- 1 Systematic Review
- 2 Validity of Dietary Assessment in Athletes: A
- 3 Systematic Review
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- Received: date; Accepted: date; Published: date
- 19 Abstract: Dietary assessment methods recognized as appropriate for the general population are
- 20 usually applied in a similar manner to athletes, despite knowledge that sport-specific factors can
- $21 \hspace{0.5cm} \text{complicate assessment and impact accuracy in unique ways. As dietary assessment methods are used} \\$
- 22 extensively within the field of sports nutrition, there is concern the validity of methodologies have
- 23 not undergone more rigorous evaluation in this unique population sub-group. The purpose was to

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systematically review studies comparing two or more methods of dietary assessment, including dietary intake measured against biomarkers or reference measures of energy expenditure, in athletes. Six electronic databases were searched for English-language, full-text articles published from January 1980 until June 2016. The search strategy combined the following keywords: diet, nutrition assessment, athlete and validity; where the following outcomes are reported but not limited to: energy intake, macro and/or micronutrient intake, food intake, nutritional adequacy, diet quality, or nutritional status. Meta-analysis was performed on studies with sufficient methodological similarity, with between-group standardized mean differences (or effect size) and 95 % confidence intervals (CI) calculated. Of the 1624 studies identified, 18 were eligible for inclusion. Studies comparing selfreported energy intake (EI) to energy expenditure assessed via doubly labelled water were grouped for comparison (n=11) and demonstrated mean EI was under-estimated by 19 % (- 2793 ± 1134 kJ/d). Meta-analysis revealed a large pooled effect size of - 1.006 (95% CI: -1.3 to -0.7; p<0.001). The remaining studies (n=7) compared a new dietary tool or instrument to a reference method(s) (e.g. food record, 24-h dietary recall, biomarker) as part of a validation study. This systematic review revealed there are limited robust studies evaluating dietary assessment methods in athletes. Existing literature demonstrates substantial variability between methods, with under and misreporting of intake frequently observed. There is a clear need for careful validation of dietary assessment methods, including emerging technical innovations, among athlete populations. Keywords: dietary assessment; food record; FFQ; biomarker; doubly labeled water; energy intake; validity; athletes; sports nutrition. 1. Introduction Adequate dietary intake is important for normal growth and development, maintaining health and

well-being, reducing the risk of illness and injury, and optimizing sports performance [1]. Individual

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dietary requirements are influenced by a range of factors such as age, gender, body mass, stature, and growth and development needs (for child or adolescent athletes) [1,2]. In addition, a range of sport-specific factors such as type of sport, training volume and intensity also influence dietary requirements which are not static due to periodization of training load across days, weeks or months of a competitive season [1,2]. Athletes have special nutrition needs which usually encompass a higher energy requirement to account for greater energy expenditure, increased protein and carbohydrate requirements to support lean mass accrual and/or maintenance, and glycogen stores, respectively as well as an increased requirement for certain micronutrients (e.g. iron, calcium, sodium) [1,2,3]. Dietary assessment is routinely undertaken by nutrition professionals to evaluate whether an individual is achieving specific health and/or sports nutrition targets [4]. However, accurate dietary assessment of athletes is complex due to the influence of sport-specific factors as well as the widespread use of rapidly evolving sports foods and supplements. Dietary assessment can be expensive (especially for large numbers of athletes or teams), time consuming and unfortunately due to these constraints, undertaken less frequently. As a result, athletes with inadequate or poor dietary intake may not be readily identified. Numerous studies have reported on dietary intake of athletes from a wide range of sports [5-9]; with many comparing self-reported intake to calculated energy requirements based on general recommendations and/or sports nutrition guidelines [10-14]. However, the discrepancy between selfreported intake and estimated energy expenditure has been reported in a number of studies involving male and female athletes, across a range of age-groups and a variety of sports [15-17]; with substantial differences (11-44 %) observed [18,19].

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Dietary assessment methodology recognized as suitable for non-athletes are usually applied in a similar manner to athletes, although specific factors such as training status, competition level and nutrition-associated beliefs and dietary practices could influence aspects of the dietary assessment process [19]. Many retrospective methods that rely on self-reporting of intake (e.g. food record, 24-h dietary recall, diet history) are susceptible to measurement error including conscious or subconscious exclusion of foods consumed [4,19-21]; and possible change in usual food intake or dietary patterns due to the nature of the dietary intervention itself [22]. By far, the most common dietary assessment method applied in sports nutrition research and practice is the food record (FR); where all food and drinks consumed are recorded by the participant for a specified number of days (i.e. 3-7 days) [4]. However, it has been widely reported that selfreported FR under-estimates intake as noted in a number of population groups including adolescents and children [23], obese individuals [24], and athletes [18,25,26]. Despite the FR being regarded a 'gold standard' in dietary assessment, it places substantial burden on participants to document information truthfully and accurately, as well as reliance on investigators to code the data correctly using appropriate databases [19,27-29]. Additional challenges facing an athlete cohort include increased burden from recording large food intakes, frequent eating occasions, potentially irregular meal patterns, difficulty estimating large portion sizes, and contribution of sports foods and supplements [19]. Recent technological developments involving the use of food photography or electronic images, wearable cameras, and various on-line tools and applications have been shown to improve participant compliance by reducing the burden of recording and enhance the accuracy of data recorded [30-33]. There is a need for high quality research to assist in identifying dietary methodologies, including emerging technologies, which are valid, as well as feasible, for use in thisunique population sub-group.

Despite a clear need for careful validation of dietary assessment methods in athletes, there has been relatively little attention directed to the applicability and validity of dietary assessment methods in this population sub-group. The purpose of this systematic review was to evaluate studies comparing nutrition intake from two or more dietary assessment methods, including intake measured against biomarkers or reference measures of energy expenditure, in athletes.

### 2. Methods

# 2.1 Search Strategy

A systematic literature search was conducted of articles published between January 1980 to June 2016 to identify studies comparing the nutrition intake assessed by two or more methods of dietary assessment, including a measure of intake against dietary biomarkers or reference measures of energy expenditure, in an athlete cohort. Databases searched were AUSPORT MED (via Informit Online), CINAHL (via EBSCO), EMBASE (via EBSCO), MEDLINE (via OvidSP), SPORTDiscus (via EBSCO) and the Web of Science. Ongoing electronic monitoring was established to assess the inclusion of eligible recent publications.

The search strategy combined the following keywords: diet (e.g. energy intake, nutritional status, nutritional requirement, food group); nutrition assessment (e.g. food frequency questionnaire, food record, diet score, diet survey, energy expenditure, biomarkers); athlete (e.g. elite athlete, team sport, collegiate, Olympic athlete); and validity (e.g. under-reporting, measurement error). In addition, the reference lists of all retrieved papers and relevant reviews were manually searched for

eligible papers. Following the search, we completed a PRISMA (Preferred Reporting Items for Systematic Reviews and Meta-Analyses) [34] informed systematic review process.

## 2.2 Selection of Studies

Study participants had to be 'athletes' (i.e. amateur, collegiate, tertiary, professional, national or international calibre) as defined by the study authors. Included studies needed to validate at least one dietary assessment method (e.g. food frequency questionnaire, 24-h dietary recall, food record) against at least one other dietary assessment method or a biological (blood or urine) measure for nutrient status (e.g. 24-hr nitrogen excretion, antioxidant concentrations). In addition, studies which compared reported energy intake to measured energy expenditure (e.g. doubly labelled water technique, activity records) were also included. Descriptive, cross-sectional or longitudinal study designs were eligible for inclusion. Studies including participants with a mean age lower than 16 years of age, papers published in a language other than English, case studies, reviews, abstracts and theses were excluded (Supplementary Material 1). After eliminating duplicates, one reviewer (LC) screened the search results against the eligibility criteria. References that could not be eliminated by title and abstract were retrieved in full and subsequently assessed against the eligibility criteria in duplicate (KB, GS) prior to inclusion.

### 2.3 Data Extraction and Conversions

Data relating to the manuscript, namely author(s), date of publication and country where the study was conducted were extracted. The institution country of the first author was used if the study country was not described in the paper. Data extracted from each publication included: participant characteristics (i.e. age, sex, sport and athlete calibre), participant anthropometric (i.e. stature, body mass) or body composition parameters (e.g. percent fat), and dietary assessment methodology (e.g.

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food record, food frequency questionnaire, 24-h dietary recall). In addition, information on reported energy and macronutrient intake, resting energy expenditure, total energy expenditure, and key statistical comparisons between dietary assessment methods and correlation coefficients, were extracted in conjunction with main study findings. Where a publication contained insufficient information, attempts were made to contact the study author(s) to obtain missing details. To aid comparison between the included papers, anthropometric parameters reported in imperial units (e.g. feet, inches, pounds) were converted to metric units (e.g. kg, cm) (1kg = 2.2 pounds; 1 cm = 0.3937 inches). Data reported in calories (e.g. energy intake, energy expenditure) were converted to kilojoules (1 Cal = 4.184 kJ). Where reported, macronutrient values were converted from grams or grams per kilogram body weight to per cent of mean energy intake using Atwater factors (i.e. protein 17 kJ/g or 4 Cal/g; fat 37 kJ/g or 9 Cal/g; carbohydrate 16 kJ/g or 4 Cal/g) [35]. Extracted data were presented as mean and standard deviation (SD) where reported. Weighted means were calculated for age, anthropometric variables and the differences between reported energy intake and measured energy expenditure.

# 2.4 Assessment of Methodological Quality

The methodological quality of all included studies were independently assessed by three researchers (LC reviewed all papers; KB and JG shared the evaluation of the other studies) using a modified assessment scale created by Downs and Black [36]. Using the scale, 18 of the 27 criteria that logically applied to the study designs were retained, while items 4, 8, 9, 13, 14, 17, 21, 23, 24 were deemed not relevant to the included studies and were excluded. Three additional items from a nutrition-specific quality criteria checklist were incorporated [37]; specifically item 8.6 *Was clinical significance as well as statistical significance reported?*; item 9 *Are conclusions supported by results with* 

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biases and limitations taken into consideration?; and item 10 Is bias due to study's funding or sponsorship unlikely? Where relevant, other items from the Academy of Nutrition and Dietetics quality criteria checklist [37] (i.e. items 2.1, 2.3, 3.4, 5.2, 7.2, 7.3, 7.4, 7.5 and 7.6) were considered concurrently to the Downs and Black [36] criteria to provide reviewer clarification when assessing quality ratings. Two variables from a third checklist described by Serra-Majem and colleagues [38] were applied specifically to validation studies to evaluate the validity of the dietary methodology used and suitability of the method for an athlete cohort. The specific variables applied to assess the quality of the validation study were statistics to assess validity (i.e. to support Downs and Black [36] item 18: Were the statistical tests used to assess the main outcomes appropriate?); and to assess whether the sample size was adequate to assess validity (i.e. to support Downs and Black [36] item 27: Did the study have sufficient power to detect a clinically significant effect where the probability value for a difference being due to chance is less than 5%? (Supplementary material 2). Each reviewer checked for internal (intra-rater) validity across items for each included publication. Disagreements in ratings were resolved by discussion and through adjudication with two separate researchers (HO, VF). No further sub-analysis was undertaken on the basis of methodological quality and none of the studies were eliminated. 2.5 Meta-Analysis Studies which involved the comparison of a dietary measure (i.e. food record) to energy expenditure as measured by DLW were sufficiently similar methodologically to enable meta-analysis to be performed. To quantitatively compare differences between reported and measured energy intake in the papers using DLW, the between-group standardised mean difference, or effect size (ES) and 95 % confidence interval (CI) was calculated. The extracted data (i.e. mean energy intake, mean

energy expenditure, SD and sample size) was transferred into Comprehensive Meta-Analysis (CMA)

version 2 software (Biostat, 2005, Englewood, IL, USA) in order to calculate the ES, standard error, variance and 95 % CIs. A forest plot was generated to display the ES and 95 % CIs from each study and the pooled estimate determined whereby a small ES was > 0.2, a medium ES > 0.5 and a large ES > 0.8 [39]. A positive ES indicated an effect favouring energy measured by DLW while a negative ES indicated an effect favouring reported EI.

### 3. Results

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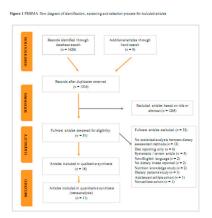
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3.1 Identification and Selection of Studies

The initial literature search identified 1624 potentially relevant papers. After removal of duplicates and elimination of papers based on title and abstract, 42 full text publications were reviewed in duplicate against the inclusion criteria. A manual search of reference lists identified an additional nine papers. Thirty-three papers were excluded because they did not meet the inclusion criteria, leaving 18 full-text articles for assessment (Figure 1).



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Figure 1 PRISMA flow diagram of identification, screening and selection process for included

191 articles

192 3.2 Demographic and Anthropometric Characteristics

Of the 18 studies included, 11 compared reported energy intake (EI) against energy expenditure (TEE) as measured by DLW (TEE<sub>DLW</sub>) [21,22,40-48]. The remaining seven papers [29,49-54] compared

195 a new dietary tool (n=3) or FFQ or modified FFQ (n=4) to a reference method(s) (i.e. food record, 24-196 h dietary recall, direct observation, biomarker) as part of a validation study. Of the 18 included 197 studies, three compared one or more dietary assessment method(s) to a biomarker (i.e. urine n=2, 198 blood *n*=1) [45,50,54]. 199 The total number of study participants was 683 (range 7-156 participants); with smaller sample 200 sizes noted in papers comparing energy intake to DLW (range 7-19 participants). A range of sports 201 were represented most of which were 'mixed sports' (6 studies; n=461 participants) with the 202 remainder reporting on athletes from 'aerobic' or 'endurance sports' (e.g. long distance running, 203 swimming, rowing, triathlon, cross country skiing) (9 studies; n=184), 'team sports' (e.g. soccer, 204 basketball) (2 studies; n=26), and 'skill' or 'aesthetic sports' (1 paper; n=12 ballet dancers). The mean 205 age was 21.8 ± 2.6 years (range 16-30.4 years). Two studies did not report participant age however it 206 was assumed the participants were older than 16 years of age due to recruitment from tertiary 207 institutions [41,53]. More than half of the studies were in male athletes (55 % participants). Athletic 208 caliber included competitive tertiary, national, international and professional level athletes. The 209 studies were conducted in Europe (n=6), North America (n=6), and across the Asia Pacific and African 210 regions (n=6) (Table 1). 211 Physique characteristics of the athletes are summarized in Table 1. The weighted mean body 212 mass and stature were  $65.3 \pm 7.2$  kg and  $173.3 \pm 6.1$  cm, respectively. One paper did not report mean 213 stature of participants [50]. Seven studies reported percentage body fat with mean levels ranging 214 from  $7.1 \pm 2.5$  to  $22.8 \pm 5.1$  % (Table 1).

Reference, country	Group (n)	Sport, calibre	Age (years)	Body Mass (kg)	Stature (cm)	BMI (kg/m²)	Body fat (%)	Comments
Baker et al. (2014), USA [49]	56 (41 M, 15 F)	Mixed sports <sup>1</sup> , competitive, tertiary	16 ± (14 - 20)	69.4 ± 14.3	174.3 ± 9.4			n=12 excluded (non-adherence)
Braakhuis et al. (2011), USA [50]	113 (56 M, 57 F)	Rowers, national competitive	22 ± 3 (17 - 36)	78.0 ± 11.0	NR			n=2 excluded (EI < 1.39*RMR)
Ebine et al. (2000), Japan [40]	9 F	Synchronised swimmers, national	19.8 ± 2.8 (16 - 23)	52.5±2.7	159.0 ± 3.0	20.7 ± 0.7		DLW
Ebine et al. (2002), Japan [21]	7 M	Soccer, professional	22.1 ± 1.9	69.8 ± 4.7	175.0 ± 5.0		13.4 ± 3.6	DLW
Edwards et al. (1993), USA [41]	9 F	Cross-country runners, highly trained, tertiary	NR#	55.3 ± 6.2	169.1 ± 5.5	19.3 ± 1.7	13.0 ± 3.2	DLW
Fogelholm & Lahti-Koski (1991), Finland [51]	84 M	Mixed sports <sup>2</sup> , recreational	24 ± 4	72.0 ± 7.0	180.0 ± 6.0			n=12 excluded (incomplete data)
Fudge et al. (2006), Kenya [42]	9 M	Middle & distance runners, elite	21 ± 2	56.0 ± 3.4	174.0 ± 2.9	18.3 ± 1.3	7.1 ± 2.5 (BIA)	DLW
Hill and Davies (1999), Australia [43]	12 F	Ballet dancers, tertiary	18.7 ± 1.2	57.3 ± 3.8	169.1 ± 4.6			DLW
Hill and Davies (2002), Australia [44]	7 F	Lightweight rowers, elite	20.0 ± 1.1	60.9 ± 2.3	168.8 ± 4.7		22.8 ± 5.1	DLW n=1 excluded (incomplete data)

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Table 1 Participant characteristics of athletes from included studies

Reference, country	Group (n)	Sport, calibre	Age (years)	Body Mass (kg)	Stature (cm)	BMI (kg/m²)	Body fat (%)	Comments
Jones and Leitch (1993), Canada [22]	(5 M, 3 F)	Swimmers, tertiary	20.1 ± 1.7	74.1 ± 9.3	186.0 ± 11.0		16.0 ± 6.4	DLW n=1 excluded (disagreement between measures)
Koehler et al. (2010), Germany [45]	12 M	Triathletes, well- trained*	30.4 ± 6.2	80.6 ± 6.5	186.0 ± 8.0	23.2 ± 1.4	11.2 ± 2.1 (BIA)	DLW n=2 excluded (change BW >3%; EI < 1.39*RMR)
Schulz et al. (1992), USA [46]	9 F	Distance runners, elite, national	26.0 ± 3.3	52.4 ± 4.1	163.0 ± 7.0		12 ± 3 (UWW) 17 ± 3 (BIA)	DLW
Scoffier et al. (2013), France^ [52]	(13 F, 9 M)	Mixed sports* Adolescent, weight sports	25.3 ± 4.7	59.7 ± 15.7*	169.8 ± 2.1*			
	20 (12 F, 8 M)	Adolescent, other sports	$26.4 \pm 5.7$	62.1 ± 10.7*	168.8 ± 2.7*			
Silva et al. (2013), Portugal [47]	19 (12 M, 7 F)	Basketball, junior national, elite	17.0 ± 0.7 (M) 16.9 ± 0.7 (F) (16 - 18)	74.7 ± 10.8	185.4 ± 11.0	21.7 ± 1.9		DLW
Sjodin et al. (1994), Sweden [48]	8 (4 M, 4 F)	Cross-country skiers, elite, national	26 ± 2 (M) 25 ± 2 (F)	75.1 ± 4.9 (M) 54.4 ± 5.1 (F)	180.0 ± 6.0 (M) 166.0 ± 2.0 (F)			DLW
Sunami et al. (2016), Japan [53]	156 (92 M, 64 F)	Mixed sports <sup>3</sup> , tertiary	NR#	68.7 ± 8.4 (M) 56.1 ± 5.9 (F)	174.7 ± 6.5 (M) 163.4 ± 5.8 (F)	22.5 ± 2.1 (M) 21.0 ± 1.5 (F)		
Ward et al. (2004), USA [29]	76 F	Mixed sports <sup>4</sup> , Division I and III NCAA, tertiary	18.8 ± 1.1 (17 - 21)	59.8 ± 7.1*	165.6 ± 6.1*	21.7 ± 2.3		

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Data are presented as mean  $\pm\,\mathrm{SD}$  or mean (range) unless otherwise indicated.

M mile, F femile, BMI body mass index, DLW doubly labeled water study, NR not reported, BIA bioelectrical impedance, BW body mass, EI energy intake RMR resting metabolic rate, UWW underwater weighing, NCAA National Collegiate Athletic Association.

Meter de porte seccer, tennis, hastrball, football, god, laureses, baseball, softball, track and fell following on the secessing, being, jee bockey, skating, dance Mittel, gener tendances specific policies porter industries of porter baskball, care seccer, hastrball, Park handlil, job, to series, vollephall. Meter sporter. baskball, care seccer, hastrball, part hastrball, care seccer, baskball, secret, vollephall. Meter deporte TAEF, cycling, archery, speed skating, short-nack skating. \*Africe calibre and recontinent source no seccession, boday, soccer, vollephall. Meter deporte TAEF, cycling, archery, speed skating, short-nack skating. \*Africe calibre and recontinent source no.

^ Data for the non-athlete adult and adolescent participants excluded. \* Anthropometric data converted to metric units. # Participants assumed >16 years due to recruitment from College sporting clubs or University team.

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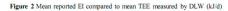
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3.3 Studies Comparing Reported Energy Intake to Energy Expenditure as Measured by DLW

Eleven studies assessed the difference between reported mean EI and mean TEE as measured by DLW (Figure 2). The most common method for measuring EI was by weighed or estimated food records (FR) over 4-10 days (mean 6.5 days) duration (Table 2). Where reported (*n*=5), the mean macronutrient intake (as percent of energy intake) was 56.5, 14.9 and 26.8 % for carbohydrate, protein and fat, respectively. One paper reported mean protein intake only (12.6% of energy intake) [45].



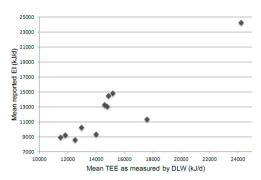


Figure 2 Mean reported EI compared to mean TEE measured by DLW (kJ/d)

Energy expenditure was measured for a mean of 8.4 days (range 6-14 days) by the DLW method (Table 2). Details on the training phase or competitive season of the participants were reported by six studies [21,22,40,42,43,47]. Six studies also recorded physical activity levels during the assessment period either by self-reporting the number of training hours via an activity record (n=5) [22,41,45,46,48], or as measured by ActiGraph<sup>TM</sup> activity monitor (n=1) [42].

Eight of the eleven studies measured dietary intake for the entire DLW measurement period [21,22,40-42,45-47]; while three studies measured intake for only 4-5 days [43,44,48]. Most studies (*n*=7) reported a stable body mass during the testing period, however four studies adjusted energy for changes noted in body mass during the testing period [43-45], or for poor dietary records as a result of participant non-compliance [22] (Table 2).

Resting metabolic rate (RMR) was measured via indirect calorimetry in four studies [40,43,46,47]; four reported RMR based on prediction calculations (i.e. Schofield, Cunningham or FAO/WHO/UNU equations) [21,42,44,48]; while three studies did not report RMR [22,41,45] (Table 2).

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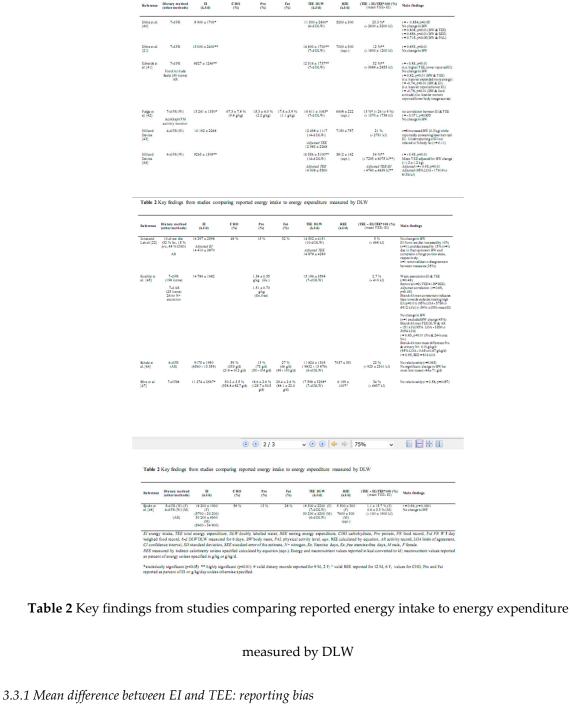
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Table 2 Key findings from studies comparing reported energy intake to energy expenditure measured by DLW

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Compared to DLW, all studies reported a lower mean EI of 19 % (range 0.4 - 36 %) with a weighted mean difference of  $-2793 \pm 1134$  kJ/d between measures (Table 3). The difference was significant in seven studies [21,40-42,44,45,47] while smaller, non-significant differences were reported in the remaining studies [22,43,46,48]. For example, Hill and Davies [44] found a very strong

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kJ/d (Table 2).

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inverse correlation of difference between TEE and self-reported EI, and the means of the two measurements (r=-0.93, p<0.01) in elite female lightweight rowers. When mean EI was adjusted for change in body mass (- 1.2 ± 1.2 kg) over the measurement period, the correlation of difference between the mean adjusted TEE and self-reported EI was also significant (r=-0.93, p<0.01). The limits of agreement (LOA) (95 % CI ± 2 SD) for the mean adjusted TEE and self-reported EI were – 17 619 to 8134 kJ. The authors also noted that reporting bias increased in those with higher TEE (i.e. TEE > 3000 kcal/d; or 12 552 kJ/d). In contrast, Schulz and colleagues [46] found no relationship between self-reported EI and TEE in female distance runners and reported a mean difference of  $-925 \pm 2301$ kJ/d. Similarly, Sjodin and colleagues [48] noted a close match between reported EI and TEE (i.e. mean difference of  $-100 \pm 1900 \text{ kJ/d}$ ) in eight cross-country skiers (Table 2). Koehler and colleagues [45] aimed to assess the validity of a food and activity record compared to reference methods such as DLW, 24-h urea nitrogen excretion and indirect calorimetry (i.e. to determine energy expenditure during an incremental running or cycling test). They found EI and TEE as measured by DLW were only weakly correlated, however following the exclusion of two implausible results (where EI < 1.39 x REE) [55] a significant positive correlation was observed (r=0.69, p<0.05). A proportional bias towards over-estimating low and under-estimating high energy intakes (p<0.01) was also noted with very wide LOA (95 % ± 2 SD) between EI and TEE of - 5736 and 4912

n	M	F	EI (kJ) (± SD)	TEE (kJ) (± SD)	(TEE-EI kJ)	Weighed mean difference (%)
9		9	8900 (1700)	11 500 (2800)	22.6 (2600 kJ)	203
7	7		13 000 (2400)	14 800 (1700)	12.2 (1800 kJ)	85
9		9	8527 (1246)	12 516 (1737)	31.9 (3989 kJ)	287
9	9		13 241 (1330)	14 611 (1043)	9.4 (1370 kJ)	84
12		12	10 192 (2268)	12 983 (2268)	21.5 (2791 kJ)	238
7		7	9263 (1309)	14 008 (5560)	33.9 (4745 kJ)	237
8	5	3	14 410 (3870)	14 878 (4289)	3.1 (469 kJ)	25
12	12		14 786 (1682)	15 196 (3598)	2.7 (410 kJ)	32
9		9	9176 (1950)	11 824 (1305)	22.4 (2648 kJ)	202
19	12	7	11 274 (2567)	17 598 (3298)	35.9 (63 24 kJ)	683
8	4	4	24 200 (3250)	24 250 (3200)	0.4 (100 kJ)	3
109	49	60				
	9 7 9 12 7 8 12 9 19	9 7 7 9 9 9 12 7 8 5 12 12 9 19 12 8 4	9 9 9 12 12 12 9 9 19 12 7 8 4 4	9 9 9 8527 (246) 9 9 8527 (246) 9 9 8527 (246) 9 9 13 241 (1330) 12 12 10 192 (256) 7 7 9203 (1399) 8 5 3 14 410 (3170) 12 12 14 756 (1652) 9 9 9176 (1650) 19 12 7 11 274 (2567) 8 4 4 24 200 (259)		N

Table 3 Calculation of weighted mean difference between reported EI and TEE measured by DLW

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3.4 Meta-Analysis

Studies which involved the comparison of a dietary measure (i.e. self-reported or weighted FR) to TEE assessed by the DLW technique were homogenous enough to conduct a meta-analysis (*n*=11). Relevant data (i.e. mean EI, SD and sample size) were used to calculate the between-group standardised mean difference, or effect size (ES) and 95 % confidence intervals (CIs). The meta-analysis revealed a large pooled ES of -1.006 (95 % CI - 1.3 to -0.7; p=0.00) where reported EI was significantly below TEE measured by DLW (mean - 2477 kJ/d) (Figure 3).

Figure 3 Meta-analysis of the pooled effect of self-reported EI compared to TEE measured by DLW

3.5 Studies Comparing Reported Dietary Intake by Two or More Methods of Dietary Assessment

Of the seven studies comparing reported intake by two or more methods of dietary assessment, four compared energy and/or macronutrient intake [49,51-53], while the remaining three studies compared the intake of a specific nutrient (i.e. antioxidants, calcium, or protein) [29,50,54]. Only two studies validated the test method (i.e. FFQ, modified FFQ) for micronutrients and food groups [51,53] (Table 4).

Four studies investigated the validity of a new self-administered, semi-quantitative or quantitative food questionnaire (i.e. FFQ or modified FFQ) in an athlete cohort. The dietary reference methods used to evaluate these new instruments included a FR (n=3) [29,50,51] or multiple pass 24-h DR (n=1) [53]. Braakhuis and colleagues [50] compared a blood biomarker of antioxidant capacity (i.e. ferric-reducing ability of plasma [FRAP]) with antioxidant intakes recorded from a 7-d weighed FR and FFQ. The remaining three studies aimed to validate a new dietary assessment instrument (e.g. web-based 24-h DR, virtual interface tool). The dietary reference methods used to evaluate these new instruments included a FR (n=1) [52]; multiple pass 24-h DR (n=2) [49,54]; and/or direct observation (n=1) [49]. Wardenaar and colleagues [54] also compared self-reported protein intake as measured by 24-h DR to urinary nitrogen excretion (Table 4).

Table 4: Studies comparing reported intake as measured by two or more dietary assessment methods

Reference	Dietary method	Reference method(s)	EI (kJ/day)	CHO (g)	Pro (g)	Fat (g)	Main findings
Baker et al. (2014) [49]	DATA ipad administered modified multiple pass 24-h DR (n=56) (pre-test DATA, n=19)	INTERVIEW 24-h DR (=56) 085ERVATION RD observed 24-h (=26)	DATA 14 656 5945 ki* OSS. 12 728 ± 5280 ki* 13 870 ± 6117 kj NTERVIEW 14 041 ± 6627 kj	DATA 475 ± 190 OBS. 426 ± 159 DATA 449 ± 205 INTERVIEW 449 ± 216	DATA 151 ± 59 OBS 139 ± 63 DATA 140 ± 62 INTERVIEW 147 ± 77	DATA 116 ± 65* OBS. 91 ± 53* DATA 112 ± 73 INTERVIEW 112 ± 61	Significant difference between DATA & Olis. Some energy, CILO, Pro. 6, aute, scalulur, micro, elevier (ICC O.7s 0.91)  A service of the control of the co
Braakhuis et al. (2011) [50]	FFQ (70 items)	FR (7d, W) (n=81 FFQ & FR) Biomarker (FRAP) (n=96 FFQ & FRAP) (n=63 FR & FRAP)	14 500 ± 5700 (7500 - 25 900)	470 ± 190 (53.5 ± 6.8 %)	150 ± 80 (17.9 ± 5.8 %)	114 ± 54 (28.0 ± 6.2 %)	Modest correlation between FR and FFQ arterioxchain intake (PG) 3 = 0.14, 90(PG) 3 = 0.14, 90(PG) 5 = 0.14, 90(PG) 5 = 0.13, 90(PG) 5 = 0.14,

Reference	Dietary method	Reference method(s)	EI (kJ/day)	CHO (g)	Pro (g)	Fat (g)	Main findings
Fogelholm & Lahti- Koski (1991) [51]	FUQ (122 items) (FUQ1 participant reported portion size; FUQ2 medium portion sizes)	FR (7d, W)	13 000 ± 2800	397 ± 123	122 ± 31	114 ± 30	Close agreement El Serveen PK & FLOI & FLOZ EF RR FLOI (95%; CL -1.70 to 1.2 to 1.77 ± 0.0) Additional finding to 0.00 to 1.77 ± 0.0) Additional finding to 0.00 to 0.00 to 1.00 to 1.
Scoffier et al. (2013)^ [52]	VSSR	FR (Id)	Adolescent athletes, weight sports VSSR 7978 ± 2513 kJ FR 7491 ± 2116 kJ Adolescent athletes, other sports VSSR 7190 ± 2004 kJ FR 7081 ± 1785 kJ				NS between VSSR & FR (adolescent athletes, weight sports) (p=0.11); or between VSSR & FR (adolescent athletes, other sports) (p=0.56)
Sunami et al. (2016) [53]	FFQ (138 food, 20 beverage, 14 seasoning items)	24-h DR (3d, non- consecutive)	24-hDR 13 332 ± 3933 kJ (M) 8962 ± 2117 kJ (F) FFQ 12 159 ± 4996 kJ (M) 8029 ± 2519 kJ (F)	24-hDR 486.6 ± 152.9 (M) 319.9 ± 81.8 (F) FFQ 452.4 ± 182.8 (M) 286.2 ± 76.0 (F)	24-hDR 100.1 ± 32.5 (M) 65.5 ± 16.4 (F) FFQ 82.0 ± 35.5 (M) 59.0 ± 22.8 (F)	24-hDR 83.7 ± 34.2 (M) 64.0 ± 18.6 (F) FFQ 76.7 ± 40.6 (M) 57.1 ± 28.6 (F)	FFG underesterated EI by 9% M and 10% F Majority natries within = 20 % range; targest difference for retinol (77% M, 32% F). Additional fidality and contenuated CC was 0.30 (0.10 to 0.5 %) Mad 0.32 c. 0.08 to 0.62; (7) For 19 food groups median destremanted CC was 0.30 (1.00 to 0.5 %) Mad 0.32 c. 0.08 to 0.62; (7) Lower difference was noted for cereative, separables fung fixed groups, while greater differences noted from super. Neurologies, seasoning and aptices.
Ward et al. (2004) [29]	RAM calcium checklist (54 items)	FR (6d)	Mean calcium (mg/d) RAM 822 ± 331 FR 823 ± 387				Test-retest reliability of RAM was moderate ICC=0.54 (p=0.0001, r=0.58) RAM had moderate agreement with FR ICC=0.41 (p=0.0067, r=0.42). RAM correctly identified 84 % with low calcium

Reference	Dietary method	Reference method(s)	EI (kJ/day)	CHO (g)	Pro (g)	Fat (g)	Main findings
Wardenaar et al. (2015) [54]	Compl-ear <sup>TM</sup> 24-h DR (3d, non-consecutive)	24-h urinary N+ excretion; Q (training load, sports foods, dietary supplements)	16 900 ± 4200 (8540 - 26 600)		24-hDR* 109.6 ± 33 (1.49 ± 0.35 g/kg/d) 24-h N+ excretion* 141.3 ± 38.2 (1.9 ± 0.39 g/kg/d)		Significant mean difference of 25.5 ± 21.3 % ( -(3.17. ± 0.9 0.9 between 24-4b.R) and 24-h.N- excretion (pc0.001) (m0.65; 95% CI 0.45-0.79) Additional findings: Underestimation of Pro related to amount of protein intake (m-0.00; 95% CI 0.46 to 0.09) Mean PII. 1.6 ± 0.4 with Pis. 7 % unbress FIS.1-7.5 indicating possible under-exporting Underreporting greater in individuals with higher protein intakes than with lower intakes.

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**Table 4** Studies comparing reported dietary intake as measured by two or more methods of dietary

300 assessment

3.5.1 Reported mean energy intake

Six of the seven studies reported mean EI and/or compared intake between dietary assessment methods (Table 4). Sunami and colleagues [53] compared mean EI of college athletes from a variety of sports from a FFQ (138 items) with 3-d non-consecutive 24-h DR and found the FFQ underestimated EI by 9 % (males) and 10 % (females). In comparison, Fogelholm and Lahti-Koski [51] found close agreement on a group level of mean EI of mixed sport athletes from a food use questionnaire (FUQ) compared to a 7-d weighed FR (95% CI -1.7, 0.1 FUQ1, 95% CI -0.1, 1.7 FUQ2); however, individual agreement was weak. Scoffier and colleagues [52] found no difference between EI measured by a virtual interface dietary assessment tool compared to a 1-d FR in two groups (i.e. weight sports, and other mixed sports) of adolescent athletes. Baker and colleagues [49] compared reported energy and macronutrient intakes measured by a web-based, 24-h DR tool (DATA) to a 24h DR (interview) and observation (by registered dietitian), and found no significant difference between the measures. They found good relative validity for group level comparisons but large variations of individual dietary intake estimates, especially in athletes with higher energy and macronutrient intakes (Table 4).

316 3.5.2 Reported mean macronutrient intake

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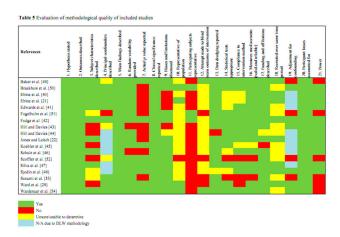
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Where reported (4 of 7 studies), the mean macronutrient intake (as percent of EI) was 53.1, 15.3 and 28.5 % for carbohydrate, protein and fat, respectively (Table 4). An additional study reported mean protein intake only (11.9 % of EI) [54]. Wardenaar and colleagues [54] reported a mean difference between 24-h DR and 24-h nitrogen excretion (25.5  $\pm$  21.3 % difference, or 31.7  $\pm$  30 g/d; p<0.001). Under-reporting of protein was related to the amount of protein intake (r=-0.20; 95% CI -0.46, 0.09) such that under-reporting was greater at a higher protein compared to a lower protein intake. In contrast, Koehler and colleagues [45] found protein intake in triathletes was strongly correlated to 24-h urinary nitrogen excretion (r=0.81; SEE=0.34g/kg/d), however considerable individual variation was observed between the two methods with very wide limits of agreement (-0.65 to 0.67 g/kg/d). 3.5.3 Other nutrients, food groups and dietary patterns reported Braakhuis and colleagues [50] found a modest correlation between energy-adjusted estimates of total antioxidant intakes in competitive rowers as measured by a quantitative FFQ (70 items) compared to 7-d weighed FR (r=0.38, 90% CI ± 0.14), and only a small correlation between the FFQ and plasma biomarker (r=0.28, 90% CI ± 0.15). However, the authors noted a trend to over or underestimate antioxidant intakes at low and high intake levels (42 % over and 73 % under-estimation, respectively). Ward and colleagues [29] compared mean calcium intake as measured by a selfadministered calcium checklist to a 6-d FR (822  $\pm$  331 mg/d and 823  $\pm$  387 mg/d, respectively) and observed no significant difference between the two methods (Table 4). In addition to comparing energy and macronutrient intake between the test method (i.e. FUQ1 and FUQ2) and a 7-d weighed FR, Folgelholm and Lahti-Koski [51] reported on selected

micronutrients (i.e. thiamin, vitamin C, calcium, magnesium, iron and zinc). They found the FUQ1

339 over-estimated reported intake of vitamin C, calcium, magnesium, iron and zinc; while the FUQ2 did 340 not differ in mean values for the selected micronutrients compared to a 7-d FR. Most of the 341 correlations by food group were reported above r = 0.24, with the exception of 'vegetable oils', 342 'cream', 'milk', 'pork', 'beef', and 'poultry' [51]. 343 One study validated the test dietary assessment method (i.e. FFQ) compared to a 3-d non-344 consecutive 24-h DR for 35 nutrients and 19 food groups [53]. While the difference in measures for 345 the majority of nutrients was within ± 20 %, the largest difference was observed for retinol (i.e. 77 % 346 for males and 32% for females). The authors found lower differences between measures for 'cereals' 347 (1 % in males, 10 % in females), 'vegetables' (9 % in males, 4 % in females) and 'fungi' (5 % in males, 348 9 % in females) food items; while greater differences were noted for 'sugar' (- 94 % in males, - 96 % 349 in females), 'beverages' (-45 % males, - 54 % females), and 'seasonings and spices' (-65 % males, -72 350 % females) food items. Overall, under and over-estimation of food group intakes was more prevalent 351 than that of nutrient intakes [53] (Table 4). 352 3.6 Evaluation of Methodological Quality 353 The methodological quality of all included studies (n=18) were evaluated with quality ratings 354 determined as fair to moderate for the majority of studies (Table 5). All papers stated the aim or 355 hypothesis, described the main outcomes to be measured, described the main findings, reported 356 clinical and statistical significance, and accounted for participant drop outs. All but two studies 357 provided estimates of the random variability [43,46]; reported on data dredging [44,48]; and all but 358 two reported sources of funding [44,45]. However, the poorest ratings were for the following items: 359 "Were actual probabilities reported?" (Item 7); "Were the subjects asked to participate representative of the 360 entire population?" (Item 10); "Were the participating subjects representative of the population they were

recruited from?" (Item 11); and "Were the subjects recruited over the same time period?" (Item 18) Only two studies [49,54] reported whether an attempt was made to blind those measuring the main outcomes of the intervention (i.e. Item 12) (Table 5).



**Table 5** Evaluation of methodological quality of included studies

### 4. Discussion

This is the first systematic review to have evaluated studies comparing two or more dietary assessment methods, including measuring intake against biomarkers or reference measures of energy expenditure, in athletes. Only 18 papers were published over the past 37 years, confirming the limited literature in this area. Most papers (*n*=11) focused on self-reported EI compared to TEE as measured by the DLW technique [21,22,40-48]. The remaining studies (*n*=7) compared a dietary assessment method to a reference method(s) (e.g. food record, 24-h DR, direct observation, biomarker) as part of a validation study [29,49-54]. Dietary assessment methods recognized as appropriate for the general population are usually applied in a similar manner to athlete groups, despite the knowledge that athlete-specific factors can complicate the assessment and impact accuracy in unique ways. As dietary assessment is used extensively in both sports nutrition clinical and research settings, it is a concern the validity of the methodologies used have not undergone more rigorous evaluation. There

is a clear need for high quality research to be undertaken to identify dietary assessment methodologies which are valid, as well as feasible for use, in an athlete population.

4.1 Studies Comparing EI to TEE as Measured by DLW

Overall, the studies (*n*=11) comparing self-reported EI to TEEdLW found mean EI was underestimated by 19% (range 0.4-36%) which is comparable to differences observed in studies in other populations [18-20,25,26,56,57]. Three DLW studies [22,45,48] reported close agreement (<5% difference between measures); which can be partially explained as participants were provided with a test diet for the duration of the DLW collection and a participant was excluded due to an inexplicable marked difference in agreement between measures [22]. Koehler and colleagues [45] adjusted mean EI following removal of participants (*n*=2) for loss of body mass (>3%) or due to implausible food records as determined by a cut-off value (i.e. EI<1.39\*RMR) [55]. While Sjodin and colleagues [48] noted a close agreement between mean EI and TEEdLW over 7 days (r=0.96; p=0.0001); no relationship was observed when authors compared EI from separate 24-h periods, indicating athletes were not in energy balance during shorter periods of time.

The remainder of the DLW studies (*n*=8) reported a greater mismatch between self-reported EI and TEEdLW (12-34% difference) attributed primarily to misreporting [21,40-44,46,47]. A variety of explanations have been proposed for lower reported EI when compared to TEEdLW [21,25,40,46]. Hill and Davies [18] suggest factors such as body size, perception of body image, restrained eating, gender, socioeconomic status, motivation, social expectations and the nature of the testing environment itself, play a role in misreporting. Nutrition-related beliefs and dietary practices of athletes present additional challenges with reporting intake accurately in this population sub-group. In addition, factors such as high energy requirements, frequency of snacking, eating away from the

home environment, applicability of 'standard' portion sizes, and the wide use of commercial sports foods, drinks and supplements can make it more difficult to quantify food intake [8,19].

## 4.1.1 Methodological Issues

The DLW technique has long been considered a 'gold standard' for measuring TEE and to validate self-reported EI [20]. While the DLW technique is considered accurate to 1%, with a coefficient of variation of 2-12%, depending on the loading dose and length of sampling period [20,58]; potential sources of error should be considered when measuring in free-living conditions including the influence of day-to-day variation of EI and TEE [58]. The optimal duration for measuring DLW is between 2-3 biological half-lives of the isotopes [59]; although a shorter period (i.e. 8-12 days) has been suggested for athlete cohorts due to higher rates of water turnover due to regular physical activity [59]. A potential limitation is the DLW measurement period of most of the included studies (*n*=8) may have been too short (i.e. 6-7 days duration) [21,40-42,45,46-48].

## 4.1.2 Assessment of Dietary Intake

Typically dietary intake is measured in DLW studies to assist in the calculation of the respiratory quotient (RQ); a required variable in calorimetry equations [60]. This calculation was reported in the methodology of only four of the included studies [22,40,46,47]. In non-athletes, a 3–4d FR is considered valid for assessment of RQ and EI for groups [61]; however dietary intake of athletes may be more variable due to the day-to-day variation in the energy cost of training. It has been suggested that recording intake for 3-7 days is a reasonable compromise between scientific rigor and practicality when estimating dietary intake of athletes to capture habitual intake and high variability in day to day energy expenditure [19,28,62]. Hence, for athletes, ideally intake would be recorded over the

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entire DLW measurement period to capture this variation. As such, a further limitation is that three studies did not record intake for the same duration as the DLW measurement [43,44,48]. In all of the included DLW studies, dietary intake was determined by weighed or estimated FR (recorded for 4-10 days). The documentation of intake via FR is considered practical and is widely used in both clinical and research settings, however the accuracy of data is reliant on subjective recording of intake, level of participant motivation, and possible fatigue from recording for longer durations [18,19,25]. Recording intake itself can modify usual eating behavior [46,63]. Investigators are responsible for improving compliance by providing clear, specific instructions to ensure records provide sufficient detail. Nine studies provided detail about instructions provided to participants and review process of dietary records upon collection [21,40-44,46-48]. Only one study supervised the weighed FR over a limited observation period in order to improve recording accuracy [48]; while another provided a test diet to all participants [22]. Given the potential for misreporting during the recording period, the plausibility of selfreported EI can be compared to a pre-determined EI:BMR value [55]. Two DLW studies applied a cut-off value to exclude implausible dietary records [44,45], and one of the validation studies [50]. However, when applying a cut-off value to assess self-reported EI of athletes, it is important to have a valid assessment of the energy demands of the athletes being assessed, as the range in TEE can be wide and is reliant on the type of sport, training phase and intensity [19,56,64]. 4.1.3 Variability of Intake and Expenditure in Athletes Although the DLW studies investigated energy balance or misreporting of intake, this was not the primary aim of all included papers. A number of the studies (n=4) aimed to determine TEE of a group of athletes (i.e. swimmers, soccer players, light weight rowers, distance runners) and compare

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to recommendations [21,40,44,46]; with the secondary outcome to evaluate the agreement between reported EI and TEE<sub>DLW</sub>. The primary aim of the remaining papers (n=7) was to compare the difference between EI and TEE<sub>DLW</sub> [22,41,45,47]; assess energy balance in a group of athletes (i.e. endurance runners, cross-country skiers) [42,48]; and assess the validity of a 4d FR in measuring EI of female classical ballet dancers [43]. While DLW is highly regarded as an accurate method for the validation of reported EI [64], a major limitation is that the technique is unable to calculate energy turnover on a daily basis. In athletes, TEE (and EI) can fluctuate substantially from day to day [65,66]. For example, Bradley and colleagues [65] found EI of rugby union players lowest when TEE was highest early in the training week, with EI increasing in preparation for competition on the weekend. Similarly, Brown and colleagues [66] noted differences in EI, macronutrient intake and energy balance of dancers between weekdays and weekend days. These recent publications highlight the day-to-day variability in energy demands of athletes over a relatively short period of time which is unable to be assessed by DLW. 4.1.4 Influence of Body Mass, Body Image and Energy Demands In most studies there was no significant change in body mass during the assessment period [21,22,40,41,45-48] indicating the difference between measures was likely due to misreporting. Conversely, Fudge and colleagues [42] attributed the difference noted between EI and TEEDLW (-13%) in elite Kenyan runners to under-eating resulting in a negative energy balance during a period of intense training. Four studies adjusted mean TEEDLW based on changes in body mass during the measurement period [22,43,44,46]. Previous research has indicated the degree of misreporting increases with increasing body mass, specifically adiposity [67,68] and BMI (body mass index) [69]. Others suggest that misreporting is independent of adiposity but linked to restrained eating, and

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body image [18,70-72]. In the current review, Edwards and colleagues [41] observed a relationship

466 between EI and body mass (r=-0.74) (i.e. heavier women reported a lower relative intake). However, 467 Hill and Davies [43] found no relationship between the extent of misreporting and percent body fat 468 of female ballet dancers (r=0.11). Similarly, Silva and colleagues [47] found no relationship between 469 EI and body mass or composition in male and female basketball players. 470 It appears the trend for misreporting increases with increasing TEE, particularly for individuals with 471 high energy needs [19,57]. Possible reasons include increased burden from reporting large volumes 472 of food and frequent eating occasions resulting in food or drink items being omitted consciously or 473 unconsciously. In the current review, a negative correlation was observed between self-reported EI 474 and TEEDLW in some studies (r=-0.854, p<0.01) (i.e. the higher the expenditure, the lower the reported 475 intake) [40,41]. While there was almost no difference between self-reported EI and TEEplw in a group 476 of triathletes, the data indicated a strong proportional bias towards under-estimating high energy 477 intakes (p<0.01) [45]. Similarly, Hill and Davies [44] reported a bias to misreporting with increasing 478 EE (i.e. TEE > 3000 kcal/d; or 12 552 kJ/d) in female lightweight rowers. This trend is supported by 479 findings in non-athletes where the magnitude of misreporting intake increases with increasing TEE 480 [57,73,74]. 481 4.2 Studies Comparing Dietary Intake by Two or More Methods of Dietary Assessment 482 The current review identified a limited number of studies (n=7) which evaluated the validity of

The current review identified a limited number of studies (*n*=7) which evaluated the validity of a novel dietary assessment tool or instrument compared to one or more reference method(s) [29,49-53] and/or a biomarker in an athlete population [50,54]. However, due to limited consistency in the pairing of study methods, it was difficult to make firm conclusions on the relative strength of the different methodologies used to evaluate dietary intake in athletes. Despite most studies reporting

acceptable validity for group level comparisons [49,51,54], individual agreement was not as strong,

488 especially for athletes with high energy intakes [49,50,54]. A variety of statistical tests were used to 489 assess study validity (e.g. paired t-tests, correlation coefficients, Bland-Altman plots), however most 490 studies used correlation coefficients [50,53] or intra-class correlation (ICC) to assess reproducibility 491 [29,49]. Correlation coefficients can be misleading, as they measure the relationship between to 492 methods, rather than the agreement between them [75]. Only one study validated the test method for 493 food groups [53], while the other studies evaluated energy and/or nutrient intakes [29,49-52,54] 494 4.2.1 Dietary Reference Methods 495 Three of the included studies determined dietary intake by (weighed or estimated) FR (6-7 days) 496 [29,50,51]; while another recorded intake for a single day only [52]. Research suggests that a longer 497 recording duration is necessary to assess habitual intake and account for day-to-day variation of 498 intake and TEE in athletes, with a 7-d FR two to three times less variable than a 1-d FR [28]. Other 499 reference methods which rely on self-reporting (e.g. 24-h DR, FFQ) are also prone to measurement 500 error associated with recall bias and awareness of portion size [19,53,76]. Three studies used the 501 multiple pass 24-h DR [49,53,54] and/or direct observation [49] to evaluate a new tool or instrument. 502 While 24-h DR has the advantage of low participant burden, data may not be representative of the 503 usual diet unless the recalls are repeated a number of times [4,19]. In addition, Wardenaar and 504 colleagues [54] suggest dietary data should be collected within a four week period to provide the best 505 insight into the accuracy of the multiple 24-h DR in athletes, due to the micro-cycle and periodization 506 of training (i.e. training programs with a variable workload and volume throughout the year). It has 507 been previously suggested that the 24-h DR is prone to underestimate EI, and that caution should be 508 taken assessing data involving high intakes [4,19,77]. In general, all dietary assessment methods are

included in the current review failed to detect true reporting bias, if both the new and established dietary assessment instruments have correlated error [20], particularly when applied to an athlete population. Researchers are encouraged to consider the relative strengths and weaknesses of the dietary reference method(s) selected, in addition to consider participant burden, cost, and validity in the study population [53].

## 4.2.2 Evaluation Using Biomarkers

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Two of the included validation studies used a biomarker to assess the accuracy of reported intake [50,54]. Wardenaar and colleagues [54] found a multiple 24-h DR acceptable for ranking protein intake compared to the reference urinary nitrogen biomarker (r=0.65). However, they noted the 'standard' portion sizes used in the 24-h DR may not be representative for athletes with high intakes of protein and therefore a potential source of error [54]. Braakhuis and colleagues [50] found only a weak association between the plasma antioxidant biomarker (FRAP) and FFQ (r=0.28). However relying on blood biomarkers to validate a nutrition questionnaire can be problematic as there is no single marker for antioxidant intake [50]. Unfortunately, the use of independent biomarkers to assess the accuracy of dietary intake is limited to energy intake or a specific nutrient only (i.e. sodium, nitrogen). Koehler and colleagues [45] indicate that their research first to validate dietary protein intake against 24-h urinary nitrogen excretion in an athlete population. They found good agreement between 24-h nitrogen excretion and dietary protein intake (r=0.81) which is consistent with results documented in non-athlete populations [56,64,76]. More recently, energy availability (EA) (i.e. EA=EI-ExEE) has been linked to biochemical indices such as cortisol, insulin,

530 growth hormone, IGF-1, leptin and thyroid hormones [45,78]. These present potential use as 531 biomarkers in future dietary assessment studies. 532 4.2.3 Nutrients, Food Groups and Dietary Patterns 533 Micronutrient intake and food group intake is less frequently reported in dietary validation 534 studies than energy and macronutrient intakes [38]. One paper evaluated calcium intake in female 535 college athletes from a range of sports (e.g. basketball, cross-country) using a 6-d weighed FR to 536 validate a self-administered calcium checklist [29]. Mean calcium estimates did not differ between 537 measures, however longer periods of recording may be required to estimate intake of key 538 micronutrients such as iron and calcium (i.e. up to 11 recording days) [19,61,62,77]. In another study, 539 Braakhuis and colleagues [50] found an FFQ was valid for estimating antioxidant intake in elite 540 rowers. 541 Two studies reported on a range of selected micronutrients [51,53] as part of validating test 542 methods (i.e. FUQ, FFQ, respectively). These same studies are the only studies in athletes to validate 543 dietary intake at the food group level. Sunami and colleagues [53] found a FFQ was useful for 544 assessing habitual dietary intake of college athletes for vitamin C, calcium, vegetables, fruits, and 545 milk and dairy products. While Fogelholm and Lahti-Kosti [51] found close agreement of EI between 546 the FR and FUQs, with most food group correlations above r=0.24, except for 'vegetable oils', 'other 547 fats', 'standard milk', 'pork', 'beef' and 'poultry'. 548 More recently, nutrition epidemiology has progressed from examining nutrients per se to 549 exploring the relationship of dietary patterns or diet quality on health outcomes [79,80]. However to 550 date, there has been limited evaluation of diet quality or dietary patterns in athletes [8,81-83]. Dietary

patterns, including the specific timing of intake over the day, have been shown to enhance health,

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training and performance outcomes [1,83] and should be addressed in future studies. Acknowledging there are challenges with assessing intake accurately, it may prove beneficial in understanding dietary patterns or diet quality of athletes to help identify athletes who may benefit from dietary input, and provide a platform to educate individuals about dietary choices for optimal health and sports performance.

# 4.3 Qualitative Assessment of Methodological Quality

Overall the studies included in this review (*n*=18) were of fair to moderate quality. Poor ratings were generally noted for items which evaluated external validity and internal validity. For example, selection criteria (i.e. inclusion and exclusion criteria) and representativeness of the participants was usually not adequately described. Ten studies included an adequate description of participant characteristics (i.e. item 3) [21,22,40-42,46,47,49,50,54]; while only six studies [21,29,42,50,51,54] clearly identified the source and how participants were recruited (i.e. item 10) and/or stated the proportion who were asked agreed to participate in the study (i.e. item 11). Potential confounding factors such as supplement use, maintenance of body mass, or maintenance of usual physical activity levels were often not discussed. The description of and adjustment for confounding factors (i.e. items 4 and 19, respectively) were either unable to be determined or deemed not relevant for most of the DLW papers [21,22,40,41,44-48] due to the nature of methodology involving DLW studies. Finally, attempts made to blind investigators to the main outcomes of the intervention were poorly rated for all but two studies [49,54]. Despite known challenges with reporting dietary intake accurately using FR; monitoring of and compliance with self-reported intake data was not always considered in the included studies. Twelve studies indicated that the expertise of qualified dietitians were involved in the collection and/or or analysis of dietary data [22,41-44,46-51,54].

# 5. Limitations, Strengths and Future Directions

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One of the key strengths of the current review includes the extensive, systematic search and evaluation of the literature which has compared two or more dietary assessment methods, including measuring intake against dietary biomarkers or reference measures of energy expenditure in athletes. The results indicate there are limited robust studies evaluating the validity of dietary assessment methods in athletes. The caliber of participants could have influenced study findings and relevance for an elite athlete population. For example, a number of studies involved participants from university sporting clubs or teams [22,29,41,43,49,53], while two studies did not provide sufficient detail about the recruitment source or participant athletic caliber [45,52]. The relatively small sample size in all but two of the eighteen papers could influence the power required to detect meaningful differences. For validation studies, Serra-Majem and colleagues [38] suggest including a sample size greater than 100 participants, or more than 50 participants when using a biomarker. However, it is not uncommon for sports science research to include a relatively small population sample which could make it difficult to detect significant change [84]. There may also be challenges in recruiting a sample that is powerful enough and representative of high level athletes due to reticence of athletes to commit to research studies perceived as time consuming and detracting from training or competition commitments. A further consideration is that the isotope deuterium used in DLW studies is expensive which could influence sample size due to cost and research feasibility. It has been suggested that combining dietary assessment methods may enhance the accuracy of assessing dietary intake [85]. For example, Rumbold and colleagues [86] found a combination of FR and 24-h DRs were effective when quantifying EI in adolescent netball players (i.e. 4.2% difference between measures indicating a slight bias towards over-reporting). In a similar study protocol, Briggs

and colleagues [87] compared the accuracy of a combined dietary data collection method (i.e. weighed FR and 24-h DR) to observed EI of adolescent male soccer players. The results showed systematic under-reporting of intake compared to observed intake (p<0.01), but the bias was small (i.e. degree of random error between dietary methods 3.1%). These publications suggest a combined dietary data collection method may provide an effective technique when quantifying energy intake in athletes.

Finally, emerging image-assisted technical innovations such as wearable cameras, handheld devices and mobile telephone technology have been shown to improve participant compliance by reducing the burden of recording and enhance the accuracy of data recorded [30-33]. Results from a systematic review by Gemming and colleagues [32] indicate that images enhance self-report by revealing unreported foods and identify misreporting errors not captured by traditional assessment methods alone. The use of an image-based FR deployed via a mobile application has recently been shown to be a positive tool for dietary monitoring and potential influence on dietary habits and behaviors [88].

#### 6. Conclusions

Adequate dietary intake is crucial for the maintenance of health and optimizing performance outcomes of athletes. However, there are unique challenges in assessing intake, including the day-to-day variation of expenditure and wide use of commercial sports foods and supplements, which can challenge the accuracy of dietary assessment methods used in athletes. Results from this review suggest self-reported food records may not be a suitable assessment method for quantifying energy expenditure, particularly for weight conscious athletes or athletes with high energy demands. Existing literature demonstrates substantial variability between dietary assessment methods, with

618 under and misreporting of intake frequently observed. As such, there is a clear need for careful 619 validation of dietary assessment methods with emerging technical innovations likely to show 620 promise as they may assist with portion quantification, reduce the burden of collection and problem 621 with missing foods, among athlete populations. 622 Supplementary Materials: The following are available online at www.mdpi.com/link, 623 Supplementary material 1: Search strategy and inclusion criteria; Supplementary material 2: Quality 624 ratings guide. 625 Acknowledgments: The authors would like to acknowledge funding support by the Queensland 626 Academy of Sport's Sport Performance Innovation and Knowledge Excellence unit, Brisbane, 627 Australia, which assisted in the preparation and submission of this manuscript. 628 The authors would also like to acknowledge the technical expertise of Ms. Elaine Tam who assisted 629 with the search of relevant databases and retrieval of articles from The University of Sydney Health 630 Sciences library. 631 Author Contributions: The primary author (LC) and co-authors (HO, VF, JG, GS, KB) were 632 responsible for the conception, design and planning of the manuscript; analysis and interpretation of 633 the data; and drafting and subsequent revisions of the manuscript. Contributing authors were also 634 involved in decisions involving final article inclusion (KB, GS, LC, HO), extraction of data (KB, LC), 635 quality ratings (JG, KB, LC, HO, VF), and revision of the manuscript and final approval for 636 publication (KB, JG, GS, VF, HO, LC). 637 Conflicts of Interest: The authors declare no conflict of interest.

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