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Draft Scientific Opinion on the safety and suitability for use by infants of follow-on formulae with a protein content of at least 1.6 g/100 kcal

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Abstract

Following a request from the European Commission, the EFSA Panel on Dietetic Products, Nutrition and Allergies (NDA) was asked to deliver a scientific opinion on the safety and suitability for use by infants of follow-on formulae (FOF) based on cow's milk intact protein with a protein content of at least 1.6 g/100 kcal (rounded value) that meet otherwise the requirements of relevant EU legislation. If the formula under evaluation is considered to be safe and suitable for use by infants, the NDA Panel is also asked to advise on whether FOF based on goat's milk intact protein, soy protein isolates or protein hydrolysates are also safe and suitable for infants under the same conditions. The Panel concludes that the use of FOF with a protein content of at least 1.6 g/100 kcal from either intact cow's milk protein or intact goat's milk protein otherwise complying with the requirements of relevant EU legislation is safe and suitable for infants living in Europe with access to complementary foods of a sufficient quality. This conclusion does not apply to IF. The Panel also concludes that the safety and suitability of FOF with a protein content of at least 1.6 g/100 kcal manufactured from either protein hydrolysates or soy protein isolates cannot be established with the available data. The same conclusion applies to IF. The NDA Panel endorsed this draft scientific opinion on 14 December 2016 for public consultation, to which stakeholders are encouraged to contribute. The document will be revised and updated according to the comments received, where appropriate.

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Summary

Following a request from the European Commission, the EFSA Panel on Dietetic Products, Nutrition and Allergies (NDA) was asked to deliver a scientific opinion on the safety and suitability for use by infants of follow-on formulae (FOF) based on cow's milk intact protein with a protein content of at least 1.6 g/100 kcal (rounded value) that meet otherwise the requirements of relevant EU legislation. If the formula under evaluation is considered to be safe and suitable for use by infants, the NDA Panel is also asked to advise on whether FOF based on goat's milk intact protein, soy protein isolates or protein hydrolysates are also safe and suitable for infants under the same conditions.

For the scientific assessment, the NDA Panel considered: a) the dietary protein requirements of infants in the second half of the first year of life, b) the protein content of breast milk during the first year of lactation, c) dietary protein intake of infants in Europe from breast milk, formula and complementary food (CF), d) the overall contribution that a FOF with a protein content of 1.6 g/100 kcal could make towards protein requirements in the target population, assuming access to CF of a sufficient quality, following established feeding guidelines in Europe (e.g. from Member States), and e) the application submitted by the food business operator, including two intervention studies in healthy term infants.

The Panel notes that:

- a) Population Reference Intakes (PRIs) of 10 g protein per day for girls and 11 g protein per day for boys aged 6 months and a PRI of 11 g protein per day for girls and 12 g protein per day for boys aged 12 months have been established,
- b) The mean content of true protein in breast milk by the end of the third month of lactation ranges between 1.3 and 1.6 g/100kcal, tends to decrease thereafter to about 1.1-1.4 g/100kcal by the end of the fourth month, and tends to remain fairly stable thereafter,
- c) The P5th and P2.5th of total protein intake in non-breastfed infants aged 6 to 12 months living in Europe was around or above the PRI for protein for that age group in all the studies and surveys available,
- d) The P5th and P2.5th of total protein intake resulting from the consumption of FOF with a protein content of 1.6 g/100kcal would remain at about or above the PRI for protein for infants aged 6 to 12 months who are not breastfed, and
- e) The two randomised, double-blind, controlled intervention studies provided by the applicant showed no differences in growth patterns between healthy term infants who consumed formulae with protein contents of 1.61 g/100 kcal and 1.65 g/100 kcal from three months of age onwards and infants who consumed formulae with protein contents of 2.15 g/100 kcal and 2.70 g/100 kcal, respectively. The control formula used in these studies contained 0.35 g/100 kcal (US study) and 0.90 g/100 kcal (Chile study) more protein than the current minimum requirement for protein content of a FOF (1.8 g/100 kcal).

The Panel also notes that the studies submitted were not specifically designed to meet the regulatory definitions for either IF or FOF laid down in Regulation (EU) No 609/2013¹, and that the information provided in relation to the type and amount of CF was not sufficient to calculate total energy and protein intake, nor the relative contribution of formulae and CF to total energy and protein intake. Therefore, the Panel considers that these studies do not provide, on their own, sufficient information to conclude on the safety and suitability of a FOF with a protein content of 1.6 g protein/100kcal.

The Panel notes, however, that:

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Regulation (EU) No 609/2013 of the European Parliament and of the Council of 12 June 2013 on food intended for infants and young children, food for special medical purposes, and total diet replacement for weight control and repealing Council Directive 92/52/EEC, Commission Directives 96/8/EC, 1999/21/EC, 2006/125/EC and 2006/141/EC, Directive 2009/39/EC of the European Parliament and of the Council and Commission Regulations (EC) No 41/2009 and (EC) No 953/2009, OJ L 181, 29.6.2013, p. 35–56



- a) the protein content of human milk tends to decrease with feeding time to about 1.1-1.4 g/100kcal by the end of the fourth month of lactation, remaining fairly stable thereafter,
 - b) that P5th and P2.5th of protein intake from all sources (breast milk, formula and CF) in European infants between 6 and 12 months of age are at or above the PRI for protein for that age group.
 - c) P5th and P2.5th of protein intake from all sources (formula and CF) in European infants between 6 and 12 months of age who are not breastfed would remain at or above the PRI for protein for that age group by assuming a protein content of 1.6 g/100kcal in all FOF.
 - d) the two human intervention studies provided by the applicant did not show an adverse impact on growth resulting from the use of a formula containing about 1.6 g of protein/100kcal as compared to control formulae containing 2.15 or 2.70 g of protein/100 kcal or the breastfed reference group.

Therefore, the Panel concludes that the use of FOF with a protein content of at least 1.6 g/100 kcal from intact cow's milk protein otherwise complying with the requirements of relevant EU legislation is safe and suitable for infants living in Europe with access to complementary foods of a sufficient quality. This conclusion does not apply to IF.

On the basis of:

- a) a previous evaluation by the Panel on the safety and suitability of goat's milk protein as a source of protein in IF and FOF (EFSA NDA Panel, 2012b), and
- b) the Panel's conclusions regarding the safety and suitability of FOF with a protein content of at least 1.6 g/100 kcal from intact cow's milk protein otherwise complying with the requirements of relevant EU legislation,

the Panel concludes that the use of FOF with a protein content of at least 1.6 g/100 kcal from intact goat's milk protein otherwise complying with the requirements of relevant EU legislation is safe and suitable for infants living in Europe with access to complementary foods of a sufficient quality. This conclusion does not apply to IF.

The Panel considers, however, that the safety and suitability of each FOF (and IF) manufactured from protein hydrolysates have to be established by clinical evaluation in the target population (EFSA NDA Panel, 2014). The Panel also considers that, given the higher minimum protein requirements established for FOF (and IF) manufactured from soy protein isolates (i.e. 2.25 g/100 kcal) and the lack of data available on the use of FOF from soy protein isolates in the target population, additional studies are required to establish the safety and suitability of FOF manufactured from soy protein isolates with a protein content of at least 1.6 g/100 kcal. Therefore, the Panel concludes that the safety and suitability of FOF with a protein content of at least 1.6 g/100 kcal manufactured from either protein hydrolysates or soy protein isolates cannot be established with the available data. The same conclusion applies to IF.



Table of contents

143

144				
145				
146		,		
147	1.		tion	
148	1.1.		und and Terms of Reference as provided by the European Commission	
149	1.1.1.		und	
150	1.1.2.		f Reference	
151	1.2.		tation of the Terms of Reference	
152	2.		d Methodologies	
153	2.1.			
154	2.2.		logies	
155	3.		ent	_
156	3.1.		protein requirements of infants in the second half of the first year of life	
157	3.2.		content of breast milk during the first year of lactation	
158	3.3.		protein intake of infants in Europe	10
159	3.4.		tion that a FOF with a protein content of at least 1.6 g/100 kcal could make	
160			protein requirements in the target population	
161	3.5.		on submitted by the food business operator	
162	3.5.1.		tion of the formulae used in the two human intervention studies	
163	3.5.2.		ntervention studies	20
164	3.6.		son between the human intervention studies provided and European dietary	
165			with respect to mean energy and protein intake from formula and complementary	
166			he target population	
167	4.		ons	23
168	4.1.		afety and suitability for use by infants of FOF with a protein content of at least	
169			0 kcal from intact cow's milk protein otherwise complying with the requirements	
170		of releva	nt EU legislation	23
171	4.2.		afety and suitability for use by infants of FOF with a protein content of at least	
172			0 kcal from goat's milk intact protein, soy protein isolates or protein hydrolysates	
173	_		e complying with the requirements of relevant EU legislation	
174			provided to EFSA	
175				
176				
177		dix A –	Absolute weight, weight gain and weight change in the "US study"	
178		dix B –	Length and head circumference in the "US study"	29
179		dix C –	Absolute weight, weight gain and weight change in the "Chile study"	30
180	Appen	dix D –	Length and head circumference in the "Chile study"	31
181			▼	



1. Introduction

1.1. Background and Terms of Reference as provided by the European 184 **Commission** 185

1.1.1. Background 186

- Commission Directive 2006/141/EC² lays down requirements for infant formulae and follow-on 187
- formulae placed on the market in the EU. Among others, it establishes that follow-on formula 188
- 189 manufactured from cows' milk intact protein shall contain at least 1.8 g protein/100 kcal (Annex II,
- 190 point 2.1).

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- Commission delegated Regulation (EU) 2016/127³ revises the rules of Commission Directive 191
- 2006/141/EC and shall repeal and replace the Directive from 22 February 2020. Annex II, point 2.1 of 192
- the delegated Regulation maintains the minimum protein content of follow-on formula manufactured 193
- from cows' milk intact protein at 1.8 g/100 kcal. 194
- 195 The Commission has received a request from a food business operator for placing on the market a
- 196 follow-on formula based on cow's milk intact protein with a protein content of at least 1.61 g/100 kcal,
- which is below the permitted levels of Directive 2006/141/EC and delegated Regulation (EU) 197
- 2016/127. In order to consider such request, the Commission needs to obtain the advice of the 198
- 199 European Food Safety Authority and has asked the food business operator to send the scientific
- dossier to the Authority for assessment. 200

1.1.2. Terms of Reference 201

- 202 In accordance with Article 29 of Regulation (EC) No 178/2002⁴, the European Commission requests
- the European Food Safety Authority to issue an opinion on the safety and suitability for use by infants 203
- 204 of a follow-on formula based on cow's milk intact protein with a protein content of at least 1.61 q/100
- 205
- 206 If the formula under evaluation is considered to be safe and suitable for use by infants, the European
- Food Safety Authority is asked to advise whether a level of at least 1.61 g protein/100 kcal would be 207
- applicable to all follow-on formulae. If this is not the case, the Authority is asked to advise on the 208
- specific criteria that need to be satisfied for the safety and suitability of such formulae to be 209
- demonstrated. 210

Interpretation of the Terms of Reference 1.2. 211

- The Panel interprets the terms of reference provided by the European Commission in the context of 212
- 213 the background information given and the application submitted. The Panel understands that the
- European Commission seeks advice on: 214
- (a) whether a follow-on formula (FOF) based on cow's milk intact protein with a minimum protein 215
- content of 1.6 g/100 kcal (rounded value) is safe and suitable for infants provided that it meets otherwise the requirements⁵ of relevant EU legislation⁶, and if so 216
- 217
- (b) whether FOF based on goat's milk intact protein, soy protein isolates or protein hydrolysates with 218
- 219 a minimum protein content of 1.6 g/100 kcal (rounded value) are also safe and suitable for infants
- provided that they meet otherwise the requirements of relevant EU legislation. 220

Commission Directive 2006/141/EC of 22 December 2006 on infant formulae and follow-on formulae and amending Directive 1999/21/EC, OJ L 401, 30.12.2006, p. 1

Commission Delegated Regulation (EU) 2016/127 of 25 September 2015 supplementing Regulation (EU) No 609/2013 of the European Parliament and of the Council as regards the specific compositional and information requirements for infant formula and follow-on formula and as regards requirements on information relating to infant and young child feeding, OJ L 25, 2.2.2016, p. 1

⁴ Regulation (EC) No 178/2002 of the European Parliament and of the Council of 28 January 2002 laying down the general principles and requirements of food law, establishing the European Food Safety Authority and laying down procedures in matters of food safety, OJ L 31, 1.2.2002, p. 1

Including the requirements with respect to the amino acid profile

⁶ Directive 2006/141/EC to be replaced by delegated Regulation (EU) 2016/127



2. Data and Methodologies

222 **2.1.** Data

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EFSA was provided with a dossier related to a FOF based on cow's milk intact protein containing a 223 minimum of 1.61 g protein per 100 kcal but otherwise complying with the compositional criteria laid 224 down in Directive 2006/141/EC. The dossier includes two intervention studies in infants, named "US 225 study" (Haves and Northington, 2014, unpublished study report #1; published as Ziegler et al., 2015), 226 227 and "Chile study" (Yao, 2014, unpublished study report #2; published as Inostroza et al., 2014). The dossier was supplemented, upon request of EFSA, with additional information provided by the 228 applicant on 28 June 2016 and on 29 September 2016. The intervention studies provided in the 229 230 dossier were designed to assess the growth pattern of infants receiving a formula with standard 231 protein content for the first three months of life and thereafter a formula with a protein content which 232 is lower than currently authorised. These studies aimed to investigate whether lower protein content 233 in formula to be fed from three to 12 months of age, in line with the decrease in the protein content of breast milk during that feeding period, would lead to growth rates closer to those of breast-fed 234 infants, as compared to infants fed a "standard" formula. 235

The Panel will also take into account in the current assessment its previous opinions on Dietary Reference Values for protein (EFSA NDA Panel, 2012a), on nutrient requirements and dietary intake of infants and young children in Europe (EFSA NDA Panel, 2013), and on the essential composition of IF and FOF (EFSA NDA Panel, 2014), as well as data on the protein content of breast milk.

2.2. Methodologies

As outlined in the Panel's previous opinion on the essential composition of IF and FOF (EFSA NDA Panel, 2014), the minimum amounts of nutrients in formulae, including protein, should be based on generally accepted scientific evidence. While for IF compositional requirements may be based on the energy and nutrient requirements of infants and on the results of intervention studies in the target population in which the formula is the only source of energy and nutrients, evidence for proposing compositional requirements for foods which are not the sole source of energy and nutrients, such as FOF, is less strong, as other foods contribute to nutrient and energy intake in variable amounts. In its previous opinion, when proposing compositional requirements for FOF, the Panel assumed that complementary food (CF) would compensate for the higher energy and nutrient requirements of older infants and for the lower formula intake during that period. This is based on the assumption that infants in the target population have access to CF of a sufficient quality, following established feeding guidelines in Europe (e.g. from Member States).

For the present assessment of whether a FOF based on cow's milk intact protein with a protein content of at least 1.6 g/100 kcal (rounded value) is safe and suitable for infants provided that it meets otherwise the requirements⁷ of relevant EU legislation⁸, the Panel will consider:

- a) Dietary protein requirements of infants in the second half of the first year of life;
- b) Protein content in breast milk during the first year of lactation;
- c) Dietary protein intake of infants in Europe from breast milk, formula and CF;
- d) The overall contribution that a FOF with a protein content of 1.6 g/100 kcal could make towards protein requirements in the target population, assuming access to CF of a sufficient quality, following established feeding guidelines in Europe (e.g. from Member States);
- e) The application submitted by the food business operator, including two intervention studies in healthy term infants.

The evaluation of the intervention studies provided by the food business operator will follow the general principles for the assessment of a modification of the composition of IFs or FOFs outside the established standards as laid down by the Scientific Committee on Food (SCF) (SCF, 2003). In addition, the recommendations for the assessment of the safety and suitability of formulae for term

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⁷ Including the requirements with respect to the amino acid profile

⁸ Directive 2006/141/EC to be replaced by delegated Regulation (EU) 2016/127



infants of the European Society for Paediatric Gastroenterology, Hepatology and Nutrition (ESPGHAN)
(Aggett et al., 2001), of the Committee on the Evaluation of the Addition of Ingredients New to Infant
Formula of the Food and Nutrition Board of the United States (US) Institute of Medicine (IoM, 2004)
and of the American Academy of Pediatrics (AAP, 1988), will be taken into account.

3. Assessment

3.1. Dietary protein requirements of infants in the second half of the first year of life

Dietary protein is an essential component of the diet, supplying the body with nitrogen (N) and amino acids as well as other non-protein metabolically active nitrogenous substances. The protein requirement of infants and young children comprises two components, the maintenance requirement and the growth requirement (EFSA NDA Panel, 2012a). In its previous opinion, the Panel established an average maintenance requirement of 0.66 g protein/kg body weight per day (105 mg N/kg body weight per day) for infants and young children aged from 6 to < 36 months, which was derived from nitrogen balance studies in adults. The average protein requirement for growth was estimated from average daily rates of protein deposition calculated from studies on whole-body potassium deposition, and adjusted by an efficiency of utilisation of dietary protein for growth of 58 %. Together, these amounts constitute an AR, to which 1.96 standard deviations were added to derive a PRI. Thus, a PRI of 10 g protein per day for girls and 11 g protein per day for boys aged 6 months and a PRI of 11 g protein per day for girls and 12 g protein per day for boys aged 12 months were established.

3.2. Protein content of breast milk during the first year of lactation

Estimating the true protein content of breast milk is challenging because of the non-protein nitrogen fraction contained in it. Total nitrogen in human milk represents both protein, about 75%, and non-protein nitrogen, which is made up of urea (up to 50% of the non-protein nitrogen), amino acids and other nitrogenous compounds (SCF, 2003; WHO/FAO/UNU, 2007). The amount of nitrogen used by infants for protein synthesis is likely to include that from true protein, free amino acids and small peptides, and a proportion of urea nitrogen. Therefore, the amount of nitrogen in breast milk used for protein synthesis by infants is between the true protein content and the crude protein calculated from total nitrogen.

A meta-analysis of 41 published studies reporting on pre-term (26 studies, 843 mothers) and term (30 studies, 2299 mothers) breast milk composition is available (Gidrewicz and Fenton, 2014). Energy was estimated in 11 studies using bomb calorimetry, and in five studies by calculation using values for the energy contributions from fat, protein, and carbohydrate. Protein was estimated based on total nitrogen in 23 studies and as a true protein estimate in 15 studies. Data on mean energy and protein content of breast milk from mothers of term infants by week of lactation is shown in **Table 1**.



Table 1: Mean energy and protein content of breast milk from mothers of term infants by week of lactation¹

	Mean ene (kcal/1		Mean prot	ein (SD) (g/100mL)	Mean protein (SD) (g/100kcal)		
Time	Bomb calorimetry	Calculated	True protein	Protein calculated from total nitrogen	True protein	Protein calculated from total nitrogen	
4-7 d	66 (9)	68 (10)	1.6 (0.3)	2.0 (0.5)	2.4 (0.5)	2.9 (0.7) -3.0 (0.8)	
2 wk	66 (9)	-	1.3 (0.2)	1.8 (0.4)	2.0 (0.3)	2.7 (0.63)	
3-4 wk	66 (8)	70 (9)	1.1 (0.2)	1.5 (0.3)	1.6 (0.3) -1.7 (0.3)	2.1 (0.4) -2.3 (0.5)	
5-6 wk	63 (7)	-	1.0 (0.1)	1.1 (0.2)	1.6 (0.2)	1.7 (0.3)	
7-9 wk	63 (7)	69 (10)	0.9 (0.1)	1.3 (0.2)	1.3-1.4 (0.2)	2.1 (0.3)	
10-12 wk	63 (8)	68 (9)	1.0 (0.1)	1.2 (0.2)	1.5-1.6 (0.2)	1.8 (0.3) -1.9 (0.3)	

¹Adapted from Gidrewicz and Fenton (2014)

The true protein content of breast milk gradually decreased from the first week of lactation, being about 1.0 g/100mL (corresponding to about 1.8-1.9 g/100kcal) by the third month.

Table 2 shows the energy (calculated) and the interval from birth according to the true protein content of breast milk samples (n=2554) donated by 224 mothers of mostly term infants to a milk bank in Denmark (Michaelsen et al., 1990). The mean true protein content of all samples combined was 0.9 g/100mL.

Table 2: Time interval from birth according to the protein content of breast milk¹

True protein (g/100mL)	Mean interval from birth (weeks)	Mean (SD) energy (kcal/100mL)	True protein (g/100kcal)	No. of samples
≥ 1.3	3-4	74.7 (9.8)	≥ 1.7	70
1.1 – 1.29	6-7	71.4 (9.8)	1.5-1.8	193
0.9 – 1.09	11-12	67.4 (9.8)	1.3-1.6	572
0.7 – 0.89	15-16	64.6 (9.8)	1.1-1.4	800
< 0.7	19-20	64.2 (9.3)	< 1.1	108

¹Adapted from Michaelsen et al. (1990)

The mean interval from birth gradually decreased with the increasing content of protein in breast milk. It was 3-4 weeks for samples containing ≥ 1.3 g/100mL and 19-20 weeks for samples containing < 0.7 g/100mL. The mean interval from birth was 11-12 weeks for samples containing 0.9-1.09 g/100mL, which is consistent with the protein content of breast milk by the third month of lactation (1.0 g/100mL) reported by Gidrewicz and Fenton (2014).

Other studies have reported on the protein content of breast milk during the first year of lactation, although the number of breast milk samples analyses in those studies is low (Allen et al., 1991; Nommsen et al., 1991; Mitoulas et al., 2002).

In the context of the DARLING study, Nommsen et al. (1991) assessed the composition of breast milk in samples taken at 3, 6, 9 and 12 months of lactation in healthy mothers of term infants. The gross energy content and the total protein content of the breast milk samples are given in **Table 3**. Protein was analysed using a modified Lowry assay with bovine serum albumin as the standard, a method which tends to result in slightly elevated values for total protein (Nommsen et al., 1991).



Table 3: Gross energy and total protein content of breast milk during the first year of lactation¹

Month of lactation	n	Mean (SD) gross energy (kcal/100mL)	Mean (SD) total protein (g/100mL)	Mean (SD) total protein (g/100kcal)
3	58	697 (97)	1.2 (0.2)	1.7 (1.5)
6	45	707 (92)	1.1 (0.2)	1.6 (1.6)
9	28	709 (74)	1.2 (0.8)	1.6 (10.8)
12	21	706 (110)	1.2 (0.2)	1.7 (1.3)

¹Adapted from Nommsen et al. (1991)

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The mean total protein at three months of lactation is comparable to the protein content calculated from total nitrogen (1.2 g/100mL, 1.8 g/100kcal) for that time period in the meta-analysis by Gidrewicz and Fenton (2014), and remained fairly stable until the twelfth month of lactation, which is in agreement with what has been reported in other studies (Allen et al., 1991; Mitoulas et al., 2002).

The Panel notes that the mean content of true protein in breast milk by the end of the third month of lactation ranges between 1.3 and 1.6 g/100kcal, tends to decrease thereafter to about 1.1-1.4 g/100kcal by the end of the fourth month, and tends to remain fairly stable thereafter.

3.3. Dietary protein intake of infants in Europe

Data on mean energy and protein intake in infants living in Europe were gathered from published studies (**Table 4**). Details about the dietary data collection and on the assessment of breast milk intake are given in **Table 5**.

Data on mean energy and protein intake in infants living in Europe were also gathered form dietary surveys for which sufficient data were available in the EFSA Comprehensive European Food Consumption Database (**Table 6**)⁹.

From the dietary surveys available in the EFSA Comprehensive European Food Consumption Database, mean energy and protein intake from formula and from CF in non-breastfed infants, and from CF only in (exclusively or partially) breastfed infants were also calculated (**Table 7**). Mean energy and protein intake by food group in non-breastfed infants are given in **Tables 8** (for infants 4 to <6 months of age) and **9** (for infants aged ≥6 to 12 months).

Details about the dietary surveys included in the EFSA Comprehensive Database are available at: https://dwh.efsa.europa.eu/bi/asp/Main.aspx?rwtrep=001



Table 4: Mean energy and protein intake in infants living in Europe from published studies

Age	Country	Study	Breastfeeding ¹ (%)	N	Mean energy (SD) ² (kcal/d) ³	Mean protein (SD) ² (g/d)	P5 th of protein intake ³	P2.5 th of protein intake ⁴	Mean E% as protein ²
			All males	262	658 (123)	15.7 (3.4)	10.1	9.0	9.5
			All females	214	604 (118)	14.5 (3.3)	9.04	8.0	9.6
			100, no solids	53	626 (-)	12 (-)	-	-	7.6
4 mo	UK	ALSPAC	0, no solids	42	583 (-)	13 (-)	-	-	8.9
			100, solids	209	646 (-)	13 (-)	-	-	8.1
			0, solids	441	640 (-)	15 (-)	-	-	9.4
			Mixed, solids	107	667 (-)	14 (-)	-	-	8.4
	France	National	0	58	668	20.0 (5.60)	10.7	-	11.9
	UK	Southampton	28	50	709 (652-818) ⁵	19.7 (17.4-23.6) ⁵	-	-	11.17
	Germany	DONALD	48	302	645 (119)	16.3 (5.4)	7.4	5.7	10.1
	Belgium				679 (659-702) ⁶	19 (17-20) ⁶	-	-	11.2
6 mo	Germany				634 (615-655) ⁶	19 (18-20) ⁶	-	-	12.0
	Italy	CHOP	0	746	731 (710-753) ⁶	23 (22-24) ⁶	-	-	12.6
	Poland				725 (702-743) ⁶	21 (20-22) ⁶	-	-	11.6
	Spain				736 (719-754) ⁶	18 (18-19) ⁶	-	-	10.0
8 mo	Belgium				734 (707-761) ⁶	24 (22-25) ⁶	-	-	13.1
2 1110	Germany	CHOP	0	625	694 (670-718) ⁶	20 (19-22) ⁶	-	-	11.5



Age	Country	Study	Breastfeeding ¹ (%)	N	Mean energy (SD) ² (kcal/d) ³	Mean protein (SD) ² (g/d)	P5 th of protein intake ³	P2.5 th of protein intake ⁴	Mean E% as protein ²
	Italy				817 (796-838) ⁶	28 (27-28) ⁶		-	13.7
	Poland				812 (783-840) ⁶	25 (24-26) ⁶	-	-	12.3
	Spain				854 (831-878) ⁶	30 (29-32) ⁶	-	-	14.1
	Finland	STRIP	0	215	842 (148)	25.0 (6)	15.1	13.2	11.9
				618 m	840 (173)	29.0 (9.0)	14.1	11.4	13.8
	UK	ALSPAC	0	513 f	784 (156)	27.0 (8.0)	13.8	11.3	13.8
	Belgium				770 (739-801) ⁶	24 (22-25) ⁶	-	-	13.5
	Germany				708 (684-731) ⁶	20 (19-22) ⁶	-	-	11.9
	Italy	CHOP	0	617	850 (826-874) ⁶	28 (27-28) ⁶	-	-	14.1
	Poland				859 (850-892) ⁶	25 (24-26) ⁶	-	-	12.6
9 mo	Spain				872 (846-899) ⁶	30 (29-32) ⁶	-	-	14.7
		1995–1996	37	80	760 (678-859) ⁵	28.0 (21.7-35.3) ⁵	-	-	14.47
	Iceland	2005	41	154	754 (629-859) ⁵	22.7 (17.8-27.6) ⁵	-	-	11.97
	Netherlands	TNO		333	970 (175)	28.8 (6.2)	18.6	16.6	11.9
	Germany	DONALD	17	332	759 (122)	22.4 (5.3)	13.6	12.0	11.8
10-12 mo	France	National	0	63	826 (160)	30.0 (11)	11.8	-	14.4

¹ Either exclusive or partial breastfeeding; ² Unless otherwise noted; ³ Calculated from the mean and the SD assuming a normal distribution of intake; ⁴ Calculated only for sample sizes n≥180; Median (interquartile range); ⁶ Mean (95% confidence interval); ⁷ Median; ⁻ = not reported or not available; m= males; f = females.



Table 5: Methods for dietary assessment, estimation of breast milk intake and food composition databases used in published studies

Country	Study	Age (months)	Publications	Dietary assessment	Breast milk intake
France	National	6, 10-12	Fantino & Gourmet, 2008	3-d weighed-DR	
Germany	DONALD	3, 6	Hilbig, 2005	3-d weighed-DR	Measured by 'test-weighing' the infant before and after each breast milk meal
UK	ALSPAC	4	Noble and Emmett, 2006	24-h recall	Duration of each breastfeeding was used to estimate the volume of milk consumed; a feed lasting ≥ 10 min was assumed to be 125 mL, or a proportion of this if the feed was of shorter duration (i.e. 12.5 mL for 1 min)
UK	Southampton	6	Marriott et al., 2008	4-d weighed-DR	Estimated using an algorithm based on length of suckling derived from published intake data
5 EU countries	СНОР	6-12	Damianidi et al., 2006	3-d weighed-DR	-
Finland	STRIP	8	Lagström et al., 1997	3-d DR	-
UK	ALSPAC	8	Noble et al., 2001	3-d DR	-
Iceland	1995–1996 2005	9	Thorisdottir et al., 2013	2-d or 3-d weighed DR	Measured by 'test-weighing' the infant before and after each breast milk meal
Netherlands	TNO	9	de Boer et al., 2006	2-d DR	NR
Germany	DONALD	9	Schwartz et al., 2010	3-d weighed-DR	Measured by 'test-weighing' the infant before and after each breast milk meal

DR = Dietary records; NR = not reported.



Table 6: Mean energy and protein intake in infants living in Europe from the EFSA Comprehensive European Food Consumption Database

Age	Breastfeeding ¹	Country	Survey	Dietary assessment	N	Mean energy intake (kcal/d)	Mean protein intake (g/d)	P5 th of protein intake (g/d)	P2.5 th of protein intake (g/d)	Mean E% as protein
	Yes	Bulgaria	NUTRICHILD	24-h recall , 3 d ^{2,5}	64	679	11.2	6.9	-	7
		Denmark	IAT 2006_07	7-d DR ^{3,6}	26	746	15.9	-	-	9
4-<6		UK	DNSIYC_2011	4-d DR ^{4,7}	27	686	15.0	-	-	9
mo		Bulgaria	NUTRICHILD	24-h recall, 3 d	88	639	16.3	9.7	-	10
	No	Denmark	IAT 2006_07	7-d DR	12	745	18.2	-	-	10
		UK	DNSIYC_2011	4-d DR	49	639	15.9	-	-	10
		Bulgaria	NUTRICHILD	24-h recall, 3 d ^{2,5}	89	905	22.2	8.3	-	10
	Yes	Denmark	IAT 2006_07	7-d DR ^{3, 6}	315	832	23.0	10.1	8.9	11
≥6-		UK	DNSIYC_2011	4-d DR ^{4,7}	264	804	24.5	13.0	12.0	12
12mo		Bulgaria	NUTRICHILD	24-h recall, 3 d	343	859	27.4	14.1	11.9	13
	No	Denmark	IAT 2006_07	7-d DR	473	933	30.0	16.2	14.7	13
		UK	DNSIYC_2011	4-d DR	1029	790	25.2	13.0	11.7	13

^{- =} $P5^{th}$ and $P2.5^{th}$ of protein intake are only provided for study groups with a sample size ≥ 60 and 180 individuals, respectively.

DR = dietary records; ¹ Either exclusive or partial breastfeeding; ² Method to estimate volume of breast milk intake per feeding occasion not reported, assumed to be 130 mL per feeding occasion; ³ Volume of breast milk per feeding occasion calculated as follows (Dewey et al., 1984): 130 mL per breastfeeding if the infant was breastfed 6 times or more per day; 89 mL per breastfeeding if the infant was breastfed 3-5 times per day; 53 mL per breastfeeding if the infant was breastfed 1-2 times per day; ⁴ Volume of breast milk per feeding occasion calculated based on the time for each feed, at 13.5g/min with a maximum of 135g per feed for infants aged 4 to 7 months and at 10g/min with a maximum of 100g per feed for infants aged 8 to 12 months; ⁵ Breast milk assumed to contain 70.0 kcal/100mL; 1.0 g of protein/100mL and 1.4 g of protein/100 kcal; ⁶ Breast milk assumed to contain 71.0 kcal/100mL; 1.3 g of protein/100mL and 1.9 g of protein/100 kcal



Table 7: Mean energy and protein intake from formula and complementary food in infants from surveys in the EFSA Comprehensive European Food Consumption Database

Age	Breastfeeding ¹	Country	N	Food group	Mean energy intake (kcal/d)	Mean protein intake (g/d)	Mean protein intake (g/100kcal) ²
		Bulgaria	64	CF	210	4.1	-
	Yes	Denmark	26	CF	255	6.0	-
		UK	27	CF	211	3.7	-
		Bulgaria	88	Formula ³	298	6.5	2.18
4-<6 mo	No			CF	341	9.8	-
		Denmark	12	Formula	495	12.0	2.42
				CF	250	6.2	-
		UK	49	Formula	493	10.9	2.21
				CF	146	6.0	-
		Bulgaria	89	CF	554	17.3	-
	Yes	Denmark	315	CF	540	17.0	-
		UK	264	CF	405	16.6	-
.		Bulgaria	343	Formula	91	2.2	2.42
≥6- 12mo				CF	749	24.8	-
121110	No	Denmark	473	Formula	225	6.1	2.71
	No			CF	708	23.9	-
		UK	1029	Formula	349	7.4	2.12
				CF	441	17.6	-

CF = Complementary food; ¹ Either exclusive or partial breastfeeding; ² Calculated from mean energy and protein intake from formula; ³ Any (infant and follow-on) formula

366367



Table 8: Mean energy and protein intake by food group in non-breastfed infants aged 4 to 6 months

	Bulgaria	a (n=88)	Denmar	k (n=12)	UK (ı	n=49)
	Mean energy	Mean protein	Mean energy	Mean protein	Mean energy	Mean protein
	(kcal/d)	(g/d)	(kcal/d)	(g/d)	(kcal/d)	(g/d)
Food group						
Animal and vegetable fats and oils	8.6	0.0	27.4	0.0	1.0	0.0
Composite food (including frozen products)	0.2	0.0	0.0	0.0	0.4	0.0
Eggs and egg products	3.1	0.2	0.0	0.0	0.4	0.0
Fish and other seafood (including amphibians, reptiles, snails and insects)	0.0	0.0	0.4	0.1	0.7	0.1
Food for infants and small children	129.5	2.4	105.5	3.0	95.1	3.0
Fruit and fruit products	7.9	0.1	14.7	0.1	11.5	0.1
Fruit and vegetable juices	15.6	0.1	0.0	0.0	0.7	0.0
Grains and grain-based products	26.6	0.6	47.5	1.1	4.7	0.1
Herbs, spices and condiments	0.0	0.0	0.0	0.0	0.3	0.0
Infant formula and follow-up formula	298.0	6.5	495.2	12.0	493.4	10.9
Legumes, nuts and oilseeds	0.4	0.0	3.7	0.3	1.9	0.2
Meat and meat products (including edible offal)	7.3	0.6	1.0	0.1	2.3	0.4
Milk and dairy products	98.1	5.5	11.1	0.7	13.5	0.8
Snacks, desserts, and other foods	2.1	0.1	0.0	0.0	3.9	0.1
Starchy roots and tubers	9.0	0.2	27.8	0.6	5.0	0.1
Sugar and confectionary	31.6	0.0	4.7	0.0	0.7	0.0
Vegetables and vegetable products (including fungi)	1.2	0.0	5.9	0.3	3.7	0.2

369



Table 9: Mean energy and protein intake by food group in non-breastfed infants aged 6 to 12 months

	Bulgaria	(n=343)	Denmarl	k (n=473)	UK (n:	=1029)
	Mean energy	Mean protein	Mean energy	Mean protein	Mean energy	Mean protein
Food group	(kcal/d)	(g/d)	(kcal/d)	(g/d)	(kcal/d)	(g/d)
Animal and vegetable fats and oils	70.4	0.0	87.0	0.0	15.8	0.0
Composite food (including frozen products)	1.3	0.1	0.0	0.0	19.2	0.9
Eggs and egg products	13.0	0.9	2.5	0.2	2.9	0.2
Fish and other seafood (including amphibians, reptiles, snails and insects)	1.3	0.2	7.9	1.0	7.2	0.8
Food for infants and small children	99.7	2.4	68.9	1.9	121.4	4.0
Fruit and fruit products	35.6	0.3	62.4	0.7	33.2	0.4
Fruit and vegetable juices	24.8	0.2	4.7	0.0	1.7	0.0
Grains and grain-based products	183.1	4.3	186.3	5.3	78.7	2.4
Herbs, spices and condiments	0.9	0.1	2.0	0.0	3.9	0.1
Infant formula and follow-up formula	90.8	2.2	225.1	6.1	349.0	7.4
Legumes, nuts and oilseeds	6.5	0.4	5.9	0.4	5.5	0.4
Meat and meat products (including edible offal)	48.3	5.2	49.4	3.7	21.9	2.9
Milk and dairy products	166.5	9.2	154.5	8.7	79.3	4.4
Snacks, desserts, and other foods	7.7	0.1	12.0	0.4	12.1	0.2
Starchy roots and tubers	24.0	0.6	34.8	0.8	20.9	0.4
Sugar and confectionary	50.6	0.0	17.1	0.0	5.9	0.1
Vegetables and vegetable products (including fungi)	14.3	0.6	12.5	0.6	10.4	0.5



Mean protein intakes from all sources were beyond the PRI for protein in all surveys from the EFSA Comprehensive Database for both breastfed and formula-fed infants aged 6 to 12 months (**Table 6**). Mean protein intake from all sources were also reported in published studies which accurately estimated breast milk intake (by weighing the infant before and after each breast milk meal) and/or which used more accurate methods for dietary assessment (3 or 4-day weighted dietary records). At 4 months of age, the lowest mean protein intake (12 g per day) were reported for infants exclusively breastfed in the ALSPAC cohort. Mean protein intakes were slightly higher (13 g per day) in breastfed infants who had already received some solid food. The lowest mean protein intake (16.3 g per day) for infants aged 6 months or older was reported in a German cohort (DONALD study), in which the proportion of breastfed infants was the highest (48%) among all the studies available (**Table 4**). Mean protein intakes from all sources were beyond the PRI for protein for infants aged 6 to 12 months in all the studies.

In the dietary surveys for which data on (exclusively or partially) breastfed infants and formula-fed infants was available separately (**Table 6**), mean protein intakes were systematically higher in formula-fed infants than in breastfed infants, as previously reported by others (Heinig et al., 1993). Breast milk was assumed to contain from 1.4 to 1.9 g of protein/100 kcal, depending on the survey. The lower mean protein intake reported for Bulgarian infants could be explained in part by the assumed lower protein content in breast milk (1.4 g/100 kcal). The protein content of formula ranged from 2.1 to 2.7 g of protein/100 kcal, depending on the survey and age category (**Table 7**). This is higher than the minimum protein content allowed by EU legislation (Directive 2006/141/EC and Commission delegated Regulation (EU) 2016/127) for (infant and follow-on) formula manufactured from intact cows' or goats' milk proteins (1.8 g/100 kcal).

In breastfed infants, mean protein intake from CF ranged between 3.7 and 6.0 g/day in infants aged 4 to <6 months, and were already well beyond the PRI for protein in infants aged 6-12 months (about 17 g/day). In formula-fed infants aged 4 to 6 months, mean protein intake from formula ranged from 6.5 to 12.0 g per day, whilst mean protein intakes from CF were about 6 g/day. In Bulgaria, where mean protein intake from formula was the lowest (6.5 g/day), mean protein intake from CF was much higher (9.8 g/day), mostly coming from cow's milk and dairy products other than formula (**Table 8**). This is due to a replacement of IF with cow's milk (rather than with FOF) at the time of the introduction of CF. In formula-fed infants aged 6 to 12 months, mean protein intakes from CF were at or beyond the PRI in all countries. The contribution of formula to total protein intake varied widely, being lower in countries (Bulgaria and Denmark) with the highest protein intake from cow's milk and dairy products and from meat and meat products (**Table 9**).

Whenever the data available allowed doing so, the 5th and the 2.5th percentiles (P5th and P2.5th, respectively) of protein intake were calculated by assuming a normal distribution of protein intake data **(Table 4)** or extracted from individual data **(Table 6)**. Otherwise, IQRs or 95% CI were considered **(Table 4)**. The Panel notes that the P5th and P2.5th of total protein intake in non-breastfed infants aged 6 to 12 months was around or above the PRI for protein for that age group in all the studies **(Table 4)** and surveys **(Table 6)** available.

3.4. Contribution that a FOF with a protein content of at least 1.6 g/100 kcal could make towards protein requirements in the target population

Consumption of a FOF with a protein content of about 1.6 g/100 kcal would provide about 9 g of protein per day in the first months of complementary feeding (assuming an intake of about 500 mL per day) and about 4.5 g of protein per day (assuming an intake of about 250 mL per day) by the end of the first year of life. This is about 1 g and 0.5 g of protein less than the estimated intake from a formula containing a minimum of 1.8 g/100 kcal, as currently authorised. The Panel notes, however, that the protein content of (infant and follow-on) formula in the European surveys available (from 2.1 to 2.7 g of protein/100 kcal, **Table 7**) was higher than the minimum authorised.

Using individual data from the three surveys which were available in the EFSA Comprehensive Food Consumption Database, total protein intake in non-breastfed infants aged 6 to 12 months was recalculated by assuming that: a) all FOF consumed by the infants contained 1.6 g of protein/100 kcal; b) the energy content of the individual FOFs did not change; c) protein intake from other sources



(IF, CF) did not change. The mean, P5th and P2.5th of total protein intake under these conditions are shown in **Table 10**.

Table 10: Protein intake in European non-breastfed infants aged 6 to 12 months, assuming a protein content of FOF of 1.6 g/100kcal

Country	N	Mean energy intake (kcal/d)	Mean protein intake (g/d)	P5 th of protein intake (g/d)	P2.5 th of protein intake (g/d)
Bulgaria	343	859	27.2	13.7	11.9
Denmark	473	933	29.4	15.8	13.8
UK	1029	790	24.4	12.6	11.2

As expected, total protein intakes resulting from the consumption of FOF with a protein content of $1.6 \, \text{g}/100 \text{kcal}$ would be lower than those reported in the original surveys **(Table 6)**. The Panel notes, however, that the P5th and P2.5th of total protein intake would remain at about or above the PRI for protein for infants aged 6 to 12 months.

3.5. Application submitted by the food business operator

The applicant provided two human intervention studies aiming to investigate whether protein content in formula to be fed from three to 12 months of age that is closer to the protein content of breast milk during that feeding period would lead to growth rates more in line to those of breast-fed infants, as compared to infants fed a "standard" formula.

3.5.1. Composition of the formulae used in the two human intervention studies

The formulae investigated in the US and Chile studies contain a minimum of 1.61 g protein per 100 kcal, based on a calculation of total nitrogen x 6.25. The protein source is based on intact proteins derived from skimmed milk and a proprietary preparation of demineralised whey. The demineralised whey is obtained from modified caseino-glyco-macro-peptide (CGMP)-reduced sweet whey produced using a patented process (Patent No PCT/EP1998/003176). The whey preparation used in the formula has a CGMP content which is reduced by at least 85%. The whey protein-to-casein ratio of the final product is 60:40. The applicant indicated that the protein sources have been used in other FOF currently marketed by the applicant, and that the use of CGMP-reduced sweet whey has allowed for a lower protein content of the FOF, while still meeting the requirements of Directive 2006/141/EC with respect to the amino acid pattern. The energy content and the amount of carbohydrates, fat, vitamins and minerals also comply with the compositional requirements laid down in Directive 2006/141/EC.

The macronutrient composition of the intervention and control formulae used in the US and the Chile studies are outlined in **Table 11**.

Table 11: Macronutrient composition of study formulae in comparison to the compositional requirements for FOF manufactured from cow's or goat's-milk proteins as laid down in Directive 2006/141/EC

	Unit	Directive 2006/141/EC	US stu	ıdy	Chile study		
			Intervention	Control	Intervention	Control	
Energy	kcal/100 ml	60-70	67.2	64.6	62.8	65.6	
Protein	g/100 kcal	1.8-3.5	1.61	2.15	1.65	2.70	
Fat	g/100 kcal	4.0-6.0	5.46	5.21	5.30	5.03	
Carbohydrates	g/100 kcal	9.0-14.0	11.10	11.13	11.41	10.98	
Cyst(e)ine	mg/100 kcal	38	28	38	28	46	
Histidine	mg/100 kcal	40	40	49	39	64	
Isoleucine	mg/100 kcal	90	95	125	100	166	
Leucine	mg/100 kcal	166	166	222	180	298	
Lysine	mg/100 kcal	113	132	185	142	234	
Methionine	mg/100 kcal	23	33	46	42	69	
Phenylalanine	mg/100 kcal	83	103	88	108	179	



Threonine	mg/100 kcal	77	94	141	84	137
Tryptophan	mg/100 kcal	32	31	31	34	57
Tyrosine	mg/100 kcal	76	52	68	69	113
Valine	mg/100 kcal	88	94	137	102	168
Methionine + Cyst(e)ine	mg/100 kcal	61 ^(a)	61	84	70	115
ratio			1.2	1.2	1.5	1.5
Tyrosine + Phenylalanine	mg/100 kcal	159 ^(b)	155	156	177	292
ratio			0.5	0.8	0.6	0.6

(a): The concentrations of cyst(e)ine and methionine may be added together if the methionine:cyst(e)ine-ratio is not >3

(b): The concentrations of tyrosine and phenylalanine may be added together if the tyrosine:phenylalanine-ratio is not >2

The applicant states that the tyrosine and phenylalanine content (calculated as sum) and tryptophan content in the intervention and control formulae of the US study were slightly lower than required by Directive 2006/141/EC, and that the histidine content in the intervention formula of the Chile study was slightly lower than required by Directive 2006/141/EC, but that the final marketed product will comply with the specifications laid down in the Directive.

In the Chile study the intervention formula also contained 2 x 10⁷ colony forming units (CFU) *Bifidobacterium lactis* (CNCM I-3446) and 2 x 10⁷ CFU *Lactobacillus rhamnosus* (CGMCC 1.3724) per
gram of powder formula, while the control formula did not contain these bacteria.

The intervention formulae in both studies had a whey protein-to-casein ratio of 60:40.

3.5.2. Human intervention studies

 The two randomised, double-blind, controlled intervention studies were conducted in Chile (Inostroza, et al., 2014) and in the US (Ziegler et al., 2015). These studies assessed growth rates in healthy term infants who consumed (low-protein, intervention) formulae with protein contents of $1.61 \, \text{g}/100 \, \text{kcal}$ (n = 97) (US study) and $1.65 \, \text{g}/100 \, \text{kcal}$ (n = 89) (Chile study) from three months of age onwards, against those of infants who consumed (control) formulae with protein contents of $2.15 \, \text{g}/100 \, \text{kcal}$ (n = 97) and $2.70 \, \text{g}/100 \, \text{kcal}$ (n = 87), respectively, and against those of a breastfed reference group (n= 76 and n = 112, respectively). In the Chile study, only infants from overweight and obese mothers were recruited.

In both studies, the primary outcome was weight gain between 3 and 6 months of age. Secondary outcomes included, amongst others, weight gain at time points beyond 6 months of age, weight changes, changes in length and head circumference, and changes in serum albumin and blood urea nitrogen (BUN). Adverse events were registered.

Statistical analyses were conducted in completers and per protocol (PP) in both studies. In the US study, 10 infants in the intervention group, 10 infants in the control group and 7 infants in the breastfed reference group discontinued the study. The numbers in the Chile study were 23, 11 and 11, respectively. Reasons for withdrawal were provided.

Despite the original protocols foreseeing exclusive formula or breastfeeding up to the age of 6 months and the introduction of CF thereafter (control formula was allowed from 6 to 12 months to the breastfed reference groups, if desired), small amounts of CF were provided to some infants from around 4 months of age onwards. In the US study, a total of 9 infants consumed >4 teaspoons of CF per day before the age of 6 months and were excluded from the PP analysis. In the Chile study, CF in amounts > 4 teaspoons per day were introduced before 6 months of age in 66 infants (28 in the intervention, 24 in the control and 14 in the breastfed reference group), who were not excluded from the statistical analysis.

The information provided in these studies did not allow the calculation of energy and protein intake from CF at any time point, and thus of the total energy and protein intake at time points in which infants consumed CF (4-12 months in the Chile study, 6-12 months in the US study).

496 Mean daily energy and protein intake from formula at different time points in the US and Chile studies 497 are given in Table 12.



Table 12: Mean daily energy and protein intake from formula at different time-points in the US and Chile studies

US Study									
	n	Mean (SD) volume intake (mL/day)	Mean (SD) energy intake (kcal/day)	Mean (SD) protein intake (g/day)					
4 months									
Low-protein formula	83	905 (216)	581 (139)	14.6 (3.5)					
Control formula	85	894 (180)	601 (121)	19.2 (3.9)					
6 months									
Low-protein formula	83	917 (232)	589 (149)	14.8 (3.7)					
Control formula	84	902 (184)	606 (124)	19.4 (4.0)					
8 months									
Low-protein formula	80	850 (208)	546 (133)	13.7 (3.3)					
Control formula	82	857 (179)	606 (124)	18.4 (3.8)					
12 months									
Low-protein formula	76	719 (239)	462 (153)	11.6 (3.8)					
Control formula	78	725 (241)	487 (162)	15.6 (5.2)					
		Chile study							
	n	Mean (SD) volume intake (mL/day)	Mean (SD) energy intake (kcal/day)	Mean (SD) protein intake (g/day)					
4 months			•	•					
Low-protein formula	75	820 (268)	515 (168)	13.5 (4.4)					
Control formula	80	868 (228)	569 (150)	23.4 (6.2)					
6 months									
Low-protein formula	62	980 (248)	615 (156)	16.2 (4.1)					
Control formula	74	957 (172)	628 (113)	25.8 (4.6)					
9 months									
Low-protein formula	55	896 (256)	563 (161)	14.8 (4.2)					
Control formula	64	869 (242)	570 (159)	23.5 (6.5)					
12 months									
Low-protein formula	47	854 (324)	536 (203)	14.1 (5.3)					
Control formula	63	747 (217)	490 (142)	20.2 (5.9)					

The mean volume of formula consumed by infants did not differ significantly between the low protein and the control groups at any time point in any of the studies. In this context, mean energy intake from formula was comparable between the two formula groups whilst mean protein intake was systematically higher in the control formula vs. the low-protein formula groups in both studies, as per study design. The Panel notes that the mean volume intake of formula in the low-protein formula and control formula groups in both studies at 9 and 12 months of age was high.

The results of anthropometric measurements in the low-protein formula, control formula and breastfed reference groups in both studies are given in Appendices A-D. In both studies, weight gain



(in g/day) was somewhat lower in the infants consuming the low protein formula than in infants consuming the control formula for the time period 3 to 6 months and 6 to 12 months of age, but this difference only reached statistical significance for weight gain between 3 and 6 months in the Chile study. Other anthropometric measures (i.e. weight, length and head circumference at different time points in absolute values and as change from baseline) generally followed this pattern in both studies.

In the US study, both formula groups showed statistically significantly higher weight gain and higher weight and length in absolute values at different time points as compared to the breastfed reference group. In the Chile study, the low-protein formula group and the breastfed reference group did not differ significantly in weight gain, weight and length. The concentrations of serum albumin and BUN remained within the normal range in all groups during the intervention in both studies. Reported adverse events were similar in the intervention and control groups.

The Panel notes that, in both studies, no differences in growth patterns were observed between infants in the control vs. the low-protein formulae, including the time period of 3 to 6 months of age when the formula was fed almost exclusively. In the US study, growth was higher in the low-protein groups as compared to the breastfed reference group, whereas in the Chile study, infants in the low-protein formula group had a similar growth pattern to breastfed infants. The control formula used in these studies contained 0.35 g/100 kcal (US study) and 0.90 g/100 kcal (Chile study) more protein than the current minimum requirement for protein content of a FOF (1.8 g/100 kcal). In both studies, at all-time points, the difference in mean protein intake from formula between the control formula group and the low-protein formula group was 4 g/day or greater.

The Panel also notes that the studies submitted were not specifically designed to meet the regulatory definitions for either IF or FOF laid down in Regulation (EU) No 609/2013¹⁰, and that the information provided in relation to the type and amount of CF was not sufficient to calculate total energy and protein intake, nor the relative contribution of formulae and CF to total energy and protein intake. Therefore, the Panel considers that these studies do not provide, on their own, sufficient information to conclude on the safety and suitability of a FOF with a protein content of 1.6 g protein/100kcal.

3.6. Comparison between the human intervention studies provided and European dietary surveys with respect to mean energy and protein intake from formula and complementary food in the target population

In the European surveys which allowed calculation of mean energy and protein intake from both (infant and follow-on) formula and CF in formula-fed infants, the protein content of formula was between 2.1 and 2.7 g/100kcal (Table 7). The lower end is close to the protein content of the control formula used in the US study (2.15 g/100cal) and the upper end is close to the protein content of the control formula used in the Chile study (2.70g/kcal). Mean energy and protein intakes from formula were, however, lower in infants aged 4-<6 months in the European surveys than in infants at 4 months of age in the two formula groups (low protein and control) in both intervention studies (US and Chile studies). In infants aged 6-12 months, mean energy and protein intakes from formula in the European surveys were about half (or lower) than in the US and Chile studies. This suggests that the contribution of formula (vs. CF) to total protein intake in the target population (infants at the time of the introduction of complementary feeding and up to 12 months of age) may be lower in Europe than in the intervention studies provided. Therefore, the impact on total protein intake of lowering the protein content of a follow-on formula to about 1.6 g/100 kcal would also be lower in Europe. However, direct comparisons regarding total energy and protein intake and energy and protein intake from CF between the European surveys and the intervention studies provided cannot be made.

Regulation (EU) No 609/2013 of the European Parliament and of the Council of 12 June 2013 on food intended for infants and young children, food for special medical purposes, and total diet replacement for weight control and repealing Council Directive 92/52/EEC, Commission Directives 96/8/EC, 1999/21/EC, 2006/125/EC and 2006/141/EC, Directive 2009/39/EC of the European Parliament and of the Council and Commission Regulations (EC) No 41/2009 and (EC) No 953/2009, OJ L 181, 29.6.2013, p. 35–56



4. Conclusions

555 4.1. On the safety and suitability for use by infants of FOF with a protein content of at least 1.6 g/100 kcal from intact cow's milk protein otherwise complying with the requirements of relevant EU legislation

The Panel considers that the two intervention studies provided by the applicant do not provide, on their own, sufficient information to conclude on the safety and suitability of FOF with a protein content of 1.6 g protein/100kcal.

- The Panel notes, however, that:
 - a) the protein content of human milk tends to decrease with feeding time to about 1.1-1.4 g/100kcal by the end of the fourth month of lactation, remaining fairly stable thereafter,
 - b) that P5th and P2.5th of protein intake from all sources (breast milk, formula and CF) in European infants between 6 and 12 months of age are at or above the PRI for protein for that age group.
 - c) that P5th and P2.5th of protein intake from all sources (formula and CF) in European infants between 6 and 12 months of age who are not breastfed would remain at or above the PRI for protein for that age group by assuming a protein content of 1.6 g/100kcal in all FOF.
 - d) the two human intervention studies provided by the applicant did not show an adverse impact on growth resulting from the use of a formula containing about 1.6 g of protein/100kcal as compared to control formulae containing 2.15 or 2.70 g of protein/100 kcal or the breastfed reference group.

Therefore, the Panel concludes that the use of FOF with a protein content of at least 1.6 g/100 kcal from intact cow's milk protein otherwise complying with the requirements of relevant EU legislation is safe and suitable for infants living in Europe with access to complementary foods of a sufficient quality. This conclusion does not apply to IF.

4.2. On the safety and suitability for use by infants of FOF with a protein content of at least 1.6 g/100 kcal from goat's milk intact protein, soy protein isolates or protein hydrolysates otherwise complying with the requirements of relevant EU legislation

On the basis of:

- a) a previous evaluation by the Panel on the safety and suitability of goat's milk protein as a source of protein in IF and FOF (EFSA NDA Panel, 2012b), and
- b) the Panel's conclusions regarding the safety and suitability of FOF with a protein content of at least 1.6 g/100 kcal from intact cow's milk protein otherwise complying with the requirements of relevant EU legislation (section 4.1),

the Panel concludes that the use of FOF with a protein content of at least 1.6 g/100 kcal from intact goat's milk protein otherwise complying with the requirements of relevant EU legislation is safe and suitable for infants living in Europe with access to complementary foods of a sufficient quality. This conclusion does not apply to IF.

The Panel considers, however, that the safety and suitability of each FOF (and IF) manufactured from protein hydrolysates have to be established by clinical evaluation in the target population (EFSA NDA Panel, 2014). The Panel also considers that, given the higher minimum protein requirements established for FOF (and IF) manufactured from soy protein isolates (i.e. 2.25 g/100 kcal) and the lack of data available on the use of FOF from soy protein isolates in the target population, additional studies are required to establish the safety and suitability of FOF manufactured from soy protein isolates with a protein content of at least 1.6 g/100 kcal. Therefore, the Panel concludes that the safety and suitability of FOF with a protein content of at least 1.6 g/100 kcal manufactured from either

602



protein hydrolysates or soy protein isolates cannot be established with the available data. The same conclusion applies to IF.





Documentation provided to EFSA

Application for the placing on the market of a follow-on formula with a new minimum protein content of 1.61 gram of protein per 100 kcal. April 2016. Nestlé Nutrition.

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603

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696 Abbreviations

AAP American Academy of Pediatrics

AR Average requirement
CF Complementary food

CGMP Caseinoglycomacropeptide

DR Dietary records

E% Percentage of total energy intake

EC European Commission

ESPGHAN European Society for Paediatric Gastroenterology, Hepatology and Nutrition

EFSA European Food Safety Authority

EU European Union

FAO Food and Agriculture Organization of the United Nations

FOF Follow-on formula IF Infant formula

IoM Institute of Medicine

NDA Dietetic Products, Nutrition and Allergies

P2.5th Percentile 2.5 P5th Percentile 5

PRI Population Reference Intake

SCF Scientific Committee on Food

US United States

WHO World Health Organization



Appendix A – Absolute weight, weight gain and weight change in the "US study"

	n	Intervention	n	Control	n	Breast-fed	Intervention vs control	Intervention vs breast-fed	Control vs breast-fed
		mean ± SD		mean ± SD		mean ± SD	mean difference (95%CI) ^(a)	mean difference (95%CI) ^(a)	mean difference (95%CI) ^(a)
	-	Weight g	jain (g/day)	•			Weight gain (g/day)	
3-6 months									
Completers	92	20.09 ± 4.62	91	20.65 ± 5.54	109	17.43 ± 4.89	-0.67 (-2.11 to 0.77)	-	-
PP	-	-	-	-		-	-0.69 (-2.20 to 0.81)	-	-
6-12 months									
Completers	86	11.54 ± 3.04	87	12.31 ± 3.08	103	10.39 ± 3.13	-0.80 (-1.73 to 0.13)	1.04 (0.12 to 1.95)	1.84 (0.93 to 2.75)
PP	78	11.45 ± 3.00	81	12.22 ± 3.05	97	10.28 ± 3.03	-0.85 (-1.80 to 0.10)	0.96 (0.03 to 1.89)	1.81 (0.90 to 2.73)
	-	Weig	jht (l	kg)	•			Weight (g)	
3 months									
Completers	97	5.96 ± 0.70	97	5.91 ± 0.68	112	5.78 ± 0.70		-	
PP	85	5.99 ± 0.68	85	5.89 ± 0.69	105	5.79 ± 0.68		-	
6 months									
Completers	93	7.62 ± 0.84	91	7.62 ± 0.89	109	7.24 ± 0.87	-71.06 (-193.4 to 51.25)	167.86 (47.71 to 288.01) ¹	238.92 (118.28 to 359.57) ²
PP	84	7.68 ± 0.84	85	7.62 ± 0.90	103	7.27 ± 0.89	-71.08 (-199.9 to 57.76)	175.72 (49.63 to 301.81)	246.80 (121.83 to 371.77)
12 months									
Completers	87	9.86 ± 1.12	87	9.97 ± 1.21	104	9.20 ± 1.10	-231.7 (-473.2 to 9.86)	385.61 (148.70 to 622.51) ¹	617.28 (380.58 to 853.98) ²
PP	79	9.89 ± 1.12	81	9.95 ± 1.21	98	9.20 ± 1.12	-239.7 (-484.6 to 5.11)	378.73 (139.63 to 617.83)	618.48 (382.08 to 854.88)
Weight change (kg)								Weight change (kg)	
6-12 months									
Completers	86	2.20 ± 0.58	87	2.35 ± 0.59	103	1.96 ± 0.59	-0.16 (-0.34 to 0.02)	$0.21 (0.03 \text{ to } 0.38)^3$	$0.37 (0.19 \text{ to } 0.54)^2$
PP	78	2.18 ± 0.57	81	2.33 ± 0.58	97	1.94 ± 0.56	-0.17 (-0.35 to 0.01)	0.19 (0.02 to 0.37)	$0.36 (0.19 \text{ to } 0.53)^2$

⁽a) adjusted for baseline and gender; 1 p \leq 0.01; 2 p \leq 0.0001; 3 p<0.05



Appendix B – Length and head circumference in the "US study"

	n	Intervention	n	Control	n	Breast-fed	Intervention vs control	Intervention vs breast-fed	Control vs breast-fed
		mean \pm SD		mean ± SD		mean ± SD	mean difference (95%CI) ^(a)	mean difference (95%CI) ^(a)	mean difference (95%CI) ^(a)
	-	Leng	jth (c	m)	•			Length (mm)	
3 months									
Completers	97	59.54 ± 2.24	97	59.44 ± 2.06	112	59.62 ± 2.04		-	
PP	85	59.62 ± 2.20	85	59.35 ± 2.03	105	59.61 ± 2.03		-	
6 months									
Completers	94	65.61 ± 2.30	91	65.75 ± 2.34	110	65.11 ± 2.32	-1.43 (-4.51 to 1.65)	6.03 (3.02 to 9.04) ¹	7.46 (4.41 to 10.51) ²
PP	85	65.72 ± 2.35	85	65.63 ± 2.30	104	65.16 ± 2.35	-1.48 (-4.63 to 1.67)	5.79 (2.73 to 8.86) ¹	7.27 (4.19 to 10.35) ²
12 months									
Completers	87	74.52 ± 2.53	87	74.71 ± 2.77	105	73.34 ± 2.69	-2.37 (-7.24 to 2.50)	12.04 (7.28 to 16.80) ²	14.41 (9.63 to 19.19) ²
PP	79	74.58 ± 2.55	81	74.55 ± 2.74	99	73.35 ± 2.74	-2.95 (-8.08 to 2.18)	11.67 (6.68 to 16.65) ²	14.62 (9.64 to 19.61) ²
		Head circui	mfere	ence (cm)				Head circumference (mm)
3 months									
Completers	97	40.27 ± 1.24	96	40.34 ± 1.27	112	40.27 ± 1.23		-	
PP	85	40.26 ± 1.16	84	40.41 ± 1.15	105	40.25 ± 1.26		-	
6 months									
Completers	94	42.99 ± 1.30	91	43.19 ± 1.29	110	42.92 ± 1.26	-0.57 (-2.24 to 1.10)	0.89 (-0.74 to 2.52)	1.46 (-0.19 to 3.11)
PP	85	43.00 ± 1.23	85	43.18 ± 1.25	104	42.88 ± 1.26	-0.09 (-1.67 to 1.49)	1.11 (-0.43 to 2.65)	1.20 (-0.35 to 2.74)
12 months									
Completers	86	46.18 ± 1.39	87	46.47 ± 1.49	104	46.04 ± 1.39	-1.36 (-3.82 to 1.11)	1.76 (-0.64 to 4.17)	$3.12 (0.70 \text{ to } 5.54)^3$
PP	79	46.23 ± 1.32	81	46.43 ± 1.44	98	45.98 ± 1.37	-0.43 (-2.73 to 1.86)	2.34 (0.10 to 4.57) ³	$2.77 (0.54 \text{ to } 5.00)^3$

^{700 (}a) adjusted for baseline and gender; 1 p <0.001; 2 p <0.0001; 3 p< 0.05



Appendix C – Absolute weight, weight gain and weight change in the "Chile study"

	n	Intervention	n	Control	n	Breast-fed	Intervention vs control	Intervention vs breast-fed	Control vs breast-fed	
		mean ± SD		mean ± SD		mean ± SD	mean difference (95%CI) ^(a)	mean difference (95%CI) ^(a)	mean difference (95%CI) ^(a)	
	_	Weight o	jain (g/day)	-			Weight gain (g/day)		
3-6 months										
Completers	66	18.97 ± 4.19	76	20.74 ± 5.01	65	20.07 ± 5.79	-2.26 (-3.88 to -0.64) ¹	-0.72 (-2.46 to 1.01)	1.54 (-0.13 to 3.21)	
PP	55	19.17 ± 4.16	68	21.02 ± 4.88	57	20.35 ± 5.83	-	-	-	
6-12 months										
Completers	54	10.97 ± 3.05	66	12.13 ± 3.03	61	10.18 ± 3.85	-0.88 (-2.10 to 0.35)	0.77 (-0.50 to 2.05)	1.65 (0.45 to 2.85) ¹	
PP	47	11.05 ± 3.03	60	12.09 ± 3.15	54	9.99 ± 3.94	-0.76 (-2.09 to 0.58)	1.16 (-0.24 to 2.57)	1.92 (0.63 to 3.21) ¹	
	_	Wei	ght (l	kg)			Weight (g)			
6 months										
Completers	66	8.03 ± 0.67	76	8.17 ± 0.95	65	8.50 ± 1.12	-142.91 (-377.33 to 91.52)	3.75 (-245.49 to 252.99)	146.65 (-93.14 to 386.45)	
PP	55	8.09 ± 0.68	68	8.21 ± 0.97	57	8.52 ± 1.11	-161.08 (-502.24 to 180.08)	71.78 (-286.92 to 430.48)	232.85 (102.29 to 568.00)	
12 months										
Completers	54	10.08 ± 0.86	66	10.36 ± 1.10	61	10.42 ± 1.27	-315.70 (-566.93 to -64.46) ³	132.25 (-129.81 to 394.32)	447.95 (199.13 to 696.77) ²	
PP	47	10.14 ± 0.87	60	10.40 ± 1.13	54	10.43 ± 1.28	-284.55 (-655.77 to 86.68)	273.83 (-112.04 to 659.70)	558.38 (211.97 to 904.79) ¹	
	-	Weight o	chang	ge (kg)				Weight change (kg)		
6-12 months										
Completers	54	2.01 ± 0.55	66	2.20 ± 0.55	61	1.89 ± 0.71	-0.14 (0.37 to 0.08)	0.11 (-0.12 to 0.35)	$0.26 (0.04 \text{ to } 0.48)^3$	
PP	47	2.03 ± 0.54	60	2.20 ± 0.57	54	1.86 ± 0.73	-0.12 (-0.37 to 0.12)	0.19 (-0.07 to 0.44)	0.31 (0.07 to 0.54) ³	

⁽a) adjusted for baseline, gender, pre-pregnancy BMI of the mother (as continuous variable), antibiotic use, introduction of complementary food prior to 6 months of age (yes/no) and ethnicity; 1 p \leq 0.01; 2 p \leq 0.001; 3 p< 0.05



Appendix D – Length and head circumference in the "Chile study"

	n	Intervention	n	Control	n	Breast-fed	Intervention vs control	Intervention vs breast-fed	Control vs breast-fed
		mean ± SD		mean ± SD		mean ± SD	mean difference (95%CI) ^(a)	mean difference (95%CI) ^(a)	mean difference (95%CI) ^(a)
		Lengt	h (cm)					Length (mm)	
3 months									
Completers	89	59.80 ± 2.01	87	60.34 ± 2.07	76	61.20 ± 1.95	-6.21 (-12.03 to -0.39) ¹	$-14.60 (-20.71 \text{ to } -8.49)^2$	-8.39 (-14.47 to -2.30) ³
PP	65	59.84 ± 1.95	73	60.30 ± 2.01	65	61.08 ± 1.93	-6.46 (-12.94 to 0.02)	-14.05 (-20.83 to -7.27) ²	-7.59 (-14.07 to -1.11) ¹
6 months									
Completers	66	66.04 ± 1.78	76	66.54 ± 2.25	65	67.09 ± 2.25	0.08 (-5.36 to 5.53)	1.86 (-3.97 to 7.68)	1.77 (-3.74 to 7.29)
PP	55	66.20 ± 1.82	68	66.56 ± 2.14	57	67.04 ± 2.18	4.12 (-3.75 to 11.99)	8.64 (0.21 to 17.06) ¹	4.51 (-3.26 to 12.28)
12 months									
Completers	54	74.23 ± 1.99	66	74.77 ± 2.34	61	75.37 ± 2.60	-0.75 (-6.58 to 5.08)	1.58 (-4.54 to 7.70)	2.33 (-3.40 to 8.05)
PP	47	74.44 ± 1.97	60	74.81 ± 2.33	54	75.36 ± 2.64	3.19 (-5.33 to 11.71)	8.33 (-0.70 to 17.35)	5.13 (-2.89 to 13.16)
		Head circum	ferenc	ce (cm)				Head circumference (mm)	
3 months									
Completers	89	40.22 ± 1.06	87	40.50 ± 1.11	76	40.66 ± 1.19	-3.19 (-6.29 to -0.10) ¹	-4.71 (-7.96 to -1.47) ³	-1.52 (-4.75 to 1.72)
PP	65	40.29 ± 0.97	73	40.46 ± 1.07	65	40.74 ± 1.15	-2.52 (-5.92 to 0.89)	-4.97 (-8.53 to -1.40) ³	-2.45 (-5.86 to 0.96)
6 months									
Completers	66	43.10 ± 1.17	76	43.27 ± 1.24	65	43.43 ± 1.36	0.73 (-1.24 to 2.69)	0.79 (-1.28 to 2.86)	0.06 (-1.92 to 2.03)
PP	55	43.22 ± 1.05	68	43.26 ± 1.17	57	43.51 ± 1.31	1.12 (-1.72 to 3.96)	1.76 (-1.25 to 4.77)	0.64 (-2.14 to 3.42)
12 months							,	•	-
Completers	54	46.02 ± 1.32	66	46.34 ± 1.09	61	46.40 ± 1.36	-0.76 (-2.87 to 1.34)	0.90 (-1.28 to 3.08)	1.67 (-0.38 to 3.71)
PP	47	46.11 ± 1.16	60	46.35 ± 0.97	54	46.50 ± 1.32	-2.19 (-5.32 to 0.93)	0.80 (-2.46 to 4.07)	3.00 (0.11 to 5.88) ¹

⁽a) adjusted for baseline, gender, pre-pregnancy BMI of the mother (as continuous variable), antibiotic use, introduction of complementary food prior to 6 months of age (yes/no) and ethnicity; 1 p < 0.05; 2 p < 0.001; 3 p < 0.01