



## Conference on ‘Nutrient–nutrient interaction’ Symposium 1: Competition and bioavailability of dietary components

### Approaches used to estimate bioavailability when deriving dietary reference values for iron and zinc in adults

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This review aims to describe approaches used to estimate bioavailability when deriving dietary reference values (DRV) for iron and zinc using the factorial approach. Various values have been applied by different expert bodies to convert absorbed iron or zinc into dietary intakes, and these are summarised in this review. The European Food Safety Authority (EFSA) derived zinc requirements from a trivariate saturation response model describing the relationship between zinc absorption and dietary zinc and phytate. The average requirement for men and women was determined as the intercept of the total absorbed zinc needed to meet physiological requirements, calculated according to body weight, with phytate intake levels of 300, 600, 900 and 1200 mg/d, which are representative of mean/median intakes observed in European populations. For iron, the method employed by EFSA was to use whole body iron losses, determined from radioisotope dilution studies, to calculate the quantity of absorbed iron required to maintain null balance. Absorption from the diet was estimated from a probability model based on measures of iron intake and status and physiological requirements for absorbed iron. Average dietary requirements were derived for men and pre- and post-menopausal women. Taking into consideration the complexity of deriving DRV for iron and zinc, mainly due to the limited knowledge on dietary bioavailability, it appears that EFSA has made maximum use of the most relevant up-to-date data to develop novel and transparent DRV for these nutrients.

#### Bioavailability: Dietary reference values: Iron: Zinc

##### Dietary reference values: concepts and terminology

Reference values for population nutrient intakes are the basis for making dietary recommendations that are consistent with good health. They are used for assessing and planning diets for individuals and groups and developing nutrition policies including food fortification. In the UK and Europe, they are collectively known as dietary reference values (DRV) and in the US/Canada, they are called dietary reference intakes. There are four main categories of values, for which different terms are used, depending

on the body setting the DRV, and each category has a different application:

- (a) Average requirement (AR)<sup>(1,2)</sup>, also called estimated average requirement<sup>(3,4)</sup>. This is the daily intake of a nutrient that meets the needs of 50 % of a healthy population, given a normal distribution of the requirement. It is the most important category for assessing the adequacy of nutrient intakes of a population group and for planning intervention strategies, such as food fortification.
- (b) Population reference intake<sup>(1)</sup>, also called the reference nutrient intake<sup>(2)</sup>, RDA<sup>(3)</sup> and recommended

**Abbreviations:** AI, adequate intake; AR, average requirement; DRV, dietary reference values; EFSA, The European Food Safety Agency.  
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nutrient intake<sup>(4)</sup>. This is the daily intake of a nutrient that meets the needs of almost all healthy individuals, namely 97.5% of the population, and is derived from the AR plus two SD. It can be used to plan the diets of individuals but not population groups. The precise SD is usually not known, and it is generally assumed to be 10–15%.

- (c) Adequate intake (AI)<sup>(1,3)</sup> or safe intake<sup>(2)</sup>. These values are derived as a last resort when data for estimating an AR are not available. The AI is the observed or experimentally determined average (median) intake in a group of apparently healthy people who are assumed to have an adequate status of the nutrient. The AI may well be higher than the actual requirements. When no population reference intake is available, the AI can be used as a guide to plan individuals' diet. The use of AI for assessing the diets of individuals or population groups is limited.
- (d) Upper level or tolerable upper intake level or upper tolerable nutrient intake<sup>(1,3–5)</sup>. These are maximum intake values for populations that if consumed chronically over time are judged unlikely to result in adverse effects; they do not apply to acute effects from high doses, e.g. supplements.

Attempts to harmonise DRV globally were initiated over 10 years ago<sup>(6)</sup> and a new umbrella term, nutrient reference value was proposed. In 2017, the National Academy of Science, Engineering and Medicine organised a workshop with WHO/FAO to discuss harmonisation of nutrient reference values<sup>(7)</sup>. An ad-hoc NAS committee prepared a report on the harmonisation process, which includes case studies on iron and zinc<sup>(8)</sup>.

The derivation of DRV requires quantitative information on the dietary requirements to prevent nutrient deficiency and maintain adequate body store or status. Where appropriate, prevention of chronic diet-related disease may also be taken into account, in which case intakes may be higher than those required to prevent deficiency. The most appropriate indicator for deriving the average nutrient requirement, which is ideally a biomarker for which a dose–response relationship has been shown, must be selected. When a nutrient does not have a useful biomarker, other approaches must be used to establish DRV, such as factorial modelling. This involves estimating physiological requirements for the absorbed nutrient and then converting this into dietary intakes using a conversion factor that takes into account bioavailability from the diet.

The definition of nutrient bioavailability is the percentage (or fraction) of total intake that is absorbed and utilised for normal body functions. Iron and zinc are two minerals that have varying (and sometimes low) bioavailability, depending on a number of dietary and host-related factors. Although isotopic labels can be used to directly measure iron bioavailability (namely utilisation) as the percentage of intake that is incorporated into Hb, this technique cannot be used to determine overall bioavailability from whole diets. Furthermore, there are no equivalent biomarkers of utilisation for zinc. Therefore, absorption is used as a surrogate measure of

bioavailability for both iron and zinc and the two terms will be used interchangeably in this review.

### Approaches used to derive dietary reference values for zinc

#### *Estimating physiological requirements*

In the absence of a suitable biomarker, dietary requirements for zinc have, in the past, been derived using the factorial approach. The European Micronutrient Recommendations Aligned (EURRECA) network undertook a series of systematic reviews, the aim of which was to develop an intake–status–health relationship model to inform the setting of DRV for zinc<sup>(9)</sup>. However, they were unable to recommend an alternative to the factorial approach as the potentially useful indicators of zinc status (plasma/serum zinc, hair zinc and urinary zinc excretion)<sup>(10)</sup> were not specific enough to characterise an intake–response relationship.

The factorial approach entails the estimation of the quantity of absorbed zinc required to replace endogenous losses of zinc. The majority of zinc lost from the body is intestinal (generally referred to as endogenous faecal zinc) and has been measured in conventional balance studies and with the use of radio- and stable isotope labels. Endogenous faecal zinc losses reflect the quantity of zinc absorbed, which is dependent on zinc intake. Non-intestinal losses include urine, dermal, menstrual (women) and semen (men). These are assumed to be constant over a wide range of zinc intake<sup>(3)</sup>, with estimates ranging from 0.30 to 0.63 mg/d in men and 0.30 to 0.44 mg/d in women<sup>(11)</sup>. Semen losses were estimated to be 0.10 mg/d<sup>(1,3)</sup> and menstrual losses about 0.01 mg/d<sup>(1)</sup>.

Estimates of total endogenous zinc losses reported by different bodies range from 1.40<sup>(4)</sup> to 3.84 mg/d<sup>(3)</sup> in adult men and 1.00<sup>(4)</sup> to 3.30 mg/d<sup>(3)</sup> in adult women. The low estimates from WHO were derived from studies in which the diets were very low in zinc. The US Institute of Medicine of the National Academy of Sciences used a linear regression approach to examine the relationship between endogenous faecal zinc and the quantity of zinc absorbed, which was then adjusted for non-intestinal losses of zinc in order to estimate total endogenous zinc losses<sup>(3)</sup>. The European Food Safety Authority (EFSA) used multiple regression analysis and found that body size was the primary predictor of the quantity of zinc absorbed (when zinc balance is null), with no sex effect. An equation relating zinc physiological requirement to body weight was derived and zinc physiological requirements were estimated to be 3.20 mg/d in men (mean body weight 68.1 kg, based on measurements made in 16 500 European men aged 18–79 years) and 2.90 mg/d in women (mean body weight 58.5 kg, based on measurements made in 19 969 European women aged 18–79 years)<sup>(12)</sup>.

#### *Converting physiological requirements into dietary intakes*

The AR is the quantity of dietary zinc that will supply the physiological needs (i.e. replace endogenous losses) of

**Table 1.** Absorption values used for zinc

Report	Absorption (with supporting evidence if provided)
DH 1991 <sup>(2)</sup>	30 %
NL 1992 <sup>(13)</sup>	25 %
IOM 2001 <sup>(25)</sup>	41 % in men 48 % in women Asymptotic regression of absorbed zinc on ingested zinc
FAO/WHO 2004 <sup>(14)</sup>	High bioavailability 50 % Moderate bioavailability 30 % Low bioavailability 15 % Algorithms for high, moderate and low bioavailability diets, and zinc content of the diet
NCM 2012 <sup>(15)</sup>	40 %
EFSA 2014 <sup>(12)</sup>	Phytate level (mg/d)    Men (%)    Women (%) 300                            42            46 600                            33            38 900                            28            32 1200                           24            28 Trivariate saturation response model (zinc absorption, zinc intake and phytate level)
ANSES 2016 <sup>(19)</sup>	Based on EFSA 2014, assuming phytate levels of 300–900 mg/day for the French diet
D–A–CH 2016 <sup>(20)</sup>	30 % from a mixed diet <sup>(38,39)</sup>

ANSES, French Agency for Food, Environmental and Occupational Health and Safety; D–A–CH, Deutschland–Austria–Confoederatio Helvetica; DH, UK Department of Health; EFSA, European Food Safety Authority; FAO, Food and Agriculture Organization; IOM, US Institute of Medicine of the National Academy of Sciences; NCM, Nordic Council of Ministers; NL, Health Council of the Netherlands.

50 % of the population. In order to convert requirements for absorbed zinc into dietary intakes, a correction for absorption has to be made. The efficiency of absorption depends on dietary composition, including zinc content and modifiers of absorption, and physiological needs. In particular, phytate (myo-inositol hexaphosphate), a phosphorus storage compound in plants, is a key determinant of zinc absorption. Values for dietary zinc absorption used by several bodies are summarised in Table 1.

The UK Department of Health assumed a value of 30 % for zinc absorption<sup>(2)</sup>, and the Health Council of the Netherlands used 25 %<sup>(13)</sup>. The US Institute of Medicine of the National Academy of Sciences<sup>(3)</sup> values were 41 % for men and 48 % for women, derived from a regression analysis using data from multiple studies on zinc absorption. However, many of the diets used in these studies were semi-purified and low in phytate, and it is likely that the inhibitory effect of phytate was underestimated.

FAO/WHO<sup>(14)</sup> estimated that 50 % of dietary zinc would be absorbed from highly refined diets. These were described as low in cereal fibre, low in phytic acid content, and with phytate–zinc molar ratio <5; adequate protein content principally from non-vegetable sources, such as meats and fish (high bioavailability). Absorption from moderate bioavailability diets was estimated to be 30 %; these diets contain animal or fish protein, and this category includes lacto-ovo,

ovo-vegetarian, or vegan diets not based primarily on unrefined cereal grains or high-extraction-rate flours. The phytate–zinc molar ratio of the diet should be within the range 5–15, or not exceeding 10 if more than 50 % of the energy intake is accounted for by unfermented, unrefined cereal grains and flours and the diet is fortified with inorganic calcium salts (>1 g Ca<sup>2+</sup>/d). Absorption from low bioavailability diets was estimated to be 15 %. Such diets were described as high in unrefined, unfermented, and ungerminated cereal grain, especially when fortified with inorganic calcium salts and when the intake of animal protein is negligible. Low bioavailability was associated with diets where the phytate:zinc molar ratio of total diet exceeds 15, and high-phytate, soya-protein products constitute the primary protein source.

The Nordic Council of Ministers assumed an absorption of 40 % from a mixed animal and vegetable protein diet; this was the same as their previous report because they considered there were no new scientific data to justify changes<sup>(15)</sup>.

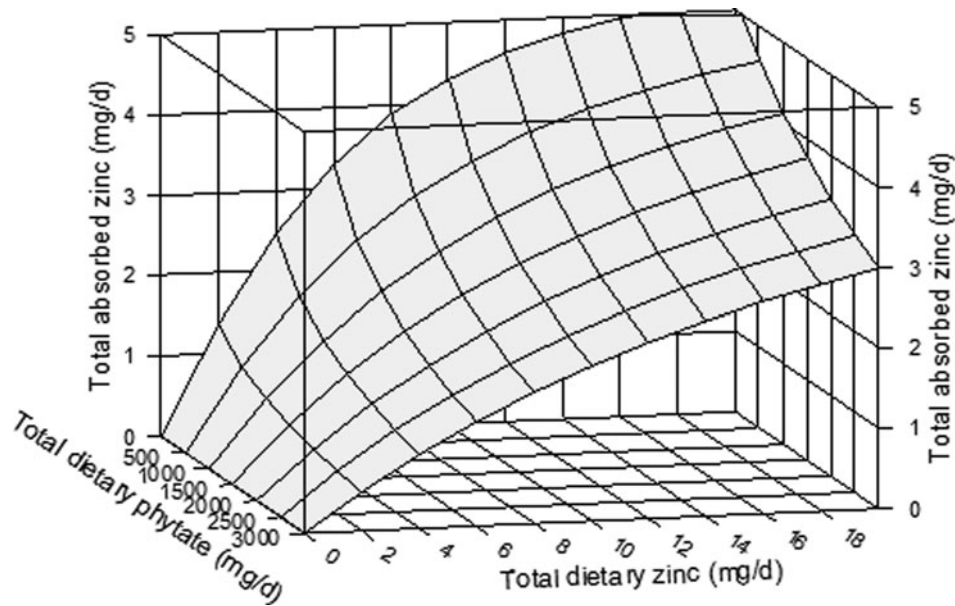
There are several models for estimating zinc absorption at different levels of phytate intake. The International Zinc Nutrition Consultative Group model<sup>(16)</sup> predicts that at phytate:zinc molar ratios of between 4 and 18, zinc absorption is 26 and 34 % in men and women, respectively, whereas, at ratios >18, absorption falls to 18 and 25 % in men and women, respectively. EFSA<sup>(12)</sup> calculated values for dietary zinc absorption at different levels of phytate (and zinc) intake using a refined trivariate model<sup>(17)</sup> (Fig. 1). This model was based on the original one developed by Miller *et al.*<sup>(18)</sup> and involved a careful and critical selection of individual values for zinc absorption from meals that were considered more representative of Western diets. The modifying effects of calcium, protein and iron were found to be insignificant and therefore were discounted in the final model<sup>(17)</sup>. Using this model, EFSA derived DRV for levels of phytate of 300, 600, 900 and 1200 mg/d (Fig. 1) which covers the usual mean/median intakes in different European countries although higher values may be found in certain countries.

ANSES<sup>(19)</sup> endorsed the approach proposed by EFSA, whereas Deutschland–Austria–Confoederatio Helvetica cited references to support their selected value of 30 % from a mixed diet<sup>(20)</sup>.

### Approaches used to derive dietary reference values for iron

#### *Estimating physiological requirements*

There are no data on the relationship between iron intake and biomarkers of the physiological requirement that can be used to derive DRV, therefore the factorial approach has been used. Obligatory (sometimes referred to as basal or endogenous) losses of iron (e.g. dermal, epithelial, intestinal and urinary) were measured in three small groups of men in the 1960s using radioisotopes, with reported mean values ranging from 0.90 to 1.02 mg/d<sup>(21)</sup>. Based on these data, basal iron losses of 0.014 mg/kg



**Fig. 1.** Refined trivariate model of dietary zinc absorption at different levels of phytate and zinc to derive dietary reference values for zinc at phytate levels of 300, 600, 900 and 1200 mg/d.

body weight daily have been used by several bodies<sup>(3,14,15)</sup>, to which estimates of menstrual iron losses had to be added in order to estimate the requirement for women of childbearing age. Using the same technique, a more recent study measured basal iron losses of 1.07 mg/d in men and 1.69–1.89 mg/d for premenopausal women<sup>(22)</sup>. The results are summarised in Table 2. EFSA developed a regression model equation from the individual data on iron turnover and daily losses from the Hunt *et al.*<sup>(22)</sup> study in order to derive distributions for iron losses in men and women, from which percentiles could be estimated as the basis for determining AR and population reference intake values<sup>(23)</sup>. For men, the 50th centile was 0.95 mg/d and for pre-menopausal women, it was 1.34 mg/d.

#### *Converting physiological requirements into dietary intakes*

The efficiency of iron absorption is determined by body iron requirements, which is related to body iron stores<sup>(24)</sup> and the properties of the diet, i.e. iron content and the presence and quantity of enhancers and inhibitors of absorption<sup>(25,26)</sup>. When deriving DRV, physiological iron requirements need to be converted into dietary intakes by applying a bioavailability factor.

DRV setting bodies have used different values for the bioavailability factor, but they are not based on primary data for iron absorption from the whole diet over an extended period of time because this is very difficult to measure. A EURRECA systematic review of iron absorption from whole diets<sup>(27)</sup> identified nineteen pertinent studies from the USA, Europe and Mexico. Large variations in mean non-haem iron absorption, ranging from 0.7 to 22.9%, were found between the studies, which were related to the iron status of the individuals.

Various algorithms have been developed to estimate iron absorption from the whole diet by taking into account the quantity of enhancers and inhibitors of iron absorption. The early algorithms used data from single meal studies in which iron absorption was measured by labelling the dietary iron with radio or stable isotopes<sup>(28)</sup>, but single meal studies tend to exaggerate the effect of enhancers and inhibitors<sup>(29)</sup>. More recently, a diet-based algorithm was developed using iron absorption data from whole diets or several meals<sup>(30)</sup>. This algorithm was used to estimate absorption from the US diet, taking into account the mean intake of inhibitors (phytate and polyphenols) and enhancers (ascorbic acid), and the proportion of haem (10%) and non-haem (90%) iron<sup>(31)</sup> in the diet. Total dietary iron absorption was calculated to be 15.5%.

Various values for dietary iron absorption (bioavailability) that are based on results from isotope absorption studies are used by different expert bodies charged with setting DRV (Table 3). Values range between 10 and 18% for Western-type diets<sup>(2,3,13–15,20,23,32)</sup>.

The US Institute of Medicine of the National Academy of Sciences (2001)<sup>(3)</sup> considered that the maximum bioavailability of iron was 18% in (non-pregnant, non-lactating) adults. This value was based on the assumption that 10% of dietary iron intake was haem iron<sup>(33)</sup>, with haem iron absorption being 25%<sup>(34)</sup>, and non-haem iron absorption being 16.8% in individuals with a serum ferritin concentration of 15 µg/l<sup>(29)</sup>. With regard to iron losses, special consideration was given to the use of oral contraceptives and hormone replacement therapy, and also to vegetarianism, intestinal parasitic infection, blood donation, and increased iron losses in exercise and intense endurance training. The decrease in menstrual blood losses in women using oral contraceptives<sup>(35)</sup> was taken into consideration, which gave an

**Table 2.** Total obligatory iron losses measured using radioisotopes in adult men and women<sup>(21,22)</sup>

Country	Participant characteristics	Daily iron losses (mg/d) ( $\pm$ sd)	Reference
USA	Men 21–61 years ( <i>n</i> 12), mean body weight 78.6 kg	Mean = 0.95 ( $\pm$ 0.30)	Green <i>et al.</i> <sup>(21)</sup>
Venezuela	Men 22–55 years ( <i>n</i> 12), mean body weight 67.6 kg	Mean = 0.90 ( $\pm$ 0.31)	
South Africa	Men 21–65 years ( <i>n</i> 17), mean body weight 62.3 kg	Mean = 1.02 ( $\pm$ 0.22)	
USA	Men 30–58 years ( <i>n</i> 29), mean body weight 92 kg	Mean = 1.07 ( $\pm$ 0.47);	Hunt <i>et al.</i> <sup>(22)</sup>
USA	Women 32–47 years ( <i>n</i> 15), mean body weight 73 kg*	Mean = 1.89 [1.07, 3.33] <sup>†</sup>	
USA	Women 32–47 years ( <i>n</i> 19), mean body weight 74 kg	Mean = 1.69 [0.98–2.92] <sup>†</sup>	

\* Pre-menopausal women, no hormonal contraceptives users.

<sup>†</sup> Geometric mean with  $-sd$  and  $+sd$  in brackets.

**Table 3.** Bioavailability factors used to convert total absorbed iron into the dietary intake

Report	Bioavailability factor (with supporting evidence if provided)
DH 1991 <sup>(2)</sup>	15 %
NL 1992 <sup>(13)</sup>	12 %
IOM 2001 <sup>(3)</sup>	18 % – mixed diet (assuming 10 % dietary iron is haem, with the absorption of 25, and 90 % is non-haem with the absorption of 10 %, overall iron absorption = 17.6 %)
	10 % – vegetarian diet
FAO/WHO 2004 <sup>(14)</sup>	5 % – developing countries (high phytate, high tannin, negligible meat/fish, low ascorbic acid intake)
	10 % – developing countries
	12 % – Western diets
	15 % – Western diet (high meat intake)
NHMRC/NZ MoH 2006 <sup>(32)</sup>	18 % – mixed western diet including animal food
	10 % – vegetarian diet
	Based on IOM 2001 <sup>(3)</sup>
NCM 2012 <sup>(15)</sup>	15 %
EFSA 2015 <sup>(23)</sup>	18 % in pre-menopausal women
	16 % in men
	–target serum ferritin 30 $\mu$ g/l <sup>(36)</sup>
D–A–CH 2016 <sup>(20)</sup>	10–15 %

D–A–CH, Deutschland–Austria–Confoederatio Helvetica; DH, UK Department of Health; EFSA, European Food Safety Authority; FAO, Food and Agriculture Organization; IOM, US Institute of Medicine of the National Academy of Sciences; NCM, Nordic Council of Ministers; NHMRC, Australian National Health and Medical Research Council; NL, Health Council of the Netherlands; NZ MoH, New Zealand Ministry of Health.

estimated reduction in losses of about 60 %. The iron bioavailability of a vegetarian diet was estimated to be about 10 % instead of 18 % for a mixed Western diet.

FAO/WHO<sup>(14)</sup> based their recommended nutrient intakes for women aged  $\geq 19$  years on the 95th percentile of the total requirements for absorbed iron (the average physiological requirement was assumed to be 1.46 mg/day). They proposed four different bioavailability figures: 15 and 12 % for Western-type diets, depending mainly on meat intake, and 10 and 5 % for developing countries.

EFSA employed values for dietary iron absorption that were derived using a novel approach developed by Dainty *et al.*<sup>(36)</sup>. Total iron (haem and non-haem) absorption was predicted from a probability model, based on measures of iron intake and status in a

representative group of men and women from the UK National Diet and Nutrition Survey. The model can provide estimates of total iron absorption from a mixed Western-style diet at any level of iron status. The EFSA panel selected a target value of 30  $\mu$ g/l for serum ferritin concentration for men and premenopausal women, and at this level, the predicted iron absorption values were 16 and 18 %, respectively<sup>(23)</sup>. Additional data from a nationally representative survey in Ireland and data collected in older people in the UK have subsequently been included in the model<sup>(37)</sup>, and an interactive tool developed for estimating total dietary iron absorption in adult populations with a selected target serum ferritin concentration<sup>(37)</sup>.

## Conclusions

Accounting for the bioavailability of iron and zinc remains one of the most challenging aspects of setting DRV. It has been established that the dietary levels of phytate and zinc are the key determinants of zinc absorption in adults, but further studies are required to examine the relationship in other population groups, especially infants, children and pregnant women. Similarly, for iron, the model used to predict dietary absorption was derived from adult data, and further work is required to develop models for other population groups. The existing model for adults needs to be adapted for lower and middle-income countries, in which intakes of iron absorption inhibitors may be higher than Western diets and haem iron intakes may be lower. Good quality representative data for iron intake and status (serum ferritin, taking due account of the presence of infection/inflammation) could be used to evaluate the validity of the existing model. Alternative interactive tools for predicting dietary iron absorption could be based on the mathematical relationship between iron intake, iron status and iron requirements (all of which can be estimated with a reasonable degree of accuracy) since iron status is determined from the difference between physiological requirements and the quantity of dietary iron that is absorbed.

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### Conflict of Interest

At the time of writing this review, S. F-T. was an appointed expert for the European Food Safety Authority Panel on Dietetic Products, Nutrition and Allergies (2009–2018) and the Working Group on DRV Minerals (2009–2015, vice-chair 2015–2018). She was also a member of an ad hoc Committee of the US National Academies of Sciences, Engineering and Medicine on the application of methodological approaches to global harmonisation of nutrient intake recommendations for young children and women of reproductive age (2017–2018). A. de S. is employed with EFSA in the Nutrition Unit that provides scientific and administrative support to the Panel on Dietetics Products, Nutrition and Allergies in the area of DRV for minerals. However, the present article is published under the sole responsibility of the authors and may not be considered as an EFSA scientific output. The positions and opinions presented in this article are those of the authors alone and do not necessarily represent the views or scientific work of EFSA.

### Authorship

The authors are jointly responsible for all aspects of preparation of this paper.

### References

1. EFSA Panel on Dietetic Products, Nutrition and Allergies (2010) Scientific opinion on principles for deriving and applying dietary reference values. *EFSA J* **8**, 1458.
2. Department of Health (1991) Dietary reference values for food and energy and nutrients for the United Kingdom. Report of the panel on dietary reference values of the committee on medical aspects of food policy. *Rep Health Soc Subj (Lond)* **41**, 1–210.
3. Institute of Medicine (2001) *Dietary Reference Intakes for Vitamin A, Vitamin K, Arsenic, Boron, Chromium, Copper, Iodine, Iron, Manganese, Molybdenum, Nickel, Silicon, Vanadium, and Zinc*. Washington, DC: National Academy Press.
4. Joint FAO/WHO Expert Consultation on Human Vitamin and Mineral Requirements (1998) Vitamin and mineral requirements in human nutrition: report of a joint FAO/WHO expert consultation, Bangkok, Thailand, 21–30 September 1998. Geneva: WHO.
5. EFSA Scientific Committee on Food and NDA Panel (2006) Tolerable Upper Intake Levels for Vitamins and Minerals [http://www.efsa.eu.int/science/nda/nda\\_opinions/catindex\\_en.htm](http://www.efsa.eu.int/science/nda/nda_opinions/catindex_en.htm) (accessed March 2018).
6. King JC & Garza C (2007) Harmonization of nutrient intake values. *Food and Nutrition Bulletin* **S3-S12**, 28 no. 1 (supplement 1).
7. National Academies of Sciences, Engineering, and Medicine (2018) *Global Harmonization of Methodological Approaches to Nutrient Intake. Recommendations: Proceedings of a Workshop*. Washington, DC: The National Academies Press.
8. National Academies of Sciences, Engineering, and Medicine (2018) *Harmonization of Approaches to Nutrient*

*Reference Values: Applications to Children and Women of Reproductive Age*. Washington, DC: The National Academies Press.

9. Lowe NM, Dykes FC, Skinner AL *et al.* (2013) EURRECA – Estimating zinc requirements for deriving dietary reference values. *Crit Rev Food Sci Nutr* **53**, 1110–1123.
10. Lowe NM, Medina MW, Stammers AL *et al.* (2012) The relationship between zinc intake and serum/plasma zinc concentration in adults: a systematic review and dose-response meta-analysis by the EURRECA Network. *Br J Nutr* **108**, 1962–1971.
11. Gibson RS, King JC & Lowe N. (2016) A review of dietary zinc recommendations. *Food Nutr Bull* **30**, S108–S143.
12. EFSA Panel on Dietetic Products, Nutrition and Allergies (2014) Scientific opinion on dietary reference values for zinc. *EFSA J* **12**, 3844.
13. Netherlands Food and Nutrition Council (1992) *Recommended Dietary Allowances 1989 in the Netherlands*. 115 pp. The Hague: Netherlands Food and Nutrition Council.
14. WHO and FAO (2004) *Vitamin and Mineral Requirements in Human Nutrition*. Geneva WHO.
15. Nordic Council of Ministers (2014) *Nordic Nutrition Recommendations 2012. Integrating nutrition and physical activity*. 627 pp. Copenhagen: Nordic Council of Ministers.
16. Brown KH, Rivera JA, Bhutta Z *et al.* (2004) International Zinc Nutrition Consultative Group technical document #1. Assessment of the risk of zinc deficiency in populations and options for its control. *Food Nutr Bull* **25**(1 Suppl 2), S99–S203.
17. Miller LV, Krebs NF & Hambidge KM (2013) Mathematical model of zinc absorption: effects of dietary calcium, protein and iron on zinc absorption. *Br J Nutr* **109**, 695–700.
18. Miller LV, Krebs NF & Hambidge KM (2007) A mathematical model of zinc absorption in humans as a function of dietary zinc and phytate. *J Nutr* **137**, 135–141.
19. ANSES (Agence nationale de sécurité sanitaire de l'alimentation, de l'environnement et du travail) (2016) Actualisation des repères du PNNS : élaboration des références nutritionnelles. Avis de l'ANSES. Rapports d'expertise collective. Available at: <http://alimentation-sante.org/wp-content/uploads/2017/01/NUT2012SA0103Ra-1.pdf>
20. Deutsche Gesellschaft für Ernährung, Österreichische Gesellschaft für Ernährung, Schweizerische Gesellschaft für Ernährung (2016) Referenzwerte für die Nährstoffzufuhr. Neuer Umschau Buchverlag, Bonn. Available at: <https://www.dge.de/wissenschaft/referenzwerte/>
21. Green R, Charlton R, Seftel H *et al.* (1968) Body iron excretion in man: a collaborative study. *Am J Med* **45**, 336–353.
22. Hunt JT, Zito CA & Johnson LK (2009) Body iron excretion by healthy men and women. *Am J Clin Nutr* **89**, 1792–1798.
23. EFSA Panel on Dietetic Products, Nutrition and Allergies (2015) Scientific opinion on dietary reference values for iron. *EFSA Journal* **13**, 4254.
24. Hallberg L, Hultén L & Gramatkovski E (1997) Iron absorption from the whole diet in men: how effective is the regulation of iron absorption? *Am J Clin Nutr*, **66**, 347–356.
25. Hurrell R & Egli I (2010) Iron bioavailability and dietary reference values. *Am J Clin Nutr* **91**, 1461S–1467S.
26. Lynch S, Pfeiffer CM, Georgieff MK *et al.* (2018) Biomarkers of nutrition for development (BOND) – iron review. *J Nutr* **148**, 1001S–1067S.



27. Collings R, Harvey LJ, Hooper L *et al.* (2013) The absorption of iron from whole diets: a systematic review. *Am J Clin Nutr* **98**, 65–81.
28. Hallberg L & Hulthén L (2000) Prediction of dietary iron absorption: an algorithm for calculating absorption and bioavailability of dietary iron. *Am J Clin Nutr* **71**, 1147–1160.
29. Cook JD, Dassenko SA & Lynch SR (1991) Assessment of the role of nonhaem-iron availability in iron balance. *Am J Clin Nutr* **54**, 717–722.
30. Armah SM, Carriquiry A, Sullivan D *et al.* (2013) A complete diet-based algorithm for predicting nonhaem iron absorption in adults. *J Nutr* **143**, 1136–1140.
31. Armah SM, Boy E, Chen D *et al.* (2015) Regular consumption of a high-phytate diet reduces the inhibitory effect of phytate on nonhaem-iron absorption in women with suboptimal iron stores. *J Nutr* **145**, 1735–1739.
32. Australian National Health and Medical Research Council/New Zealand Ministry of Health (2006) Nutrient Reference Values for Australia and New Zealand Including Recommended Dietary Intakes. Available at: <https://www.nhmrc.gov.au/guidelines-publications/n35-n36-n37>
33. Raper NR, Rosenthal JC & Woteki CE (1984) Estimates of available iron in diets of individuals 1 year old and older in the nationwide food consumption survey. *J Am Diet Assoc* **84**, 783–787.
34. Hallberg L, Rossander-Hultén L (1991) Iron requirements in menstruating women. *Am J Clin Nutr* **54**, 1047–1058.
35. Nilsson L & Solvell L (1967) Clinical studies on oral contraceptives – a randomized, doubleblind, crossover study of 4 different preparations (anovlar mite, lyndiol mite, ovulen, and volidan). *Acta Obstet Gynecol Scand* **46**(Suppl 8), 1–31.
36. Dainty JR, Berry R, Lynch SR *et al.* (2014) Estimation of dietary iron bioavailability from food iron intake and iron status. *PLoS One* **9**(10), e111824.
37. Fairweather-Tait SJ, Jennings A, Harvey LJ *et al.* (2017) Modeling tool for calculating dietary iron bioavailability in iron-sufficient adults. *Am J Clin Nutr* **105**, 1408–1414.
38. Milne DB, Canfield WK, Mahalko JR *et al.* (1983) Effect of dietary zinc on whole body surface loss of zinc: impact on estimation of zinc retention by balance method. *Am J Clin Nutr* **38**, 181–186.
39. Taylor CM, Bacon JR, Aggett PJ *et al.* (1992) Homeostatic regulation of zinc absorption and endogenous losses in zinc-deprived men. *Am J Clin Nutr* **53**, 755–763. Erratum in: *Am J Clin Nutr* (1992), 56, 462.