

Young children formula consumption and iron deficiency at 24 months in the general population: A national-level study

Anne-Sylvia Sacri, Alain Bocquet, Mariane de Montalembert, Serge Hercberg, Laurent Gouya, Béatrice Blondel, Amandine Ganon, Pascale Hebel, Catherine Vincelet, Franck Thollot, et al.

▶ To cite this version:

Anne-Sylvia Sacri, Alain Bocquet, Mariane de Montalembert, Serge Hercberg, Laurent Gouya, et al.. Young children formula consumption and iron deficiency at 24 months in the general population: A national-level study. Clinical Nutrition, 2021, 40, pp.166 - 173. 10.1016/j.clnu.2020.04.041 . hal-03493162

HAL Id: hal-03493162 https://hal.science/hal-03493162v1

Submitted on 2 Jan 2023

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers. L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



Distributed under a Creative Commons Attribution - NonCommercial 4.0 International License

Version of Record: https://www.sciencedirect.com/science/article/pii/S0261561420302156 Manuscript f1f2eda4e8b7805e58174b1366fd56ef

Young children formula consumption and iron deficiency at 24 months in the general population: a national-level study

3

4 Authors' full name and academic degrees:

- Anne-Sylvia Sacri, MD, PhD ^{a,b}; Alain Bocquet, MD ^{c,d}; Mariane de Montalembert, MD,
 PhD ^{a,b}; Serge Hercberg, MD, PhD ^{e,f}; Laurent Gouya, MD, PhD ^g; Béatrice Blondel, PhD ^a;
 Amandine Ganon, Pharm D ^h; Pascale Hebel, PhD ⁱ; Catherine Vincelet, MD ^j; Franck
 Thollot, MD ^k; Massimiliano Rallo, MD ¹; Piotr Gembara, MD ^m; Corinne Levy, MD ^{c,n,o};
 Martin Chalumeau, MD, PhD ^{a,b}
- 10

11 Authors' affiliations:

- 12 ^a Centre of Research in Epidemiology and Statistics Sorbonne Paris Cité, Obstetrical,
- 13 Perinatal and Pediatric Epidemiology Research team (EPOPé), Inserm UMR 1153; Labex
- 14 GR-Ex; Paris Descartes University, France
- ^b Department of General Pediatrics and Pediatric Infectious Diseases, Necker-Enfants malades
- 16 hospital, AP-HP; Paris Descartes University; Paris, France
- 17 ^c Association Française de Pédiatrie Ambulatoire (AFPA), Talence, France
- 18 ^d Comité de nutrition de la Société Française de Pédiatrie, Paris, France
- ^e Centre of Research in Epidemiology and Statistics Sorbonne Paris Cité, Nutritional
- 20 Epidemiology Research Team (EREN), Inserm UMR 1153, Paris Descartes University,
- 21 Sorbonne Paris Cité, France
- ^f Département de Santé Publique, AP-HP, Hôpital Avicenne, Bobigny
- ^g Centre de recherche sur l'inflammation, Inserm UMR 1149, Université Paris Diderot ; ERL
- 24 CNRS 8252, Faculté de Médecine site Bichat, Paris, France
- 25 ^h Laboratoire CERBA, Saint-Ouen l'Aumône, France

26	ⁱ Centre de Recherche pour l'Etude et l'Observation des Conditions de vie (CREDOC), Paris,
27	France
28	^j Centre régional de coordination des dépistages des cancers Ile-de-France, Lieusaint, France
29	^k Cabinet de pédiatrie, 4 Rue Parmentier, Essay Les Nancy, France
30	¹ Cabinet de pédiatrie, 3 Rue de la Béraudière, Nantes, France
31	^m Cabinet de pédiatrie, 4 Avenue Pierre Coulon, Vichy, France
32	ⁿ Association Clinique et Thérapeutique Infantile du Val-de-Marne (ACTIV), Saint-Maur des
33	Fossés, France
34	^o Université Paris Est, IMRB- GRC GEMINI, 94000 Créteil, France
35	
36	
37	Short title: Iron-fortified formula use and iron deficiency
38	
39	Corresponding author:
40	Anne-Sylvia Sacri, Inserm UMR1153, Épidémiologie Périnatale, Obstétricale et Pédiatrique
41	(ÉPOPé). Maternité de Port Royal, 53 avenue de l'Observatoire, 75014 PARIS, France.
42	Phone: +33142345580. Fax: +33143268979. E-mail: anne-sylvia.sacri@inserm.fr
43	

45 Abstract (286/300 words)

Background and aims: Iron deficiency (ID) is considered the most frequent micronutrient deficiency in industrialized countries where strategies for its primary prevention vary widely and are insufficiently evaluated. We aimed to study the effectiveness for iron status of a national iron deficiency prevention strategy based on recommendations for young-child formula (YCF) use after age 12 months, taking into consideration other sources of iron and the family's socio-economic status.

52 **Methods:** In a cross-sectional observational study conducted in primary care pediatrician 53 offices throughout France from 2016 to 2017, infants aged 24 months were consecutively 54 included for a food survey and blood sampling. Associations between YCF consumption and 55 serum ferritin (SF) level were studied by multivariable regression after adjustment on 56 sociodemographic, perinatal and dietary characteristics, notably other intakes of iron.

57 Results: Among the 561 infants analyzed, the ID prevalence was 6.6% (37/551; 95%) confidence interval [CI] 4.7-9.0). Daily iron intake excluding YCF and total daily iron intake 58 including YCF were below the 5-mg/day recommended average requirements for 63% and 59 18% of children, respectively. ID frequency was significantly decreased (or SF level was 60 independently higher) with any YCF consumption since age 10 months (odds ratio 0.15, 95%) 61 CI 0.07-0.31), current YCF consumption at age 24 months (median SF level 29 vs 21 µg/L if 62 none), prolonged YCF consumption (28 μ g/L if > 12 months vs 17 μ g/L if none), and 63 increasing daily volume of YCF consumed at age 24 months from a small volume (e.g., 29 64 μ g/L if <100 mL/day vs 21 μ g/L if none). 65

66 Conclusions: Current or past YCF use was independently associated with a better iron status
67 at age 24 months than non-use. The strategy recommending YCF use at weaning after age 12
68 months seems effective in the general population.

69 ClinicalTrials.gov identifier: NCT02484274

- **Keywords:** Iron deficiency; primary prevention; young-child formula; socioeconomic factors,
- 72 effectiveness; nationwide cohort study

73 Introduction

Iron deficiency (ID)¹ is considered the most frequent micronutrient deficiency worldwide, 74 including in industrialized countries, and is strongly suspected to be associated with adverse 75 short- and long-term neurocognitive effects when it occurs in neonates and infants [1]. In this 76 context, ID is a target of various primary preventive strategies supported by medical societies 77 and public health authorities, based on advice to parents for consumption of naturally iron-78 79 rich foods during complementary feeding, iron fortification (e.g., formula or cereals) or iron supplementation (drops) in their children [2-4]. In some countries like France, the preventive 80 strategy involves both parental education related to complementary feeding and the 81 82 recommendation for consumption of iron-fortified formulas (IFFs) at breastfeeding cessation [5]. IFFs after age 6 months include "follow-on formulas" and "young-child formulas" 83 (YCFs, sometimes inadequately called "growing-up formula") between age 6 and 10-12 84 85 months and after 10-12 months, respectively, to help to ensure the recommended average requirement (AR) of 5 mg/day at this age [5, 6]. The European Food Security and Authority 86 (EFSA) does not recommend YCF as the main source of iron after age 12 months and has 87 criticized YCF-based strategies given the lack of evidence supporting their superiority as 88 compared with balanced diet strategies [7, 8]. In some industrialized countries, balanced diets 89 90 were recently shown to be rarely achieved at this age at the population level [9, 10].

91 The efficacy of IFFs has been evaluated by eight clinical trials, seven of which found a 92 statistically significant effect of IFFs use on iron status [11-17]. The efficacy of IFFs seems 93 well established at the scale of randomized trials, but this conclusion cannot be extrapolated 94 to their effectiveness, notably after age 12 months, at the population level. Indeed, the ability 95 of IFF-based strategies to reach disadvantaged families (with low economic and/or education

¹**Abbreviations:** AR: average requirements; EFSA: European Food and Security Authority; ID: iron deficiency; IFF: iron-fortified formula; SF: serum ferritin; YCF: young children formula.

level) [18-21], who are at greatest risk of ID [2-4, 22], is poorly known and should be 96 evaluated. Moreover, to be reliable, such demonstration must take into consideration between-97 country and between-region variability in iron intake from non-dairy food. Thus, the 98 evaluation of the effectiveness of IFF-based prevention strategies requires large populations 99 with a diversified recruitment collecting sociodemographic, biological and dietary data. Only 100 three studies with a closely approaching design are available and were performed in Ireland, 101 The Netherlands, and the United Kingdom [23-25]. They all found a protective effect of YCF 102 103 on iron status at age 12 to 24 months (e.g., adjusted odds ratio [OR] for ID was 5.5 for lack of use of YCF [24]). However, these studies had limitations that prevent definitive conclusions 104 105 on the effectiveness of YCF-based strategies: selection bias (single-center recruitment [25], children requiring surgery [24]), strong attrition [23, 25], dietary survey not following EFSA 106 107 recommendations [24], and/or lack of multivariate analyses to take into consideration socio-108 economic status or non-dairy iron intake [23].

A non-biased evaluation of the independent effect of YCF at the national level is needed to produce evidence to discourage or support YCF-based strategies after age 12 months versus other strategies for ID primary prevention. Here we aimed to evaluate the effectiveness of a strategy for ID prevention based on the recommendation of YCF use after age 12 months at a national level by analyzing its association with iron status in children aged 24 months, taking into account other sources of iron and the family's socio-economic status.

115

116 Methods

117 *General methodology*

118 We conducted a cross-sectional observational study in primary care paediatric private 119 practices throughout France from January 2016 to December 2017, named the CARMA 120 study. The signed consent from at least one parent was obtained before inclusion. The study protocol was approved by local ethics and administrative authorities (CPP IDF III no. 3295)
and was registered (ClinicalTrials.gov identifier: NCT02484274). We used the Strengthening
the Reporting of Observational Studies in Epidemiology (STROBE) guidelines to report this
study (Supplemental Table 1).

125

126 Participant selection

Investigators were recruited from the directory of the French Ambulatory Pediatricians 127 Association (the national association of primary care pediatricians, n≈1600). Invitations to 128 participate were sent to all pediatricians. Among the respondents, a selected sample of about 129 130 120 pediatricians stratified on the 21 French continental regions was recruited to take in consideration the between-region variability. Investigator recruitment was secondarily 131 extended to all members of the association given difficulties in recruiting participants in this 132 133 complex study. This recruitment in primary care pediatrician offices and different regions of the country aimed at including infants from the general ambulatory population at a national 134 level. Each pediatrician was asked to include 10 consecutive children, aged 22 to 26 months, 135 usually living in France (Supplemental Figure 1), who had health insurance coverage, and 136 were not known to be affected by at least one of the following diseases that may affect iron 137 metabolism: transfusion(s) since birth, celiac disease, inflammatory bowel disease, cystic 138 fibrosis, other enteropathy (except allergy to cow's milk protein), enteral nutrition of more 139 than 15 days in the last 6 months, chronic hemolytic diseases, chronic renal failure, 140 hemophilia, hemochromatosis, malignant diseases and lead poisoning. 141

142

143 Data collection

The study phases are described in detail in Supplemental Methods. In summary, current
sociodemographic data, past and current main dietary habits and medical data were collected

by the investigator at inclusion. During the month after inclusion, a dietary survey was 146 147 performed according to the EFSA reference methodology. Parents recorded all ingesta per 24hr period over three non-consecutive days (two during the week and one during the weekend) 148 149 by using paper diaries, as in previous studies [26]. Investigators and on-line research assistants helped parents complete the survey when needed. Dietary data from these diaries 150 were double-entered, tested for consistency, and transformed by using the ANSES-Ciqual 151 152 nutrient composition table for non-infant foods [27] and the infant-specific table designed for the French Children's Food Union [28]. With this process, we obtained the ingesta and their 153 source for each individual and each nutrient. 154

After the dietary survey, a fasting blood sample was taken; serum ferritin (SF) and Creactive protein (CRP) were measured centrally in a unique laboratory after refrigerated transport of blood samples at +2 to +8°C. SF was measured by immunochemiluminescence (COBAS -Roche Diagnostics- COBAS 6000 E601, measurement range: 0.5-2000 μ g/L, intraand inter-series coefficients of variation <1.1% and <5.7%, respectively) and CRP was measured by immunoturbidimetric assay.

In case of fever or any other condition that could alter the ingesta or biological iron status (e.g., bronchiolitis, gastroenteritis), the dietary survey and blood sampling were postponed to 15 days after discontinuation of symptoms. Children for whom no blood sample was taken, whose parents did not complete the dietary survey, or with a CRP level ≥ 10 mg/L were excluded from the statistical analyses.

166

167 *Statistical analyses*

We first described the general characteristics of the studied and analyzed population and compared them to those of the 2016 French National Perinatal Survey, a national representative sample of births (n=13 148) [29]. Second, we characterized iron intake

specifically and its distribution considering all sources from the whole diet including YCF or 171 excluding YCF. Third, we described the distribution of SF and used multivariable linear 172 regression models (after logarithmic transformation of SF [30]) to study the crude then 173 174 adjusted association of SF with past dietary habits (including declared YCF consumption) and current ingesta (including declared and measured YCF consumption) and perinatal and 175 sociodemographic characteristics. The specific sources of iron evaluated in the analyses are 176 listed as dietary characteristics in Table 1. YCF consumption was defined in several ways: 177 "any consumption after 10 months" and "current consumption at age 24 months" (two binary 178 [no/yes] declarative variables), "daily volume consumed" in five categories (measured 179 180 variable in the dietary record) to evaluate a "dose effect", "consumption duration" (declarative variable) used as continuous variable, then in three categories to evaluate a "time effect". We 181 evaluated the duration of consumption of any IFF since age 6 months by a continuous 182 183 variable corresponding to the sum of consumption duration for both follow-on formula and YCF. This variable represented the cumulative iron intake from dairy food at about age 6 184 months. The statistical handling of other variables is detailed in Supplemental Methods. 185 Finally, SF was dichotomized with a $12-\mu g/l$ threshold to define ID, as suggested [2, 4, 31]. 186 The prevalence of ID and its crude associations with potential determinants were studied by 187 univariate analysis. No multivariable analysis was used given the low number of ID cases. 188

No multi-level model was used in these analyses despite recruitment in different practices because the variance of a multilevel model with random intercept including only the center of recruitment was low, 7%. For all continuous variables, the lack of deviance to linearity was searched and a fractional polynomial of degree 2 at most was used if appropriate. Several sensitivity analyses were performed by repeating the univariate analyses with a $10-\mu g/l$ threshold of SF to define ID [2, 4, 32], repeating regressions analyses without sociodemographic characteristics or studying specific associations of perinatal and sociodemographic variables with iron status. Missing data (≤2%) were not imputed. The
number of participants required (n=1000) was calculated with a very conservative hypothesis
(Supplemental Methods); the recruitment was stopped prematurely for financial reasons.
The analyses involved use of Stata/SE v13.1 (StataCorp, College Station, TX, USA).

200

201 **Results**

202 *Participants*

A total of 830 young children were included by 58 pediatricians from 16 French regions; 15 were excluded because of non-eligibility criteria, 212 (26%) because of no blood sample, 8 (1%) because of lack of dietary survey, 31 (5%) because of CRP level \geq 10 mg/L, and 3 (0.5%) because of missing data for SF level (**Supplemental Figures 1 and 2**), leading to an analyzed population of 561 infants. The analyzed population was less disadvantaged than the 2016 French National Perinatal Survey population but did not differ from the non-analyzed population (**Supplemental Table 2**).

210 In the analyzed population, the mean age of children was 24 months (SD 0.6), 49% were girls, 3.6% had cow's milk protein allergy, and 10% had received iron supplementation 211 according to parents. YCF was currently used in 73% of the population (408/561, 95%) 212 confidence interval [CI] 69-76) and cow's milk in 33% (186/561, 95% CI 29-37). The mean 213 and median total iron intake for the population was 8.2 mg/day (95% CI 7.8-8.6) and 7.6 214 mg/day (interquartile range [IQR] 5.6-9.8). The mean iron intake for the population from all 215 food excluding YCF and from YCF only was 4.7 mg/day (95% CI 4.6-4.9) and 3.5 mg/day 216 (95% CI 3.1-3.8). The median iron intake for the population from all food excluding YCF and 217 from YCF only was 4.3 mg/day (IQR 3.4-5.7) and 3.1 mg/day (IQR 0-5.2). Daily iron intake 218 219 for the population from all food excluding YCF and total daily iron intake including YCF was below the 5 mg/day AR for 63% (95% CI 59-67) and 18% (95% CI 15-22) of children
(Figure 1).

Median SF was 26 μ g/L (IQR: 20-37, mean: 31 [SD 21]) (Supplemental Figure 3). After dichotomization of SF at thresholds <12 and <10 μ g/L, ID prevalence was 6.6% (37/561, 95% CI 4.7-9.0) and 3.9% (22/561, 95% CI 2.5-5.9) (Table 1 and Supplemental Table 3).

- 226
- 227 Dietary factors associated with iron status

228 On univariate analysis, SF was significantly (p <0.05) lower with cow's milk use beginning 229 before age 24 months (median 23 vs 28 μ g/L if not) or its current use at 24 months (23 vs 230 28 μ g/L if not) and with increased iron intake from all food excluding YCF (β coefficient =

-0.03, 95% CI -0.05-[-0.01]) (Table 1). We found no significant association between SF and

232 exclusive breastfeeding beyond 6 months or cow's milk consumption duration (Table 1). SF was significantly higher with follow-on formula use between age 6 and 12 months (27 vs 233 234 23 µg/l if not), YCF use since age 10 months (27 vs 17 µg/L if not) or current use at 24 months (29 vs 21 µg/L if not), daily volume of YCF consumed (29 µg/L if <100 mL/day vs 235 21 μ g/L if none), increasing YCF consumption duration (28 if > 12 months vs 17 μ g/L with 236 no YCF), and increasing IFFs consumption duration (β coefficient = 0.05, 95% CI : 0.03-237 0.07; i.e., 28 if > 12 months vs 17 μ g/L if no IFF, this last result is not shown in Table 1) 238 (**Table 1**). For the 58 children with a high daily volume of YCF consumed (≥450 mL; i.e., the 239 90th percentile), we found no significant increase in mean SF level (34 vs 31 µmol/L with 240 daily YCF volume <450 mL, p=0.23) or median SF level [90th percentile] (26 [51] vs 27 241 µmol/L [58] with daily YCF <450 mL). 242

In several multivariable linear models adjusting for co-variables of interest (Table 2)
and after checking for no significant collinearity between YCF and cow's milk consumption,

SF was positively associated with YCF consumption, whether described by a binary variable (any use after age 10 months), volume consumed or duration of consumption. SF was also associated with duration of IFFs consumption since age 6 months (**Table 2**). In all multivariable models, unfavorable iron status was associated with a medical history of prematurity, an elevated rank among siblings and disadvantaged social situation: low father's education level (**Table 2**, **Supplemental Table 6**) as in univariate analyses (**Supplemental Tables 4 and 5**).

After dichotomizing SF at the 12- μ g/L threshold, ID was significantly associated (p <0.05) with the same markers as SF used as a continuous variable (**Table 1**) except iron intake from all food excluding YCF. For example, ID was significantly decreased with YCF use since age 10 months (25% vs 5% with YCF use, OR 0.15, 95% CI 0.07-0.31).

Sensitivity analyses at the 10- μ g/L threshold found a statistically significant association between ID (p <0.05) and the same variables as with the 12- μ g/L threshold, except for infant formula use, which was significant at the 10- μ g/L threshold only (**Supplemental Table 3**). Similar results were found in the sensitivity analyses for the multivariate models without adjustment for sociodemographic variables comparatively to those adjusted for sociodemographic variables (**Supplemental Table 6**).

262

263 **Discussion**

264 *Main results*

With a simultaneous collection of medical, biological and dietary data at the national scale, a design rarely achieved in the literature [23, 24], we found a protective role of YCF use on ID in children at age two years in the general population. This protective role was strong with, for example, a reduction of one-fifth of ID with YCF use since age 10 months (OR 0.15, 95% CI 0.07-0.31). This protective role was consistent according to the different definitions of

YCF consumption (amount, duration), measured either retrospectively by asking parents 270 about the past consumption or prospectively by quantifying current consumption with a 271 dietary survey. This protective role is consistent with results from clinical trials [11-17] and 272 273 previous observational studies [23-25, 33] and was independent of potential confounders such as disadvantaged social situation, non-dairy iron intake and iron supplementation, which were 274 not always taken into account in previous studies [23, 24]. This protective role is easy to 275 conceive theoretically because iron given to young children in any form is supposed to 276 277 increase their iron stores in the absence of concurrent loss or illness, given the almost closedcircuit metabolism of iron [4, 31, 34, 35]. 278

279 Our study provides information on how YCF-based strategies could help improve iron status in the general population. First, the mean total iron intake was 8.2 mg/day, higher than 280 the 5 mg/day AR recommended in Europe by the EFSA, in 82% of children [6]. The mean 281 282 intake from all sources excluding YCF was 4.7 mg/day and from YCF only was 3.5 mg/day for the population analyzed. Added to the EFSA recommended strategy based on naturally 283 iron-rich food, YCF use allowed for a reduction from 63% to 18% of children with below the 284 5-mg/day AR. Thus, YCF use added to the rest of the diet allowed 45% of children to reach 285 the daily AR. Furthermore, the YCF impact on iron status was detectable even for a small 286 daily volume consumed, but the increase >500 mL/day was not associated with a significant 287 increase in SF. Results from several studies support that a small volume of YCF (100-300 288 mL/day) allows for reaching the recommended AR [36-38]. Iron bioavailability mechanisms 289 are complex, and the amount of iron ingested is not equal to the amount of iron absorbed. 290 This absorption depends notably on the origin of iron (haem or not), the type of other 291 concomitant ingested food and the period of life, notably acceleration of growth [2-4, 31, 39]. 292 293 Thus, even a small amount of iron with IFFs could improve iron status [40, 41].

Second, another finding helps in understanding how YCF use could help improve iron 294 status: the association observed between increased SF and consumption duration of IFFs since 295 age 6 months (i.e., follow-on formula, then YCF). This "time effect" implying the duration of 296 297 exposure to IFFs suggests a role of long-term, "cumulative", consumption of IFFs in iron status at age 24 months. This finding, as the previous one about a non-linear effect of YCF 298 volume consumed on iron status, can probably be explained in part by the role of hepcidin in 299 300 regulating iron absorption and the prevention of iron overload if iron reserves are already high. Indeed, this peptide synthetized by hepatocytes participates in regulating iron absorption 301 by enterocytes: when iron reserves are high, hepcidin synthesis increases and induces the 302 303 degradation of ferroportin on enterocytes and macrophage membranes, thus limiting iron release in blood circulation [35, 39, 42], whereas when iron reserves are low, hepcidin 304 synthesis is downregulated, thereby increasing iron bioavailability. The potential risk of iron 305 306 excess with fortification use can be raised. Use of IFFs has been found associated with iron intake above the tolerable upper intake level but not excess iron stores based on SF 307 308 distribution in infants [38, 43]. In our analyzed population, we found no significant difference 309 in SF distribution with high daily volume consumed. This study suggests that small amounts of iron ingested regularly over time could be more beneficial to prevent ID than high amounts 310 over a short period without leading to iron excess stores. 311

In this study, we found other determinants of iron status at 24 months usually identified in the literature: medical history of prematurity and factors linked to deprived situation (low father's education level) or family situation (high rank in siblings). The independent association found between iron status and a history of prematurity, remaining even after taking into account iron supplementation, suggests that this sub-population at high risk of ID should benefit from more specific prevention and monitoring of iron status, as previously found [44, 45]. "Exclusive" breastfeeding beyond 6 months is a classical risk factor for ID [46], but it was not independently associated with iron status in our study, probably because assessment of iron status occurred late, at 24 months, and many other nutritional factors may have intervened between 7 and 24 months.

322

323 *Strengths and limitations*

Our study has several strengths. The measurements of iron biomarkers were standardized and centralized as recommended by the World Health Organization [47], as was the evaluation of nutritional intake with a dietary survey performed on three non-consecutive days to improve the accuracy of intra-individual variability, as recommended by the EFSA [26]. Biological and dietary data were measured independently, with blinding. A specific strength of this study is the consideration of all sources of iron besides YCF in multivariate analyses in contrast to some other national studies [23, 24].

There were several limitations. First, the analyzed population presented a selection 331 bias as shown by less frequent disadvantaged situations than in all births. This bias can be 332 explained mainly by recruitment by pediatricians and the significant attrition due notably to 333 failure to obtain a blood sample (32% of the studied population). Indeed, only 20% of French 334 children at age 24 months are followed by a pediatrician [48] and those who accepted to 335 participate in this complex study were probably familiar with clinical research. The study 336 possibly selected families interested in the topic of nutrition with good knowledge and 337 practices regarding ID prevention. This selection bias probably led to underestimating the ID 338 prevalence, which was regardless close to the most recent national estimations [49]. 339 Furthermore, this potential bias could not have altered the associations between iron status 340 and YCF consumption in the analyzed children. Second, the recruitment was stopped before 341 the calculated sample size was achieved, and the attrition may have led to loss of statistical 342 power. However, the sample size was calculated on the basis of a very conservative 343

hypotheses on expected associations that were not confirmed by the study (associations were 344 much stronger than expected). To manage the lack of power, the main analyses were finally 345 based on ID defined by SF level as a continuous variable. Finally, as in other studies, some 346 inaccuracies may have occurred during the declarative food intake data collected by the 347 pediatrician and regarding the quantities assessed by the parents during the survey. However, 348 they may have been limited by the checks of a well-trained dietician as detailed in 349 Supplemental Methods. Besides, these inaccuracies occurred with blinding to iron status, so 350 they may have led to only a non-differential bias and a loss of statistical power. 351

352

353 *Implications*

In this first national-level study taking into account non-dairy iron intake and iron 354 supplementation and the family's socio-economic status, YCF consumption allowed for 355 356 reaching the required iron AR in 45% of children who were not reaching it otherwise by dietary intake. YCF use was strongly and independently associated with better iron status than 357 non-use. The biological benefit of YCF was obtained even for a limited daily amount, about 358 200 mL/day, especially when implemented over a long period. These findings support 359 national strategies for ID prevention based on the recommendation for YCF consumption 360 after age 12 months [7, 8]. YCF consumption seems a simple way to supply young children 361 with the recommended iron intake and reach the recommended AR added to the rest of the 362 diet in practice, according to the family's and country's culture. Indeed, a well-balanced diet 363 by solid food is possibly difficult to achieve at this age [9, 10]. These findings should trigger 364 measures to enhance YCF accessibility to at-risk families (i.e., those in disadvantaged 365 situations). 366

368 Acknowledgments

The authors thank URC-CIC Paris Descartes Necker Cochin (project manager: Nelly Briand and clinical research associates: Camille Boucheny, Charlène Ambvouli and Emilie Le Roux) for the implementation and monitoring of the study. The authors thank ACTIV network clinical research associates: Claire Prieur, Deborah Kern. They thank the participant families and the pediatricians who recruited them:

374 Jean-Marie André, Pierre Bakhache, Sophie Barrois, Bénédicte Blanc, Marie-Emmanuelle Bouloumié, Fabienne Cahn-Sellem, Marie-Hélène Cavert, Anne Cheve, Christian Copin, 375 Muriel Cornic, François Corrard, Armelle Courboin, Véronique Dagrenat, Caroline De 376 Backer, Pascal Dechelette, Jean-François Delobbe, Michèle Dudreuilh, Corinne Dumas, 377 Christine Duvant-Fay, Annie Elbez, Chantal Gayot, Nathalie Gelbert, Patricia Giraud-378 Escoffier, Geneviève Granat, Cécile Guiheneuf, Frédéric Hassid, Kai Kassmann, Sylvie 379 380 Labarge, Sophie Langlais, Hélène Le Scornet, Dominique Lemarié, Claudia Litzler, Christine Magendie, Delphine Menet, Bénédicte Nold, Jean- Claude Oilleau, France Paratte, Brigitte 381 Pinçant, Sophie Ravilly, Olivier Romain, Bechara Saade, Catherine Salinier, Claire Sarreau, 382 Ruth-Caroline Schweitzer, Anne Sengier-Njinou, Philippe Simon, David Somerville, Joachim 383 Stoller, Marie-Pierre Streicher, Catherine Tuberg-Romain, Jean-Christophe Vigreux, Pascale 384 385 Vurpillat-Almanza, Andreas Werner, Alain Wollner, Morched Zouari.

386

387 Sources of funding:

This work was supported by Paris Diderot University - Sorbonne Paris Cité for a doctoral
grant for Dr. AS Sacri, Secteur Français des Aliments de l'Enfance, and the French Ministry
of Health ["DGOS PHRC régional 2014 no. AOR14053"]. The sponsor was Assistance
Publique – Hôpitaux de Paris (Délégation à la Recherche Clinique et à l'Innovation [DRCI]).

393 Conflict of interest statement:

394 The funders had no role in the study design, data collection and analysis, decision to publish,

395 or preparation of the manuscript. The authors have no patents, products in development or

396 marketed products to declare.

398 **References**

Lozoff B, Jimenez E, Wolf AW. Long-term developmental outcome of infants with
 iron deficiency. N Engl J Med 1991;325: 687-694.

Baker RD, Greer FR. Diagnosis and prevention of iron deficiency and iron-deficiency
anemia in infants and young children (0-3 years of age). Pediatrics 2010;126: 1040-1050.

403 3. EFSA. European Food Security Authority. Panel on Dietetic Products, Nutrition and

404 Allergies (NDA). Scientific Opinion on nutrient requirements and dietary intakes of infants

and young children in the European Union. European Food Security Authority (EFSA)

406 Journal [serial online] 2013;11: 3408-3511.

407 4. Domellof M, Braegger C, Campoy C, Colomb V, Decsi T, Fewtrell M, et al. Iron
408 requirements of infants and toddlers. J Pediatr Gastroenterol Nutr 2014;58: 119-129.

409 5. Ghisolfi J, Vidailhet M, Fantino M, Bocquet A, Bresson JL, Briend A, et al. Lait de

410 vache ou lait de croissance : quel lait recommander pour les enfants en bas age (1-3 ans) ?

411 [French]. Arch Pediatr 2011;18: 355-358.

412 6. EFSA. Dietary reference values for nutrients: Summary report. EFSA supporting

413 publication 2017: published online Dec 8. DOI: 10.2903/sp.efsa.2017.e15121. (Available

414 from: https://www.efsa.europa.eu/en/supporting/pub/e15121.)

415 7. 'Growing-up' formula: No additional value to a balanced diet, says EFSA. Parma:

416 European Food Safety Authority (EFSA), 2013. (Accessed November 2018, at:

417 https://www.efsa.europa.eu/en/press/news/131025.)

418 8. EFSA Panel on Dietetic Products, Nutrition and Allergies (NDA). Scientific Opinion

419 on nutrient requirements and dietary intakes of infants and young children in the European

420 Union. EFSA Journal 2013: published online Oct 25. DOI: 10.2903/j.efsa.2013.3408.

421 (Available from: https://www.efsa.europa.eu/en/efsajournal/pub/3408.)

422 9. Chouraqui JP, Tavoularis G, Simeoni U, Ferry C, Turck D. Food, water, energy, and
423 macronutrient intake of non-breastfed infants and young children (0-3 years). Eur J Nutr
424 2019.

425 10. Grimes CA, Szymlek-Gay EA, Campbell KJ, Nicklas TA. Food Sources of Total
426 Energy and Nutrients among U.S. Infants and Toddlers: National Health and Nutrition
427 Examination Survey 2005-2012. Nutrients 2015;7: 6797-6836.

428 11. Fuchs GJ, Farris RP, DeWier M, Hutchinson SW, Warrier R, Doucet H, et al. Iron
429 status and intake of older infants fed formula vs cow milk with cereal. Am J Clin Nutr
430 1993;58: 343-348.

431 12. Moffatt ME, Longstaffe S, Besant J, Dureski C. Prevention of iron deficiency and
432 psychomotor decline in high-risk infants through use of iron-fortified infant formula: a
433 randomized clinical trial. J Pediatr 1994;125: 527-534.

13. Daly A, MacDonald A, Aukett A, Williams J, Wolf A, Davidson J, et al. Prevention of
anaemia in inner city toddlers by an iron supplemented cows' milk formula. Arch Dis Child
1996;75: 9-16.

437 14. Gill DG, Vincent S, Segal DS. Follow-on formula in the prevention of iron deficiency:
438 a multicentre study. Acta Paediatr 1997;86: 683-689.

Morley R, Abbott R, Fairweather-Tait S, MacFadyen U, Stephenson T, Lucas A. Iron
fortified follow on formula from 9 to 18 months improves iron status but not development or
growth: a randomised trial. Arch Dis Child 1999;81: 247-252.

16. Maldonado Lozano J, Baro L, Ramirez-Tortosa MC, Gil F, Linde J, Lopez-Huertas E,

et al. [Intake of an iron-supplemented milk formula as a preventive measure to avoid low iron

444 status in 1-3 year-olds]. An Pediatr (Barc) 2007;66: 591-596.

445 17. Akkermans MD, Eussen SR, van der Horst-Graat JM, van Elburg RM, van Goudoever

446 JB, Brus F. A micronutrient-fortified young-child formula improves the iron and vitamin D

- status of healthy young European children: a randomized, double-blind controlled trial. Am J
 Clin Nutr 2017;105: 391-399.
- Attree P. A critical analysis of UK public health policies in relation to diet and
 nutrition in low-income households. Matern Child Nutr 2006;2: 67-78.
- 451 19. Barker D, Barker M, Fleming T, Lampl M. Developmental biology: Support mothers
- to secure future public health. Nature 2013;504: 209-211.
- 453 20. Fleurbaey M, Luchini S, Schokkaert E, Van de Voorde C. Évaluation des politiques de
- 454 santé : pour une prise en compte équitable des intérêts des populations. Economie et
- 455 Statistique / Economics and Statistics Insee 2012;455-456.
- 456 21. Méthodes d'évaluation des politiques publiques actes du séminaire [séminaire
- 457 Dress]. Paris, France: Ministère des Solidarités et de la Santé, 2011. (Accessed May 2019, at:
- 458 http://drees.solidarites-sante.gouv.fr/etudes-et-statistiques/publications/recueils-ouvrages-et-
- 459 rapports/ouvrages-thematiques/article/methodes-d-evaluation-des-politiques-publiques-actes-460 du-seminaire.)
- 461 22. Sacri AS, Hercberg S, Gouya L, Levy C, Bocquet A, Blondel B, et al. Very low
 462 prevalence of iron deficiency among young French children: A national cross-sectional
 463 hospital-based survey. Matern Child Nutr 2017 May 03. doi: 10.1111/mcn.12460. [Epub
 464 ahead of print].
- 23. Sidnell A, Pigat S, Gibson S, O'Connor R, Connolly A, Sterecka S, et al. Nutrient
 intakes and iron and vitamin D status differ depending on main milk consumed by UK
 children aged 12-18 months secondary analysis from the Diet and Nutrition Survey of
 Infants and Young Children. J Nutr Sci 2016;5: e32.
- 469 24. Uijterschout L, Vloemans J, Vos R, Teunisse PP, Hudig C, Bubbers S, et al.
- 470 Prevalence and risk factors of iron deficiency in healthy young children in the southwestern
- 471 Netherlands. J Pediatr Gastroenterol Nutr 2014;58: 193-198.

472 25. McCarthy EK, Ni Chaoimh C, Hourihane JO, Kenny LC, Irvine AD, Murray DM, et
473 al. Iron intakes and status of 2-year-old children in the Cork BASELINE Birth Cohort Study.

474 Matern Child Nutr 2017;13.

475 26. EFSA. General principles for the collection of national food consumption data in the
476 view of a pan-European dietary survey. EFSA Journal 2009;7.

477 27. Table de composition nutritionnelle des aliments Ciqual [Internet database]. Agence

478 nationale de sécurité sanitaire de l'alimentation, de l'environnement et du travail (Anses),

479 2017. (Accessed January 2018, at: https://ciqual.anses.fr/.)

480 28. Tavoularis G. Nutri-Bebe 2013 Study Part 1. Presentation and methodological

481 considerations. Arch Pediatr 2015;22: 10s11-16.

482 29. Blondel B, Coulm B, Bonnet C, Goffinet F, Le Ray C, National Coordination Group

483 of the National Perinatal S. Trends in perinatal health in metropolitan France from 1995 to

2016: Results from the French National Perinatal Surveys. J Gynecol Obstet Hum Reprod
2017;46: 701-713.

486 30. Bland JM, Altman DG. Transformations, means, and confidence intervals. BMJ
487 1996;312: 1079.

488 31. Zimmermann MB, Hurrell RF. Nutritional iron deficiency. Lancet 2007;370: 511-520.

489 32. Dallman PR, Siimes MA, Stekel A. Iron deficiency in infancy and childhood. Am J
490 Clin Nutr 1980;33: 86-118.

491 33. Vincelet C, Foucault C. [Measuring iron levels relative to the type of milk consumed

within a population of 16 to 18 month old French infants]. Sante Publique 2005;17: 339-346.

493 34. de Montalembert M, Bresson JL, Brouzes C, Ruemmele FM, Puy H, Beaumont C.

494 [Diagnosis of hypochromic microcytic anemia in children]. Arch Pediatr 2012;19: 295-304.

495 35. Loreal O, Bardou-Jacquet E, Jouanolle AM, Gandon Y, Deugnier Y, Brissot P, et al.

496 [Iron metabolism and tools for the iron status assessment]. Rev Med Interne 2012;33 Suppl 1:497 \$3-9.

Walton J, Flynn A. Nutritional adequacy of diets containing growing up milks or
unfortified cow's milk in Irish children (aged 12-24 months). Food Nutr Res 2013;57.

500 37. Ghisolfi J, Fantino M, Turck D, de Courcy GP, Vidailhet M. Nutrient intakes of

501 children aged 1-2 years as a function of milk consumption, cows' milk or growing-up milk.

502 Public Health Nutr 2013;16: 524-534.

503 38. Eussen SR, Pean J, Olivier L, Delaere F, Lluch A. Theoretical Impact of Replacing

504 Whole Cow's Milk by Young-Child Formula on Nutrient Intakes of UK Young Children:

Results of a Simulation Study. Ann Nutr Metab 2015;67: 247-256.

506 39. de Montalembert M, Bresson J, Brouzes C, Ruemmele F, Puy H, Beaumont C.

507 Exploration d'une anemie microcytaire chez l'enfant. Arch Pediatr 2012;19: 295-304.

Fomon SJ, Ziegler EE, Serfass RE, Nelson SE, Frantz JA. Erythrocyte incorporation
of iron is similar in infants fed formulas fortified with 12 mg/L or 8 mg/L of iron. J Nutr
1997;127: 83-88.

511 41. Walter T, Pino P, Pizarro F, Lozoff B. Prevention of iron-deficiency anemia:

comparison of high- and low-iron formulas in term healthy infants after six months of life. J
Pediatr 1998;132: 635-640.

42. Ganz T. Hepcidin and iron regulation, 10 years later. Blood 2011;117: 4425-4433.

43. van der Merwe LF, Eussen SR. Iron status of young children in Europe. Am J Clin
Nutr 2017;106: 1663S-1671S.

517 44. McCarthy EK, Dempsey EM, Kiely ME. Iron supplementation in preterm and low-

518 birth-weight infants: a systematic review of intervention studies. Nutr Rev 2019;77: 865-877.

45. Akkermans MD, Uijterschout L, Abbink M, Vos P, Rovekamp-Abels L, Boersma B,
et al. Predictive factors of iron depletion in late preterm infants at the postnatal age of 6
weeks. Eur J Clin Nutr 2016;70: 941-946.

46. Atkins LA, McNaughton SA, Campbell KJ, Szymlek-Gay EA. Iron intakes of
Australian infants and toddlers: findings from the Melbourne Infant Feeding, Activity and
Nutrition Trial (InFANT) Program. Br J Nutr 2016;115: 285-293.

525 47. Iron deficiency anaemia. Assessment, prevention and control. A guide for programme

526 managers. Geneva: World Health Organization (WHO), 2001. (Accessed December 2019, at:

527 http://www.who.int/nutrition/publications/en/ida_assessment_prevention_control.pdf.)

528 48. Bocquet A, Chalumeau M, Bollotte D, Escano G, Langue J, Virey B. [Comparison of

529 prescriptions by pediatricians and general practitioners: a population-based study in Franche-

530 Comte from the database of Regional Health Insurance Fund]. Arch Pediatr 2005;12: 1688-

531 1696.

532 49. Sacri AS, Hercberg S, Gouya L, Levy C, Bocquet A, Blondel B, et al. Very low

533 prevalence of iron deficiency among young French children: A national cross-sectional

based survey. Matern Child Nutr 2018;14.

535 50. Cochrane Handbook for Systematic Reviews of Interventions. London: The Cochrane

536 Collaboration, 2011. (Accessed June 2018, at: https://www.cochrane.org/news/new-

537 cochrane-handbook-systematic-reviews-interventions.)

538 51. Chouraqui JP, Turck D, Tavoularis G, Ferry C, Dupont C. The Role of Young Child

539 Formula in Ensuring a Balanced Diet in Young Children (1-3 Years Old). Nutrients 2019;11.

540 52. World Economic and Financial Surveys: World Economic Outlook Database.

541 Washington, D.C.: International Monetary Fund Research Dept., 2017. (Accessed September

542 2019, at: http://www.imf.org/external/pubs/ft/weo/2016/02/weodata/index.aspx.)



Figure 1. Distribution of mean daily iron intake (mg/day) according to deciles and quartiles
of iron intake stratified by iron sources (n=561): from all food (red line), from all food
excluding young-child formula (YCF, blue line), and YCF only (green line).

548 Deciles and quartiles were calculated separately by iron sources. At a consequence, for each
549 decile or quantile, mean daily iron intake from all food is not equal to the addition of mean
550 daily iron intake from all food excluding YCF + YCF only.

551 In children who did not reached the required iron average requirements recommended from 552 all food without YCF, 45% reached this threshold if iron intake from YCF was considered.

553 Horizontal orange line: threshold of iron intake of 5 mg/day corresponding to the average

requirements recommended by the European Food Security and Authority (EFSA) at age 24

- 555 months [6]
- 556
- 557

Table 1. Dietary characteristics of children and univariate associations with serum ferritin 558 level and iron deficiency (ID) at a threshold of 12 µg/L (by milk according to the age of 559

consumption) 560

561

Characteristics	n (%) ^a	Serum	ferritin level (µ	ID, serum ferritin level < 12 μg/L				
		Median ^b	IQR ^b	p°	ID ^a (%)	No ID ^a (%)	OR ^d	95% CI °
	n = 561				n = 37	n = 524		
"Exclusive" breastfeedi	ng > 6 months							
No	484 (86)	27	20-37	-	5	95	-	-
Yes	77 (14)	26	17-35	0.07	16	84	3.39	1.62-7.07
Infant formula: consump	otion 0 to 6 month	IS						
No	128 (23)	26	19-35	-	10	90	-	
Yes	433 (77)	27	20-37	0.11	6	94	0.52	0.26-1.05
Late food diversification	> 6 months							
No	546 (98)	26	20-37	-	7	93	-	-
Yes	10(2)	30	23-38	0.83	10	90	1.57	0.19-12.77
Follow-on formula: cons	umption 6 to 10-1	2 months						
No	68 (12)	23	15-35	-	21	79	-	-
Yes	492 (88)	27	20-37	< 0.01	5	95	0.19	0.09-0.39
Follow-on formula: cons	umption duration	n (months) ^f						
	5.7 (1.5)	0.01	-0.03-0.04	0.65	5.8 (1.7)	5.7 (1.4)	1.05	0.80-1.37
YCF: consumption after	10 months							
No	48 (9)	17	11-30	-	25	75	-	-
Yes	512 (91)	27	21-37	< 0.001	5	95	0.15	0.07-0.31
YCF: current consumpti	ion at 24 months							
No	153 (27)	21	14-29	-	18	82	-	-
Yes	408 (73)	29	22-40	<0.001	2	98	0.10	0.05-0.22
Current YCF: daily volu	ime consumed	21	14.20		10	02		
No consumption	150 (27)	21	14-30	-	18	82	-	-
< 100 mL / day	1/5 (31)	29	22-43	<0.001	1	99	0.05	0.01-0.23
100-300 mL/ day	103 (18)	28	21-35	<0.001	/	93	0.33	0.14-0.80
300-500 mL / day	101 (18)	30	21-40	<0.001	1	99	0.05	0.01-0.34
\geq 500 mL / day	32(6)	27	22-42	<0.001	0	100	0.14	0.01-0.91
YCF: consumption dura	11.2(2.0)	0.04	0.02.0.05	<0.001	95(47)	114(20)	0.82	0.74.0.00
VCE: consumption dura	11.5 (5.0)	0.04	0.02-0.03	N0.001	8.3 (4.7)	11.4 (2.9)	0.82	0.74-0.90
No consumption	48 (0)	17	11.30		27	73		
< 6 months	61(11)	21	14-34	0.09	15	85	0.47	0.18-1.21
≥ 0 months > 6 and ≤ 12 months	235(44)	21	21.36	<0.09	3	07	0.47	0.13-1.21
> 12 months	197 (36)	28	21-30	<0.001	3	97	0.08	0.03-0.22
IFFs: consumption (since	e age 6 months) di	uration (month	s) ^f	40.001	5	71	0.00	0.05 0.24
III I S. Consumption (since	15.3(5.3)	0.05	0.03-0.07	<0.001	8.9 (7.3)	15.8 (4.8)	0.85	0.81-0.89
Cow's milk: consumption	n beginning befor	e 24 months	0102 0107	.01001	012 (112)	1010 (110)	0.00	0101 0109
No	356 (64)	28	21-40	-	3	97	-	-
Yes	201 (36)	23	15-34	< 0.001	13	87	5.98	2.75-13.00
Cow's milk: current con	sumption at 24 m	onths						
No	375 (67)	28	21-39	-	3	97	-	-
Yes	186 (33)	23	16-35	< 0.001	14	86	5.38	2.59-11.15
Cow's milk: consumption	n duration (mont	hs) ^f						
	7.4 (4.4)	-0.01	-0.04-0.01	0.15	9.4 (4.7)	7.1 (4.3)	1.13	1.02-1.25
Dietary iron intake from	all food excludin	g YCF f (mg/da	y)					
	4.7 (2.2)	-0.03	-0.05-(-0.01)	0.01	5.2 (2.8)	4.7 (2.2)	1.10	0.96-1.25
Iron supplementation sin	ce birth ^g							
No	503 (90)	26	20-36	-	7	93	-	-
Yes	57 (10)	29	22-41	0.17	4	96	0.49	0.11-2.08

^a or mean (SD) if the "characteristics" variable is continuous

562 563 564 565 566 567 568 569 570 571 572 573 573 ^b or β coefficient of a simple linear regression after logarithmic transformation (instead of medians) and 95% confidence interval (CI, instead of interquartile range [IQR]) if the « characteristics » variable is continuous

^c Degree of significance of the Student test, Kruskal-Wallis test or of the β coefficient of a simple linear regression (univariable analyses after a logarithmic transformation of the dependent variable: serum ferritin level)

^d For cells with a zero value, 1 was added to all values of the contingency table as suggested [50]

e 95% CI of odds ratio (OR) in multivariable analyses

^f no deviance to linearity in the linear model or the univariate logistic regression model

^g no change in the results taking into account time interval between iron supplementation and serum ferritin level measure

Abbreviations: IFF = iron fortified formula, YCF = young-child formula

formula (YCF) consumption, other dietary, perinatal and sociodemographic characteristics 576

577

	n (%)"		Serum ferritin level (µg/L)							
	n = 561	β ^b	95% CI ^f	β ^c	95% CI ^f	β ^d	95% CI ^f	β ^e	95% CI ^f	
YCF considering di	fferent definitio	ons								
Consumption of Y	CF after 10 mo	nths								
No	48 (9)	-	-	-	-	-	-	-	-	
Yes	512 (91)	0.28	0.12-0.45	-	-	-	-	-	-	
Daily volume of YC	CF consumed (r	mL/day)								
No YCF	150 (27)	-	-	-	-	-	-	-	-	
< 100	175 (31)	-	-	0.34	0.20-0.47	-	-	-	-	
100-300	103 (18)	-	-	0.25	0.10-0.39	-	-	-	-	
300-500	101 (18)	-	-	0.33	0.18-0.48	-	-	-	-	
≥ 500	32 (6)	-	-	0.39	0.17-0.60	-	-	-	-	
CF consumption	duration (mon	ths) ^g								
	11.3 (3.0)	-	-	-	-	0.03	0.01-0.04	-	-	
Duration of consun	nption of IFFs	(since age 6 1	nonths) ^g							
	15.3 (5.3)	-	-	-	-	-	-	0.03	0.01-0.06	
Other dietary chara	cteristics									
Exclusive » breas	tfeeding > 6 m	onths								
No	484 (86)	-	-	-	-	-	-	-	-	
Yes	77 (14)	-0.08	-0.21-0.06	-0.12	-0.24-0.02	0.04	-0.11-0.18	0.06	-0.08-0.21	
ate food diversific	ation > 6 mont	ths								
No	546 (98)	-	-	-	-	-	-	-	-	
Yes	10(2)	0.13	-0.21-0.46	0.13	-0.20-0.46	0.07	-0.31-0.45	0.14	-0.18-0.47	
Cow's milk consum	ption beginnin	ng before 24	months							
No	356 (64)	-	-	-	-	-	-	-	-	
Yes	201 (36)	-0.17	-0.26-(-0.07)	-0.05	-0.16-0.07	-0.12	-0.23-(-0.01)	-0.12	-0.22-0.01	
Dietary iron intake	from all food	excluding YO	CF ^h (mg/day)							
Continuous	4.7 (2.2)									
< 3.5	161 (29)	-	-	-	-	-	-	-	-	
3.5-7	337 (60)	-0.05 ^h	-0.21-0.12	-0.06 ^h	-0.22-0.11	-0.06	-0.17-0.04	-0.06	-0.16-0.04	
7-10.5	48 (9)	-0.10 ^h	-0 29-0 10	-0.11 ^h	-0 30-0 09	-0.01	-0 19-0 17	-0.02	-0 19-0 15	
> 10.5	15(3)	-0.12 ^h	-0.32-0.09	-0.13 ^h	-0.33-0.07	-0.17	-0.51-0.16	-0.16	-0.46-0.14	
ron supplementation	on since birth ⁱ	0.12	0.52 0.05	0.12	0.00 0.07	0.17	0.51 0.10	0.10	0.10 0.11	
No	503 (90)	-	-	-	-	-	-	-	-	
	57 (10)	0.13	0.03.0.20	0.14	-0.02-0.29	0.12	-0.03-0.29	0.14	0.02.0.30	

587 588 589 590 591 592 593 594 595 596 597 ^e β coefficient of a multiple linear regression model using the variable duration of consumption of iron-fortified formula since 6 months-old ^f95% CI of the β coefficient of a multiple linear regression model corresponding

^g no deviance to linearity

^h creation of a fractional polynomial degree 2 for this variable in a few models because of deviance to linearity (as shown by italics). Results for dietary iron intake excluding YCF are presented by categories for ease of reading

ⁱ no change in the results taking into account time interval between iron supplementation and serum ferritin level measure

Perinatal and sociodemographic variables in these models are not shown: in all models a lower SF was independently associated with a medical history of prematurity, a high rank in siblings, and a low father's education level

Abbreviations: CI = confidence interval, IFF = iron fortified formula