparator groups. Variance was calculated from the SD and SE of
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182 183 184 185 186 187 188	4.4. Statistical analysis The overall treatment effect of physical activity on primary and secondary outcomes was calculated using the difference between either the end of intervention values or change scores for the intervention and comparator groups. Variance was calculated from the SD and SE of end of intervention values or change scores, or from the confidence intervals (CI) where these values were not available [25]. In crossover studies, the mean and SD, SE or CI of
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182 183 184 185 186 187 188 189 190 191 192 193 194	4.4. Statistical analysis The overall treatment effect of physical activity on primary and secondary outcomes was calculated using the difference between either the end of intervention values or change scores for the intervention and comparator groups. Variance was calculated from the SD and SE of end of intervention values or change scores, or from the confidence intervals (CI) where these values were not available [25]. In crossover studies, the mean and SD, SE or CI of intervention and control periods were extracted and analyzed separately [26]. Where intervention endpoint data was unable to be obtained, the results were described narratively. Meta-analysis was performed where outcomes were reported in at least two studies using Revman (Version 5.3; Cochrane Collaboration). Outcome data was converted to the same units prior to meta-analysis (kcal/day) and was reported as the mean difference (MD)[27]. A

^{12/05/2020,} available online: http://www.tandfonline.com/10.1080/02640414.2020.1754716.

random-effects model was used to produce a pooled estimate of the MD, and the fixed-effects model was used to check for robustness and potential outliers. Inconsistencies between studies were assessed using the I² statistic, where significant heterogeneity was defined as I² $\geq 50\%$.

199

Post hoc subgroup analyses were undertaken for primary and secondary outcomes that were
reported in at least two studies in each subgroup. Post hoc subgroup analyses included:
intervention types (aerobic and resistance training), exercise-alone versus combined dietexercise interventions, changes in total body mass (TBM) during the study period (increased;
decreased; stable; and not reported). These were categorised (decreased, versus stable, versus
increased) where a significant change in body composition was reported.

206

In studies including multiple, separate arms involving different exercise interventions, the interventions were pooled together for the overall meta-analysis, with a weighted average of the intervention arms and study variance calculated [28]. In the subgroup analyses exploring the effect of different intervention types on RMR, the interventions were analysed separately based on their respective intervention types

212

Significant outliers were determined by visual inspection as well as through a study-by-study
sensitivity analysis, where each study was sequentially omitted, and the remaining data reassessed. If a study contributed to over 30% heterogeneity (based on changes to the I²
statistic) then it was removed from the analysis in the sensitivity analysis [27]. Funnel plots
were generated for outcomes where at least 10 studies were included in the meta-analysis
[29] and reporting bias detected by assessment of funnel plot asymmetry by visual inspection.

5. RESULTS

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221	The literature search identified 1669 articles; the PRISMA Diagram in Figure 1 summarises
222	the results of the literature search. 22 studies were included in the qualitative analysis and 18
223	studies provided enough information to be included in the meta-analysis.

- 224
- 225

5.1. Study characteristics

226

227 The general characteristics of trials included in the systematic review are summarised in

Table 1. A total of 822 participants were captured in 22 studies; with most including less than

45 participants with the exception of Scharhag-Rosenberger et al. [30], Frey-Hewitt et al.

[31], Jennings et al. [32] and Gomersall et al. [33] which included 74, 85, 103 and 107

participants, respectively. One study by Hunter et al. [34] did not specify the exact number of

232 participants but reported the inclusion of at least 140 participants. The meta-analysis included

data from 392 participants and 270 controls. Most of the studies were a parallel study design

except for one cross-over study design [35]. The majority of studies were conducted in

overweight/obese populations that were predominantly sedentary [5, 31, 32, 34-44], two in

type-2 diabetic populations [32, 40], one in a population with metabolic syndrome [37],

several in predominantly normal-weight and/or healthy sedentary populations [17, 30, 33, 45-

48] and one in active, healthy populations [20]. All studies captured were in adult

populations, with several predominately focussing on females [5, 34, 36, 39, 42-44, 46, 48],

240 males [17, 20, 31, 38, 41, 47], a combination of both [30, 32, 33, 35, 40, 45] or gender was

not reported [37].

242

243 Several interventions were exercise only; with either a predominant focus on aerobic exercise

244 [17, 31, 40], resistance exercise [5, 30, 35, 38, 46, 48] or a combination of both exercise

245 modalities [32, 33]. Many studies used a combined dietary and exercise intervention; with

four studies using predominantly aerobic exercise [36, 37, 45, 47], two in resistance exercise

[20, 39] and five using a combination of both exercise modes [34, 41-44]. The shortest

248	interventior	was 10 days [47]; while several studies were conducted over 2-6 weeks [20, 33,	
249	39, 40, 43].	The majority of interventions were conducted over 12 weeks [17, 36, 37, 41, 42,	,
250	44-46] whil	e several longer interventions spanned 20-24 weeks [5, 32, 35, 38] and the longe	st
251	study interv	rention was 12 months [31]. While some studies did not measure or report body	
252	composition	n assessments [33, 37]; the majority of studies used Dual-Energy X-Ray	
253	Absorption	netry (DEXA) [20, 34-36, 39, 40, 45, 48], anthropometry/skinfolds [30, 38, 43,	
254	46], hydros	tatic weighing, underwater weighing/air-displacement plethysmography [5, 17,	
255	31, 41, 44,	47] or bio-electrical impedance (BIA) [32, 42].	
256			
257 258	5.2.	Meta-analysis	
259	Eighteen st	udies were able to be meta-analysed. Four studies were not included in the meta-	
260	analysis as	they only presented data in graphs or with no means/variance reported [37, 42],	
261	did not con	ain specific participant numbers [34] or did not report outcome data in units that	
262	were able to	be reliably converted for meta-analysis [30].	
263			
264	Across the	18 intervention studies pooled into meta-analysis, exercise (aerobic and resistance	e
265	exercise con	nbined) did not significantly increase RMR (MD: 74.16 kcal/day [95% CI: -	
266	13.01, 161.	33], $P = 0.10$; Figure 2). There was high heterogeneity ($I^2 = 96$ %); with two	
267	studies cont	ributing as outliers [31, 36]. Neither study contributed over 30% toward the total	
268	heterogenei	ty, with 7% (21) and 22% (26), respectively. However, removal of these two	
269	studies fron	n the analysis reduced the heterogeneity to 20%, and the overall finding became	
270	significant	(MD: 61.45 kcal/day [95% CI: 27.46, 95.44], <i>P</i> =0.0004).	
271			
272	Aerobic exe	ercise did not significantly increase RMR compared to the control group (MD:	
273	81.65 kcal/o	lay [95% CI: -57.81, 221.10], <i>P</i> = 0.25, Figure 2), however there was high	
274	heterogenei	ty ($I^2 = 98\%$)Resistance exercise significantly increased RMR compared to the	

control group (MD: 96.17 kcal/day [95% CI: 45.17, 147.16], P = 0.0002; Figure 2) with minimal statistical heterogeneity ($I^2 = 0\%$).

- 277
- 278 **5.3.** Subgroup analyses
- 279

280

Subgroup analysis comparing the effects of exercise-only interventions with combined

exercise and dietary interventions showed that showed that both types of interventions led to

a similar effect, with neither exercise-only (MD: 46.79 kcal/day [95% CI: -9.52,103.09], P =

283 0.10, Figure 3) nor exercise and diet (MD: 74.16 kcal/day [95% CI: -13.01, 161.33], P =

284 0.12, Figure 3) subgroups having a significant effect on RMR.

285 Subgroup analysis comparing exercise intervention in individuals based on anthropometric

changes in TBM had a significant effect on RMR. Studies that reported a stable body mass

throughout the intervention period showed exercise increased RMR (MD: 66.17 kcal/day

288 [95% CI: 2.95, 129.38], P = 0.04, Figure 4). Studies that reported either an increase in body

289 mass or failed to report on body mass, showed RMR was not different as it was just outside

290 the P <0.05 pre-determined criteria (MD: 70.61 kcal/day [95% CI: -3.58,144.81], P =0.06,

291 Figure IV and MD: 89.27 kcal/day [95% CI: -3.20,181.74], *P* =0.06, Figure 4). There was no

effect of exercise on RMR in studies that reported a decreased body mass (MD:

293 84.59kcal/day [95% CI: -77.37, 246.54], *P* =0.31, Figure 4).

294 295

5.4. Comparison of study methods

296

297 The methodologies that were used and reported for measuring RMR are summarised in

298 Supplementary File 2. Of the studies that reported RMR methodology; several studies

reported using a ventilated hood [17, 33, 40, 43-45, 47] and several used a mouthpiece with

300 one-way valve/nose clip [31, 39, 46, 48]. Most studies reported measuring RMR for 30 – 45

301 minutes [5, 17, 20, 30, 32-34, 36, 39, 41, 45, 46]; with some reporting shorter durations of 10

302	- 25 minutes [31, 40, 42-44, 48] while others did not report RMR measurement duration [35	,
303	37, 38, 47]. Many studies did not report acclimation or familiarisation to the test protocol bu	t
304	of the available data acclimation was undertaken between 15 - 30 minutes duration [5, 17, 3	1-
305	34, 39-44, 46] While many studies did not report a fasting duration prior to measurement of	
306	RMR studies that provide detailed methods show participants were fasted 10 hours [41], 12	
307	hours [17, 31-33, 39, 40, 43, 46] or overnight prior to commencing the test [20, 34, 48]. Son	ıe
308	studies reported time in recovery/rest following a previous exercise bout; either 12 hours [3]	,
309	33, 47], 24 hours [30, 42], 36 hours [5], 48 hours [17, 32, 48] or 72 hours [35] – however	
310	most did not report the intensity or mode of the last exercise session. The RMR was typicall	y
311	derived from measurements of resting oxygen uptake (VO ₂), carbon dioxide production	
312	(VCO_2) and RER (VCO_2/VO_2) using the Weir formula [49]. Some, but not all, studies	
313	reported the test environment and conditions during which the measurement was undertaken	
314	(e.g. thermo-neutral; low-light). RMR data was reported in a range of units e.g. mJ/d, kJ/d,	
315	kJ/min and was generally reported as an absolute change.	
316		
317	The studies reported several methods of body composition assessment including Dual-Energy	;y
318	X-Ray Absorptiometry [20, 35, 36, 39, 40, 45, 48], Hydrostatic weighing or Air-displaceme	nt
319	plethysmography [5, 17, 31, 41, 44, 47], Bio-electrical impedance [32, 42] or	
320	skinfolds/anthropometry [30, 38, 43, 46]. Several studies reported TBM but did not report	
321	FFM [30, 38, 43, 46] and several studies did not report TBM or FFM [33, 37, 47].	
322		
323 324	5.5. Risk of Bias	
325	The risk of bias was unclear for many of the studies for random sequence generation,	
326	allocation concealment, participant/personnel blinding and selective reporting	
327	(Supplementary File 3). The risk of bias was low for blinding of outcome assessment,	
328	moderate for incomplete outcome data and moderate-high for other bias.	
		13

22% of studies adequately reported random sequence generation to support a low risk of bias 330 assessment and allocation concealment [30, 32, 33, 35, 48]. For all studies, the risk of bias for 331 blinding of the participants to their condition was unclear and the risk of bias for blinding of 332 the outcome was low. For incomplete outcome data; 22% of studies had a high risk of bias 333 [34, 35, 38, 42, 43], 22% had an unclear risk of bias [5, 31, 36, 41, 45] and 55% had a low 334 risk of bias [17, 20, 30, 32, 33, 37, 39, 40, 44, 46-48]. For selective reporting, 9% had low 335 [30, 33], 86% had an unclear [5, 17, 20, 31, 32, 34-48]; while only one study had a high risk 336 of bias [36]. Only a single study was judged as high risk of bias for 'other bias' [34] because 337 it didn't report on participant numbers, with 32% of studies judged as low risk of bias [30-33, 338 38, 40, 47], with the remainder judged to be unclear. 339

340 6. DISCUSSION

341

The primary findings from the review were 1) resistance exercise significantly increased 342 343 RMR in comparison to a control group as measured by indirect calorimetry, 2) aerobic exercise and exercise-combined (i.e. resistance exercise and aerobic exercise) did not 344 significantly increased RMR in comparison to a control group, 3) a lack of comparable body 345 346 composition assessment data meant it was unclear how changes in body composition interacted with changes in RMR and 4) while there were a large proportion of studies which 347 did not report key aspects of their methodology that would represent best practice and/or 348 there was inconsistency in methodology between studies, this meta-analysis only included 349 350 studies with a control group thus limiting the impact of their methodological differences on 351 the meta-analysis

The meta-analysis captured data from 392 participants and 270 controls (total 662
participants) and in large part addresses the inherent limitation of small-scale or single-arm

studies. This systematic review provides new information to show a resistance exercise 354 program has the capacity to increase RMR. A primary adaptation associated with resistance 355 training is upregulation of anabolic processes within skeletal muscle resulting in hypertrophy 356 and increased muscle cross sectional area [50]. It is generally well-accepted that increases in 357 fat-free/lean mass and total body mass may induce an increase in RMR due to greater volume 358 of metabolically active tissue, skeletal muscle remodelling and increasing the fat free-to-total 359 360 body mass ratio [51-53]. Moreover, fat-free mass has been shown to make a substantial contribution (25-70%) to individual variations in RMR [10, 11]. While the findings of the 361 meta-analysis support such a contention, the sub-analyses did not support a clear association 362 between changes in body composition and RMR. Unfortunately, total body mass was not 363 reported on all occasions and while some studies used body composition assessment 364 365 measures that more accurately measure compartmental body mass (i.e. fat mass and fat-free mass) others, such as DEXA, used derived or predicted values to determine reported 366 compartmental body mass. Moreover, there is an increasing awareness of the deficiencies in 367 368 the 2-compartment (FFM and FM) profile of body composition in explaining variance in RMR and in RMR changes, and that the future may lie in an operational quantitative dynamic 369 organ-system RMR model [54]. 370

While the data clearly show resistance exercise is effective for increasing RMR, a similar 371 outcome was not apparent for aerobic exercise. Interestingly, aerobic exercise has the 372 373 capacity to induce modest hypertrophy but the effect may be dependent on the mode and intensity of aerobic exercise and the physical activity status of the participant [55]. In 374 addition, our meta-analysis showed the overall effect of aerobic and resistance exercise 375 combined on RMR was not significant. Therefore, we suggest the addition of higher quality, 376 methodologically sound studies are warranted to better determine the effects of different 377 378 exercise modalities on RMR. While no study contributed greater than 30% heterogeneity; two clear outliers reported a significant increase in RMR following aerobic exercise 379

compared to a control group [31, 36]. As it was not explicitly stated - and the methodological 380 reporting was broad - it was not clear whether the studies adhered to best-practice protocols 381 for the measurement of RMR. Interestingly, when these studies were removed from the 382 analysis there was a significant, positive effect of combined exercise modalities on RMR. 383 A potential confounding factor within the literature that may influence this meta-analysis is 384 the effect of preceding exercise when study cohorts progress from sedentary to exercising 385 status. Specifically, baseline RMR testing may be undertaken without preceding exercise 386 while post-intervention testing may occur with limited recovery after the final exercise bout 387 388 which may artificially inflate the measurement of RMR. It is important that studies follow best practice protocols which prescribe cessation from exercise or vigorous physical activity 389 390 for a standardized period prior to the measurement of RMR. Compher et al. [18] recommend 391 2 hours of abstention from moderate aerobic exercise (Grade II - fair) and 14 hours for vigorous exercise (Grade III - limited) and Fullmer et al. [19] recommend 12-48 hours after 392 light to vigorous intensity physical activity. As many of the participants were untrained and 393 were potentially doing exercise that would generate post-exercise oxygen consumption 394 (EPOC) and due to the potential for micro-trauma and repair of muscle damage, it has also 395 been suggested that longer periods of abstinence up to 72 hours may be warranted [53]. Many 396 studies in the current meta-analysis did not report abstinence from physical activity prior to 397 the measurement of RMR. If exercise was performed in this time this could artificially inflate 398 the measurement and thus the authors could conclude an effect of the exercise intervention on 399 RMR; however as there was a methodologically-comparative control group in each study the 400 overall effect in this meta-analysis would not be impacted. In addition, while our inclusion 401 criteria allowed for interventions that both included or did not include dietary interventions, 402 and energy balance is one consideration that may influence RMR independent of training 403 [12], these were only included where the diet only intervention served as the control group. 404

The sub-analysis confirmed that the effect of exercise on RMR was similar between exercise-only and combined dietary-exercise studies.

The methodology characteristics table (Supplementary File 2) highlighted several gaps in the 407 408 included study methodologies when compared to best practice guidelines. While many studies reported a fasting period in-line with best-practice guidelines, other areas of 409 standardisation including familiarisation, time-of-day, room conditions, body position, the 410 control for stimulants or supplements and physiological conditions (illness, medications) 411 prior to measurement was minimal. Other key aspects of RMR methodology, including the 412 413 calculation of steady-state and calibration procedures were not routinely reported despite being important aspects of evidence-based practice [18, 19]. The risk of bias was moderate-414 high for some of the studies. While most studies did not report random sequence generation 415 416 or allocation concealment, this is difficult in small-scale studies that include an exercise intervention. 417

418 This systematic review and meta-analysis clearly shows that resistance exercise generates increases in resting metabolic rate while aerobic and combined resistance and 419 aerobic exercise fail to induce a robust effect on changes in RMR. While some limitations of 420 421 this systematic review have already been discussed, it should also be noted that number of 422 observations can impact statistical significance and there were less resistance exercise 423 studies. In addition, the overall effect had wide confidence intervals suggesting a high variability in data. The systematic review included exercise interventions of any type and 424 duration, excluding single exercise bouts, and thus compared different study designs and 425 methodologies. For example, while there was a clear effect of resistance exercise on RMR, 426 427 differences in the type of resistance exercise and its' overarching aim (i.e. changes in power, strength or muscular endurance) were beyond the scope of this review. As well, the effect of 428 exercise was most evident when total body mass remained stable during the intervention 429 period, but lack of comparable data means it was unclear how changes in body composition 430

interacted with changes in RMR. Despite this, a strength of this systematic review and metaanalysis is that it addresses the inherent limitation of small-scale or single-arm studies as it
included a range of studies in comparison to control group. It is strongly recommended that
future studies to adhere to best-practice protocols in the measurement of RMR and body
composition assessment and to ensure that methodology is adequately reported to permit
replication and appropriate interpretation [18, 19].

437

7. AUTHORSHIP CONTRIBUTION

KMS, NB and VC contributed to the study design concept and protocol. KMS, DS and JK
contributed to the initial and updated literature search and screening, data extraction and risk
of bias. KMS drafted the manuscript with contribution from DS and JK. All authors
performed critical analysis and revision of manuscript and approved the final version.

442 **8.** A

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446 9. CONFLICT OF INTEREST

447 Authors K. MacKenzie-Shalders, J.T. Kelly, D, So, V.G, Coffey & N.M. Byrne declare they
448 have no conflicts of interest.

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594	
595	Figure Legends
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597	Figure 1: Flow diagram of studies evaluated in the systematic review.
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599 600 601 602 603 604 605 606	Figure 2: Forest plot of randomized controlled trials in adults comparing interventions involving exercise and physical activity training with non-exercising control group comparators. The overall effect of exercise and physical activity is presented (1.2.1). Additionally, sub-group effects based on the specific type of exercise training are also presented: aerobic (1.2.2) and resistance (1.2.3). Data are presented as means and SDs of RMR at the end of intervention. Effects of trials are presented as kilocalorie per day and MD (95% CI). CI, confidence interval; IV; inverse variance; MD, mean difference; RMR, resting metabolic rate; SD, standard deviation.
607	
608	Figure 3: Forest plot of randomized controlled trials in adults comparing interventions

involving exercise and physical activity training with non-exercising control group

610 comparators. Studies are sub-grouped by whether the exercise and physical activity training

611 was delivered alone (1.14.1) or in combination with dietary modifications (1.14.2). Data are

bresented as means and SDs of RMR at the end of intervention. Effects of trials are presented

as kilocalorie per day and MD (95% CI). CI, confidence interval; IV; inverse variance; MD,

614 mean difference; RMR, resting metabolic rate; SD, standard deviation.

615

Figure 4: Forest plot of randomized controlled trials in adults comparing interventions

617 involving exercise and physical activity training with non-exercising control group

618 comparators. Studies are sub-grouped based on the mean reported changes in total body mass

of participants during the study period, categorised as: stable (1.6.1); increased (1.6.3);

620 decreased (1.6.4); and not reported (1.6.6). Effects of trials are presented as kilocalorie per

day and MD (95% CI). CI, confidence interval; IV; inverse variance; MD, mean difference;

622 RMR, resting metabolic rate; SD, standard deviation.