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# Risk-Benefit Assessment of Cereal-Based Foods Consumed by Portuguese Children Aged 6 to 36 Months—A Case Study under the RiskBenefit4EU Project

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**Abstract:** Cereal-based foods, including breakfast (BC) and infant cereals (IC), are among the first solid foods introduced to infants. BC and IC are sources of macro and micronutrients that have beneficial effects on health, but can also be sources of harmful chemical and microbiological contaminants and nutrients that may lead to adverse health effects at high consumption levels. This study was performed under the RiskBenefit4EU project with the aim of assessing the health impact associated with consumption of BC and IC by Portuguese children under 35 months. Adverse effects associated with the presence of aflatoxins, *Bacillus cereus*, sodium and free sugars were assessed against the benefits of fiber intake. We applied a risk–benefit assessment approach, and quantified the health impact of changes in consumption of BC and IC from current to various alternative consumption scenarios. Health impact was assessed in terms of disability-adjusted life years. Results showed that moving from the current consumption scenario to considered alternative scenarios results in a gain of healthy life years. Portuguese children can benefit from exchanging intake of IC to BC, if the BC consumed has an adequate nutritional profile in terms of fiber, sodium and free sugars, with levels of aflatoxins reduced as much as possible.

**Keywords:** public health; risk–benefit assessment; cereal-based foods; children; mycotoxins; *Bacillus cereus*; sodium; fiber; free sugars

## 1. Introduction

Cereal-based foods, including breakfast cereals (BC), i.e., ready-to-eat processed grains; and infant cereals (IC), i.e., processed cereal-based foods that are or have to be reconstituted with water, milk, or other appropriate liquids, are among the first solid foods introduced to infants (age 0–1 ([www.cdc.gov](http://www.cdc.gov) (accessed on 19 August 2021))) and are important components of the diet for toddlers (age 0–3 ([www.cdc.gov](http://www.cdc.gov) (accessed on 19 August 2021))) [1,2]. They are sources of nutrients necessary for a healthy development, including macronutrients (e.g., fiber) and essential micronutrients (e.g., vitamin B2, potassium, magnesium). Cereal-based foods are also an important source of whole grains [3,4]. Additionally, some BCs and ICs may be fortified with vitamins such as folate and thiamine. However, cereal-based foods can also be a vehicle for hazardous contaminants or other components that can potentially lead to adverse health effects. Cereals can be contaminated with harmful chemicals such as mycotoxins [5–8], heavy metals [9–13], acrylamide [14], furans [15] and polycyclic aromatic hydrocarbons (PAHs) [16] that are associated with acute and chronic toxic effects, such as gastroenteritis, growth impairment, immune-toxicity and cancer. Despite being considered low moisture foods (LMF), food monitoring has shown that cereals may be contaminated with *Bacillus cereus*, a Gram-positive, rod-shaped, spore-forming bacterium. *B. cereus* can lead to two types of illness in humans: emetic (caused by preformed toxins in food) and diarrheal (caused by ingestion of a large number of bacterial cells), and foodborne outbreaks have identified cereals as a vehicle [17]. Depending on their recipes, cereal-based foods may also be a key source of sodium and free sugars, which, at high consumption levels, increase the risk of type 2 diabetes, cardiovascular disease and cancer [18,19].

In Portugal, previous studies have reported that ICs and BCs constitute a considerable proportion of the diet of infants and toddlers (children under three years of age) [5,6,20–22]. ICs are traditionally considered suitable as the first solid foods introduced to infants at 6 months of age. BCs are also marketed for infants but considered more suitable for toddlers of older age. According to products available on the Portuguese market, BCs usually contain several types of grains, more fiber and less sodium and added sugar than ICs [5,6,22]. Previous studies have demonstrated the occurrence of mycotoxins in both BCs and ICs available on the Portuguese market [5–7,23,24] and the risk associated with exposure to aflatoxins through consumption of cereal-based foods [6]. Likewise, the consumption of IC and BC by young children could represent a significant contribution to the daily intake of sodium and sugar, but also to overall energy intake (recommended energy intake for children up to 36 months is, according to Food and Agriculture Organization (FAO), up to 427 KJ/kg bodyweight per day [25]), and act as risk factors for the development of future non-communicable diseases. Therefore, it is not straightforward to predict the health impact following a change in consumption of BC and IC, and an approach allowing for the integration of associated health risks and benefits will be needed in order to inform dietary recommendations that maximize health benefits for children.

Risk–benefit assessment (RBA) of foods aims to assess the combined adverse and beneficial health effects associated with foods. RBA integrates chemical, microbiological and nutritional risk and benefit assessments [26–29] and is used to inform food safety and public health strategies, including updating dietary advice [30]. “RiskBenefit for EU—Partnering to strengthen the risk–benefit assessment within EU using a holistic approach” (RB4EU) was a knowledge translation project funded by the European Food Safety Authority (EFSA), which gathered a multidisciplinary team and developed a framework for capacity building in RBA through case studies [26,31].

The aim of this case study was to perform an RBA to quantify the overall health impact in terms of DALYs from changes in the consumption of BC and IC by Portuguese children below 36 months of age. Disability-Adjusted Life Years (DALYs) are a composite health metric commonly applied to integrate risks and benefits in an estimate of overall health gain or loss following dietary changes [32]. We followed the harmonized framework developed under the RiskBenefit4EU project to answer the question: how many healthy

life years are gained or lost by changes in the consumption of BC and IC by Portuguese children below 35 months of age?

## 2. Materials and Methods

### 2.1. Selection of Relevant Components and Health Effects

A two-step literature search was carried out to identify the BC and IC components (contaminants/nutrients) and associated health effects to be considered in the study: step (1) a “food–component” literature search, to identify components of interest for the food products considered; step (2) a “component–health effect” literature search, to identify the health effects associated with specific food components. The search strategies, identified components and health effects are summarized in Appendix A. Table 1 summarizes the selected food components, their associated health effects and the type of analysis performed.

**Table 1.** Food components and health effects considered for the risk–benefit assessment.

	Food Components	Health Effects	Type of Analysis
Nutrition	Total fiber	Type 2 Diabetes mellitus Cardiovascular diseases	Quantitative <sup>a</sup>
	Sodium	Type 2 diabetes	Semi-quantitative <sup>b</sup>
	Free sugars	Cardiovascular disease Cancer (different organs)	
Toxicology	Aflatoxins (AFB <sub>1</sub> ) <sup>c</sup>	Hepatocellular carcinoma	Quantitative <sup>a</sup>
Microbiology	<i>Bacillus cereus</i>	Gastrointestinal disease	Quantitative <sup>a</sup>

<sup>a</sup> Risks and benefits quantified in DALY; <sup>b</sup> Intake compared with the dietary reference values; <sup>c</sup> AFB<sub>1</sub>: aflatoxin B<sub>1</sub>.

Mycotoxins are widely occurring secondary metabolites produced by fungi that can contaminate foods and feeds. Aflatoxins, a subgroup of mycotoxins, are genotoxic, carcinogenic and immunosuppressive substances. The four major aflatoxins are B<sub>1</sub>, B<sub>2</sub>, G<sub>1</sub> and G<sub>2</sub>. Aflatoxin B<sub>1</sub> (AFB<sub>1</sub>) is the most potent naturally occurring chemical liver carcinogen known. These toxins cause hepatocellular carcinoma (HCC) [33] and thus have been classified as Group 1 carcinogens (carcinogenic to humans) by the International Agency for Research on Cancer [34]. HCC is the third leading cause of cancer deaths worldwide and was estimated to have been responsible for nearly 746,000 deaths in 2012 (9.1% of all cancer deaths that year) [35].

*B. cereus* is a foodborne pathogen that causes diarrhea and vomiting [36], and is the most frequent causative agent of foodborne illnesses in cereals and grains. It was responsible for foodborne outbreaks associated with exposure to cereal products in different regions of the United States, Australia, New Zealand, Asia and Europe [17]. Some of the reported outbreaks involved IC [36] and ready-to-eat cereals [37,38].

Dietary fiber may contribute to the prevention of childhood obesity, maintenance of normal blood glucose, lipids and blood pressure, however, the scientific evidence is rather contradictory for some of these outcomes [39]. On the other hand, there is convincing evidence that dietary fiber intake promotes normal gastrointestinal function, especially laxation [40], and convincing/probable evidence that it reduces the risk of colon cancer [41–43]. Moreover, probable evidence shows that fiber intake reduces the risk of cardiovascular disease (CVD) and type 2 diabetes (DM2) [40,41,43,44]. Children with higher intakes of dietary fiber tend to consume diets that are more rich in nutrients and are more likely to meet recommended daily intakes for key nutrients [45]. Cereal-based foods with a high content of wholegrains constitute an important source of dietary fiber.

Sodium is an essential cation that is necessary for normal cell function and for neurotransmission. However, high sodium intake is associated with increased blood pressure among adults and children and with a high risk of cardiovascular diseases among adults. The World Health Organization (WHO) recommends a maximum sodium intake of 2 g/day in adults. For children, this value should be adjusted downward based on the energy re-

quirements of children when compared to adults [18]. Accordingly, EFSA recommends an intake of 1.1 g of sodium per day as safe and adequate for children aged 1–3 years old [46].

The WHO defines free sugars as all monosaccharides and disaccharides added to foods (i.e., added sugars), and sugars naturally present in honey, syrups, and fruit juices. Moderate-quality evidence suggests that reduced intake of free sugars is associated with reduced body weight in children [19]. However, higher rates of dental caries are observed when free sugar intake exceeds 10% of total energy intake [19]. Thus, the WHO recommends that free sugars should not exceed 10% of total energy intake, with the conditional recommendation to reduce free sugar intake to below 5% of total energy intake [19].

Finally and according to Table 1, total aflatoxin-HCC, *B. cereus*-gastrointestinal disease and dietary fiber-DM2 & CVD were included in the quantitative RBA along with sodium and free sugars, in a semi-quantitative approach comparing consumption scenarios using dietary reference values (DRV).

## 2.2. Investigated Consumption Scenarios

In order to assess health impact due to changes in BC and IC consumption, we investigated four scenarios:

1. 100% BC scenario, simulating that all infants consume only BC;
2. 100% IC scenario, simulating that all infants consume only IC;
3. Optimal BC scenario, simulating that all infants consume only BC at an optimal composition; and
4. Worst IC, simulating that all infants consume only IC with the worst composition.

Each of the above scenarios was analyzed relative to the current amount consumed, in which only IC, BC or both (i.e., the reference scenario) are consumed by children aged 6 to 35 months, as reported by the National Food, Nutrition and Physical Activity Survey of the Portuguese General Population 2015–2016 (IAN-AF 2015–2016) [22]. Current consumption is defined as the consumption of IC and BC reported in the latest survey of Portuguese dietary habits [22], as described below under 2.3. Specifically, the consumption of IC and BC in scenario 1 and 2 was modelled, by substituting the current consumption of IC by BC and BC by IC, respectively, based on the observed distribution of consumed IC and BC [22]. For scenarios 3 and 4 the best BC and worst IC products were identified among the BC and IC products available on the market, by scoring each product according to their content of fiber, sodium and free sugar per 100 kcal (Appendix A). To create the score, the variables fiber, sodium and free sugar per 100 kcal were first standardized and then summed. Fiber was considered a positive parameter; sodium and free sugar negative ones. The score was calculated for each product as follows:

$$Score = z_{fiber} - (z_{sodium} + z_{free\ sugar}) \quad (1)$$

The best BC was the one with the highest score among all BC products and the worst IC was the one with the lowest score among all IC products (Table 2). Scenarios 3 and 4 were modelled by substituting the current consumption of IC and BC by the BC with the highest score (“Best BC”) and by the IC with the lowest score (“Worst IC”), respectively. Substitutions in all four scenarios were iso-caloric.

**Table 2.** Nutritional composition of “Best Breakfast cereal (BC)” and “Worst Infant cereal (IC)” products selected among the cereal products available on the Portuguese market.

	Fiber (g/100 kcal)	Sodium (mg/100 kcal)	Free Sugar (g/100 kcal)
Best BC	3.9	1.4	0
Worst IC	0.2	51.4	4.2

## 2.3. Data Used in the Model

Food consumption data were collected from the National Food and Physical Activity Consumption Survey (IAN-AF 2015–2016), a representative sample of the Portuguese

general population aged between 3 months and 84 years old, with a total of 5811 participants [20–22]. In this study, food consumption data from 779 children aged 6 to 36 months from the IAN-AF 2015–2016 sample was included. The sample size of participants was determined, considered in the power analysis, and presented in detail in the methodological publication of the IAN-AF 2015–2016 survey [21]. Data on food consumption were collected according to European guidelines [47]. For children aged under 10 years old, dietary intake was assessed by two non-consecutive one-day food diaries that were filled in by the main caregiver, followed by a face-to-face interview conducted with the caregivers to collect additional details on food description and quantification. The survey used an extended version of the Portuguese Food Composition Table [48] to convert the reported food items into nutrients, covering 38 BCs and 87 ICs.

Concentration data of aflatoxins in ICs and BCs available on the Portuguese market were collected from published surveys [5–7]. Detailed information about analytical conditions and method performance of aflatoxin determination were described elsewhere [6,7]. Occurrence levels of aflatoxins in both BCs and ICs were below legislative limits (infant cereals: 0.100 µg/kg for AFB<sub>1</sub>; BC: 2.0 µg/kg for AFB<sub>1</sub> and 4.0 µg/kg for the sum of AFB<sub>1</sub>, AFB<sub>2</sub>, AFG<sub>1</sub> and AFG<sub>2</sub>) [49].

Prevalence and level of contamination of *B. cereus* in ICs and BCs were modeled using results from the Portuguese National Sampling Plan (PNCA) for BC [50] and available data from the scientific literature for IC [37]. Between 2018 and 2019, the PNCA analyzed 50 samples of BC, corresponding to 30 different batches, for which 5 boxes were tested. Fourteen out of 150 samples had positive levels of *B. cereus*. For IC, data from another study carried out in Portugal in 2007 were applied, in which 35 samples were analyzed, including 15 positives [37]. For both IC and BC, samples below the limit of detection (LOD) (<10 colony forming units (CFU)/g), were assumed to have 5 CFU/g. For BC, positive samples below 40 CFU/g were reported as <40 CFU/g in the database; these were assumed to have 25 CFU/g, i.e., the median between 10 and 40.

Regarding DALY, this metric combines information on disease incidence, severity, duration and mortality in one number and enables comparison across diseases [51,52]. The epidemiological and toxicological data used to derive the correspondent DALY are presented in Table 3.

**Table 3.** Epidemiological and toxicological data used for the model.

	Dose–Response (Cases/100,000/year/ng/kg AFT bw/day)	Incidence <sup>2</sup> (Cases/100 k) (Mean, 95% CI)	Risk Ratios (Mean, 95% CI)	DALY Rate <sup>2</sup> (DALY/100 k) (Mean, 95% CI)
Hepatocellular carcinoma (HCC)	0.01 <sup>3</sup>	NA	NA	9.96 (7.77–12.61)
Cardiovascular disease (CVD)	NA	25.07 (21.36–29.12)	0.78 <sup>1</sup> (0.68–0.90)	60.78 (49.67–73.44)
Diabetes mellitus 2 (DM2)	NA	168.07 (125.70–215.47)	0.85 <sup>1</sup> (0.82–0.89)	22.4 (12.71–34.47)

NA = Not applicable; CI = Confidence interval; DALY = Disability-adjusted life years; AFT = Aflatoxins; <sup>1</sup> per 8 g/day of fiber [39]; <sup>2</sup> Global Burden of Disease 2017 study (GBD) [53]; <sup>3</sup> WHO (2017) [54].

#### 2.4. Exposure Assessment, DALY Calculation and Integration of Risks and Benefits

We applied different approaches for exposure and health impact assessment for each component–health effect pair.

All models were developed and implemented in the @Risk<sup>®</sup> software for Microsoft Excel version 6 (Palisade Corporation, Ithaca, NY, USA). We defined model variables as probabilistic distributions to quantify uncertainty. The approach applied for each component is described below and all models are documented in the Supplementary material (Table S1). SPADE software [55] was used to estimate the intake distributions of total fiber, sodium and free sugars in the current and each alternative scenario.



#### 2.4.1. Total Fiber, Sodium and Free Sugars

In order to estimate the relative risk (RR) of the current and each alternative intake scenario, we performed the following calculations. The distribution of fiber intake of children aged 6 to 35 months, which was derived from the National Food and Physical Activity Consumption Survey [22], was divided into quartiles, with each quartile representing a consumption class (1–4). The median of each class represented the intake of each class, respectively. RRs for DM2 and CVD derived from the literature (Table 3) were used to estimate an RR for each class, assuming an RR of 1 at zero exposure and a log-linear association between exposure and RR [56] (Appendix B and Table S2). Thus, the log-linear slope,  $\beta$ , and RR for each class,  $j \in \{1, 2, 3, 4\}$ , in each scenario,  $i \in \{1, 2, 3, 4, 5\}$  (i.e., the reference and the four alternative scenarios), were calculated according to the following equations:

$$\beta = \frac{\ln RR_{literature}}{Dose} \quad (2)$$

$$RR_i = \exp(\beta \cdot exposure_i) \quad (3)$$

where  $RR_{literature}$  is the RR for DM2 and CVD per  $Dose$  reported in the literature (Table 3), and  $RR_i$  and  $exposure_i$  are the RR and intake of fiber in each scenario.

The potential impact fraction (PIF) is a measure of the proportional change in disease burden after a change in exposure to a related risk factor—in this case changes in exposure to fiber from the current to alternative scenarios—and was calculated for each alternative scenario [57]. We applied RR shift methodology [57], which assumes that the change in exposure is described by a change in the RR of the categories, while keeping the proportion in each category constant:

$$PIF = \frac{\sum_{j=1}^4 RR_{alt} - \sum_{j=1}^4 RR_{ref}}{\sum_{j=1}^4 RR_{ref}} \quad (4)$$

where  $RR_{ref}$  is the RR of the reference scenario and  $RR_{alt}$  is the RR of each alternative scenario.

We calculated the incidence and DALYs attributed to the change in IC and BC consumption in each scenario by multiplying incidence and DALY rates for DM2 and CVD for Portugal, obtained from the Global Burden of Disease 2017 study (GBD) [53] (Table 3), by the PIFs for each alternative scenario.

A semi-quantitative analysis was performed for sodium and free sugars. The DRVs used were an upper-limit (UL) of 1500 mg/day for sodium [58] and a recommended intake (RI) of 5% and 10% of total energy intake (TEI) for free sugars [19]. The prevalence of individuals above or below the DRVs was estimated for each scenario using SPADE software [55].

#### 2.4.2. Aflatoxins

We estimated the exposure to aflatoxins from BC and IC consumption by:

$$\text{AFT intake } (\mu\text{g}/\text{kg bw}/\text{day}) = \frac{\text{Conc}_{\text{AFT}} \times \text{Consump}}{\text{bw}}, \quad (5)$$

where  $\text{Conc}_{\text{AFT}}$  is the concentration of aflatoxins determined in BC and IC in  $\mu\text{g}/\text{kg}$ ;  $\text{Consump}$  is the consumption of BC and IC in  $\text{kg}/\text{day}$ ; and  $\text{bw}$  is the body weight of the considered population in  $\text{kg}$ .

We used the total concentration of aflatoxins measured in cereal samples, assuming that all aflatoxins have equal carcinogenic potency—a conservative approach previously applied by EFSA [59]. Values below the LOD can be replaced via different approaches as recommended by EFSA: lower-bound (LB,  $<\text{LOD} = 0$ ), middle-bound (MB,  $<\text{LOD} = 1/2 \text{ LOD}$ ) and upper-bound (UB,  $<\text{LOD} = \text{LOD}$ ) approaches [60]. The different approaches were applied for aflatoxin concentrations in scenarios as follows: (i) an MB approach was used for

the reference, 100% IC and 100% BC scenarios; (ii) a LB approach was used for the Best BC scenario, since the best breakfast cereals would have the minimum possible contamination by aflatoxins; (iii) a UB approach was used for the Worst IC scenario, since the worst infant cereal would have the maximum possible contamination by aflatoxins.

We estimated the number of extra HCC cases caused by exposure to aflatoxins from BC and IC in the reference scenario and each alternative scenario by combining the DRV related to aflatoxin exposure and the risk for HCC (Table 3) [54]. The DALYs attributed to the estimated aflatoxin exposure were calculated by multiplying the estimated extra HCC cases by a DALY/case rate, derived from WHO incidence and the DALY envelope [53].

#### 2.4.3. Bacillus Cereus

A beta distribution was used to approximate prevalence based on the percentage of positive values; it was expressed as a beta of “number of positive + 1; total sample—number of positive + 1”. For positive samples, to represent the level of *B. cereus*, a cumulative distribution was built for BC and a triangular one for IC. Simulations were run to represent the level of *B. cereus* in IC and BC, giving a value of 5 CFU/g when considered to be non-contaminated (<10 ufc/g) and selecting randomly for positive sample levels from the cumulative or uniform distribution of BC and IC, respectively.

To estimate exposure to *B. cereus* (in CFU/day) through consumption of BC and IC in the considered population group, contamination data in CFU/g and consumption data in g/day were multiplied. This was done by multiple simulations to combine the different possible exposure values (observed for the reference scenario or estimated for alternative scenarios). All variables were defined as probability distributions, informed by the available data.

Quantitative DR for *B. cereus* were not available in the literature. Consequently, a threshold approach was used considering a potential case of illness when the product contamination exceeded a threshold of  $1.0 \times 10^3$  CFU/g [61] or when the overall level of daily exposure to bacteria exceeded a threshold of  $5.0 \times 10^4$  CFU/day [36].

#### 2.4.4. Integration of DALY

The difference in DALYs over each health outcome attributed to a change in IC and BC consumption was summed to estimate the overall health impact ( $\Delta$ DALY) of each scenario.  $\Delta$ DALY > 0 implies a health loss due to the change in consumption.  $\Delta$ DALY < 0 implies a health gain due to the change in consumption.

### 3. Results

#### 3.1. Intake and Exposure Assessment

The estimated median daily consumption of BC and IC by Portuguese children between 6 and 35 months of age (reference scenario) was 0.0 g and 8.7 g, respectively (Table 4). The substitution of the current consumption of IC and BC by the Best BC scenario resulted in the highest daily consumption (14.5 g/day) among the alternative scenarios.

**Table 4.** Distribution of consumption represented by the median (P5-P95) of BC and IC consumption by Portuguese children between 6 and 35 months of age, in reference and alternative scenarios.

	Reference	100% BC	100% IC	Best BC	Worst IC
BC (g/day)	0.0 (0.0–20.8)	13.4 (0.0–51.1)	-	14.5 (0.0–55.5)	-
IC (g/day)	8.7 (0.0–42.7)	-	12.5 (0.0–50.1)	-	13.7 (0.0–52.3)

P5: 5th percentile; P95: 95th percentile; BC: breakfast cereals; IC: infant cereals.

The Best BC scenario resulted in a mean fiber intake of 11.3 g/day, i.e., around 2 g/day higher than the other scenarios (Table 5). Regarding sodium and free sugars, intake was estimated to be lower in the Best BC scenario when compared to the remaining scenarios (Table 5). The Best BC scenario also resulted in the lowest mean exposure to aflatoxins. This scenario also reflects a full reduction in exposure to *B. cereus*, as no contaminated samples

were found where it could be targeted, contrary to the Worst IC scenario, which led to around 3.3 log CFU/day (Table 5).

**Table 5.** Usual intake distribution of fiber (g/day), sodium (g/day), free sugars (g/day) and exposure to aflatoxins (ng/kg/day) and *Bacillus cereus* (log CFU/day) in Portuguese children between 6 and 35 months of age, in reference and alternative scenarios.

		Reference	100% BC	100% IC	Best BC	Worst IC
Fiber (g/day)	Mean	9.3	9.5	9.2	11.3	8.9
	Median	9.0	9.2	9.0	11.0	8.7
	(P25–P75)	(7.2–11.1)	(7.3–11.4)	(7.1–11.0)	(8.8–13.5)	(6.8–10.7)
Sodium (g/day)	Mean	1.17	1.18	1.16	1.15	1.18
	Median	1.15	1.17	1.14	1.13	1.16
	(P25–P75)	(0.79–1.50)	(0.80–1.51)	(0.79–1.49)	(0.77–1.47)	(0.80–1.51)
Free sugars (g/day)	Mean	19.5	20.0	19.4	16.3	19.0
	Median	17.8	18.4	17.7	14.2	17.4
	(P25–P75)	(11.8–25.2)	(12.4–25.8)	(11.8–25.1)	(8.8–21.5)	(11.6–24.6)
Aflatoxins (ng/kg bw/day)	Mean	0.065	0.073	0.065	0.052	0.090
	Median	0.049	0.054	0.047	0.038	0.066
	(P25–P75)	(0.0–0.95)	(0.0–0.109)	(0.0–0.094)	(0.0–0.078)	(0.0–0.136)
<i>Bacillus cereus</i> (log CFU/day)		2.5	2.1	2.6	<0 *	3.3

P25: 25th percentile; P75: 75th percentile; bw: body weight; BC: breakfast cereals; IC: infant cereals; \* <0 log ufc/g equals <1 ufc/g.

### 3.2. Incidence and DALYs

For DM2 and CVD, the Best BC scenario presented the highest number of cases prevented by change in fiber intake from the reference scenario. The Worst IC scenario presented the highest number of extra cases caused by change in fiber intake (Table 6). The highest number of extra cases of HCC was estimated for the Worst IC scenario (Table 6). Regarding *B. cereus*, no cases of gastrointestinal illnesses were estimated as all scenarios provided exposure levels far below both thresholds of illness; all simulations estimated exposure levels below the threshold of  $5.0 \times 10^4$  CFU/day [36] and product concentrations below the maximum limit of  $1.0 \times 10^3$  CFU/g [61].

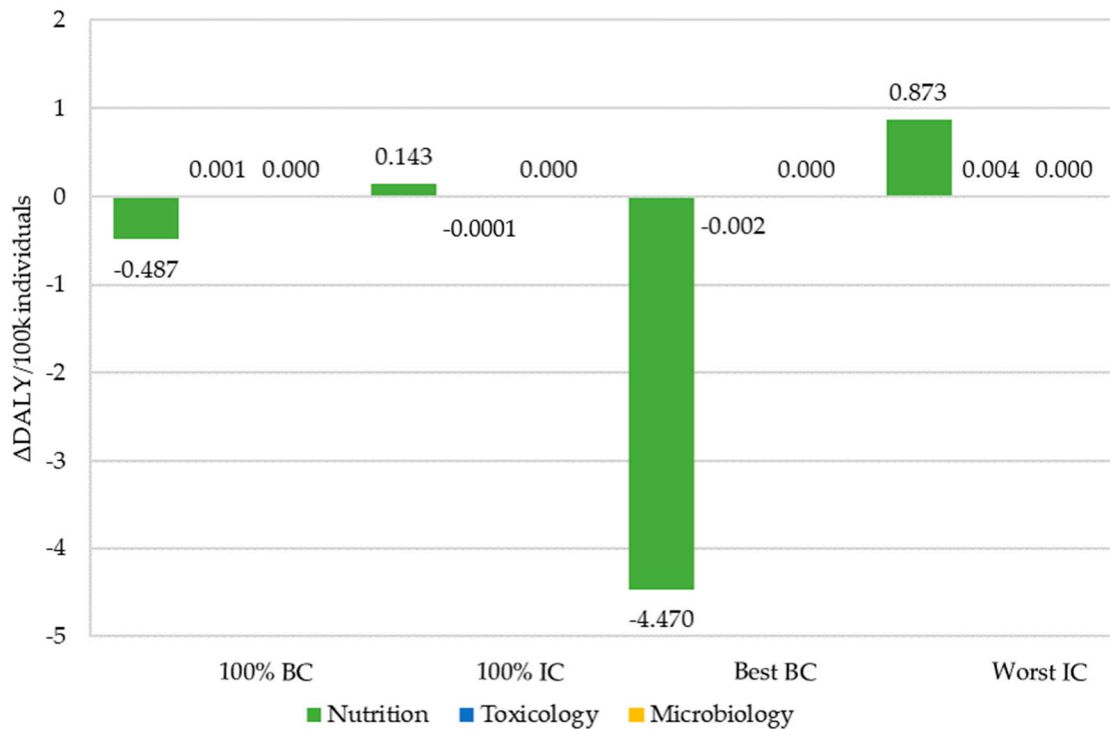
**Table 6.** Number of prevented or extra cases of type 2 diabetes (DM2), cardiovascular diseases (CVD) and hepatocellular carcinoma (HCC) due to consumption of BC and IC by Portuguese children between 6 and 35 months of age in each alternative scenario.

	100% BC	100% IC	Best BC	Worst IC
DM2 (fiber)				
Number of cases/100,000/year *	−0.714	0.211	−6.618	1.272
(95% CI)	(−0.998; −0.482)	(0.143; 0.295)	(−9.229; −4.489)	(0.858; 1.782)
CVD (fiber)				
Number of cases/100,000/year *	−0.160	0.047	−1.469	0.288
(95% CI)	(−0.253; −0.072)	(0.021; 0.074)	(−2.288; −0.668)	(0.127; 0.458)
HCC (AFB <sub>1</sub> )				
Number of cases/100 k/year **	0.00073	0.00065	0.00077	0.00090
(95% CI)	(0; 0.0027)	(0; 0.0023)	(0; 0.0028)	(0; 0.0033)
Number of Gastrointestinal disease ( <i>Bacillus cereus</i> )	0	0	0	0

\* Difference in incident cases between alternative and reference scenarios; BC: breakfast cereals; IC: infant cereals, \*\* Number of extra incident cases due to aflatoxin exposure in each scenario.



Figure 1 presents the  $\Delta$ DALY across the health outcomes accounted for in each domain, i.e., nutrition (fiber), toxicology (aflatoxins) and microbiology (*B. cereus*) for the different scenarios assessed. As no cases of gastroenteritis due to *B. cereus* were identified in any of the scenarios, no DALYs were likewise estimated. The 100% BC and Best BC scenarios resulted in negative  $\Delta$ DALY for fiber, indicating a health gain associated with a change in intake to these scenarios from the current scenario. On another hand, the 100% IC and Worst IC scenarios resulted in a positive  $\Delta$ DALY, indicating a health loss due to the change in intake (Figure 1). Contribution from aflatoxins was negligible, accounting for a  $\Delta$ DALY close to zero in all the considered scenarios.



**Figure 1.** Health impact per domain of nutrition, toxicology and microbiology in each alternative scenario expressed in DALYs/100,000/year. BC: breakfast cereals; IC: infant cereals.

When integrating  $\Delta$ DALY over the domains, an overall  $\Delta$ DALY of 0.143 and 0.877 per 100,000 individuals in the 100% IC scenario and Worst IC scenario, respectively, was estimated. In contrast, the 100% BC and Best BC scenarios resulted in an overall negative  $\Delta$ DALY of  $-0.486$  and  $-4.473$ , respectively (Table 7).

**Table 7.** Overall DALY difference ( $\Delta$ DALY) estimates for each alternative scenario, integrating risks and benefits associated with consumption of breakfast cereals (BC) or infant cereals (IC) by Portuguese children aged between 6 and 35 months.

	100% BC	100% IC	Best BC	Worst IC
Sum of $\Delta$ DALY (per 100 k individuals) (95% CI)	$-0.486$ ( $-0.727$ ; $-0.262$ )	$0.143$ ( $0.078$ ; $0.213$ )	$-4.473$ ( $-6.614$ ; $-2.449$ )	$0.877$ ( $0.471$ ; $1.317$ )

BC: breakfast cereals; IC: infant cereals.

### 3.3. Comparison to DRVs

Table 8 presents the prevalence of inadequate intake of sodium and free sugars when compared to DRVs. These results reflect the intake of sodium and free sugars from all dietary sources, and not only the contribution of IC and BC in all the scenarios considered. We estimated that 25% of Portuguese children between 6 and 35 months of age exceed the upper-limit (UL) intake for sodium in the reference scenario, where BC and IC contribute

0.7% and 1.3% to total sodium, respectively. This prevalence was increased slightly in the 100% BC and Worst IC scenarios compared to the reference scenario (from 24.8% to 25.7% and 25.4%, respectively). In the Best BC scenario, the prevalence was reduced to 24% over the UL. Currently, 29% have an intake of free sugars higher than 10% of TEI. The prevalence of individuals exceeding the RI was lowest (16%) in the Best BC scenario. For all scenarios, the prevalence of exceeding the RI of free sugars considerably increased if 5% TEI was considered as the cut-off.

**Table 8.** Prevalence of Portuguese children aged 6–35 months with an intake of sodium and free sugars, from all dietary sources, above the DRV in the reference and alternative scenarios (semi-quantitative approach).

DRV	Reference	100% BC	100% IC	Best BC	Worst IC
<b>Sodium (UL)</b>					
1500 mg/day	24.8%	25.7%	24.3%	23.4%	25.4%
<b>Free sugars (RI)</b>					
5% TEI	80.0%	81.9%	79.8%	69.7%	79.0%
10% TEI	29.4%	31.6%	29.2%	16.0%	27.7%

DRV: Daily Reference Value; UL: upper-limit; RI: recommended intake; TEI: total energy intake; BC: breakfast cereals; IC: infant cereals.

## 4. Discussion

### 4.1. Consumption of Breakfast Cereals Instead of Infant Cereals Could Result in a Gain of Healthy-Life Years

This study assessed the risk–benefit balance of consuming different proportions of BC and IC by Portuguese children aged between 6 and 35 months. The health risks associated with the presence of aflatoxin and *B. cereus* were evaluated as well as the benefits of the intake of fiber, by comparing four different scenarios of BC and IC consumption with current consumption. Overall health impact was expressed in DALYs. The change in the intake of sodium and free sugars from current to alternative consumption scenarios were evaluated by a semi-quantitative approach. We found that the substitution of current consumption by 100% BC and Best BC would lead to a gain in healthy-life years. The Best BC scenario improved the percentage of the considered population exceeding the UL and RI for sodium and free sugars, respectively. To our knowledge, this is the first RBA of cereal-based foods intended for children aged between 6 and 35 months.

Our results suggest that children can benefit from exclusive BC consumption if product formulations are optimized in terms of fiber, sodium and free sugar content, and at the same time, aflatoxin levels are reduced as much as possible. Choosing the best BC available in the Portuguese market would benefit population health in terms of prevention of DM2 and CVD and a decrease in the prevalence of exceeding the UL and RI of sodium and free sugars, respectively. The consumption of this cereal-based product could prevent a loss of about 4.5 healthy life years per 100,000 individuals annually. However, BCs with high fiber and low sodium and free sugars content could lead to low palatability and consequently a low adherence by consumers, especially for the considered age group. Our results also showed that the negative impact on health due to exposure to aflatoxins was minor in all scenarios considered. However, previous studies performed in Portugal [5,6] and abroad [62] have stressed the risk associated with the consumption of cereal-based foods, including BC and IC, due to exposure to aflatoxins present in these food products—showing, for the highest percentiles of intake, a margin of exposure below 10,000, representing a potential health risk [5,6]. Cereal-based products constitute one of the main contributors of human exposure to mycotoxins and the Food and Agriculture Organization of the United Nations (FAO) estimated that at least 25% of cereals produced in the world are contaminated by mycotoxins, acknowledging that this value could be higher than 50% if emerging mycotoxins are considered, but of which limited data are so far available [63,64]. Mycotoxins are natural contaminants and cannot be completely

prevented. Consequently, significant efforts should be considered, from farm to fork, to improve the safety and quality of foods, reducing mycotoxin exposure—particularly to carcinogenic aflatoxins.

Recognizing the potential benefits, there is a growing interest in the reformulation of breakfast cereal products [65]. European countries have been developing and establishing programs to reduce the intake of sodium and sugar. In Italy, in an attempt to improve the nutritional characteristics of food products, one objective was to reformulate breakfast cereals by reducing sugar and salt down to mean contents of 30 and 0.4 g per 100 g, respectively, and increasing fiber to 4.5 g per 100 g [66]. In Portugal, the Integrated Strategy for the Promotion of Healthy Eating was adopted in 2017, which also considered promotion of the reformulation of some food categories, including breakfast cereals, by monitoring the salt and sugar in breakfast cereals [67]. In the effort to reformulate BCs, attention should also be paid to aflatoxin levels in these food products. Despite being consumed by this age group in Portugal, breakfast cereals are not recognized as infant foods in a legislative context [49]. If BC products were included in the category “infant foods”, stricter limits would be applicable, and consequently extra efforts to produce cereal-based foods presenting lower contamination levels of mycotoxins would be developed. Other studies based on qualitative assessments argued that the risk of exposure to aflatoxins through cereal-based product consumption is negligible compared to benefits related to nutritional compounds [68,69]. The present study confirms this perspective in a quantitative assessment. However, the ALARA (as low as reasonably achievable) principle needs to be considered to ensure that levels of aflatoxins in foods are kept as low as possible.

Strategies related to the future reformulation of food products should preferably be evaluated in quantitative assessments of the reformulations’ impact on public health. In this study, we included the impact of changes in fiber and aflatoxin exposure in a full quantitative assessment. Due to a lack of data, the health impact associated with intake of free sugars and sodium was assessed semi-quantitatively, not included in the overall DALY estimate. Nevertheless, the potential health effects associated with these two food components are likely important.

#### 4.2. Sources of Uncertainty

In the present study, the analysis of the obtained results should be considered with regard to all sources of uncertainties that were identified. These sources of uncertainty could influence the precise estimate of the health impact of the considered scenarios. Identified sources of unquantified uncertainty are listed below:

1. **Selection of relevant food components and associated health effects:** Despite an extensive literature search being performed to support decisions in the selection of food components and associated health effects, other beneficial (e.g., micronutrients used for fortification or saturated fats) or hazardous (e.g., other mycotoxins such as deoxynivalenol and zearalenone, heavy metals, furans, acrylamide) components could be present in BC and IC; also, food allergies are relevant health outcomes when investigating the health impact of infant and toddler foods, but have been disregarded in this assessment.
2. **Data used in the exposure assessment:** Concentration of *B. cereus* and aflatoxins in BC and IC were based on reported levels of analyzed samples or extracted from the Portuguese Food Composition Table (sodium, free sugars and fiber). These data are necessarily affected by uncertainty that impacted (over and/or underestimated) the exposure assessment.
3. **Risk of CVD and T2D later in life based on exposure that occurred in the first years of life:** Due to the restrictions of the data, CVD and T2D cases were estimated, taking into account the available dose–response data. However, it was assumed that exposure occurring in the first years of life will contribute to and determine cases later in life.

## 5. Conclusions

This study demonstrated that changes in consumption patterns of BC and IC in children below 35 months of age in Portugal would lead to health gains in the population. These results can be considered an alert for industry and risk managers, motivating strategies and measures for reduction of aflatoxins in breakfast cereals, and supporting higher content of fiber. Efforts must also continue to maintain low levels of *B. cereus* contamination, and to further reduce sodium and free sugar content in recipes. A combination of these actions will contribute to improve children's health during childhood as well as in the long-term. Application of the RB4EU framework for the risk–benefit assessment proved useful, and we encourage its use to address other risk–benefit questions.

**Supplementary Materials:** The following are available online at <https://www.mdpi.com/article/10.3390/nu13093127/s1>, Table S1: Summary of the probabilistic approach performed, including the probabilistic distributions assigned to each variable. Table S2: Median fiber intake classes based on the quartiles of intake distribution for Portuguese children aged between 6 and 35 months and risk ratio (RR) estimates (95% CI) for type 2 diabetes (DM2) and cardiovascular disease (CVD) for the different scenarios considered.

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## Appendix A. Search Strategy

The literature search strategy applied in the study involved two steps. Literature searches were conducted in the period between September 2018 and December 2018 and covered scientific papers, reports and official assessments published by expert groups in food safety and nutrition domains (European Commission, European Food Safety Authority, World Health Organization, International Agency for Research on Cancer, World Cancer Research Fund International), as well as the Portuguese Economic and Food Safety Authority (ASAE) Database and the European Rapid Alert System for Food and Feed Portal (RASFF).

The first step was conducted to identify relevant “food–component pairs”. This process resulted in the identification of the following nutritional components in BC and IC: added and free sugars, salt, saturated fat, dietary fiber, and micronutrients present due to fortification [18,19,40,70].

The following pathogenic microorganisms were identified in BC and IC and in outbreaks where breakfast cereals were the culprit foods: *Salmonella* spp. [37,71,72] and *Bacillus cereus* [36,37,61,73,74]. In the ASAE's Database, no data concerning the occurrence of pathogenic microorganisms in cereal-based products was registered. Therefore, a new breakfast cereal sampling and analysis was performed in the Microbiology Laboratory of

the ASAE, with a focus on the determination of *Bacillus cereus* in breakfast cereals on the Portuguese market. Regarding chemical contaminants, the literature review revealed the occurrence of mycotoxins [5–8], heavy metals [9,11–13,75], acrylamide [14], furans [15] and PAHs [16] in breakfast cereals and infant cereals.

The second step was conducted for the association “health effect–component” pairs and concerned the identification of potential health effects of each nutrient/contaminant/microbiological agent identified. The selection of relevant health effects was based on official assessments by international authorities, regulatory agencies, food composition databases, and scientific papers available. Considered health effects included Diabetes Mellitus 2 and cardiovascular diseases, with an emphasis on evidence related to children [76–82]. Major health risks identified related to exposure to *Salmonella* were diarrhea, enteric fever, enterocolitis and chronic arthritis [83]. For *Bacillus cereus*, emesis (nausea and vomiting), diarrhea, fulminant liver failure and death were identified as health effects [84–86]. For chemical contaminants, health effect selection was based on critical effects identified for the establishment of health-based guidance values, i.e., the no/lowest-observed adverse effect levels or benchmark dose levels [54].

To perform a risk–benefit analysis investigating the substitution of IC and BC products, differences in nutritional composition between these types of products must be evaluated. A comparison of the nutritional content of IC and BC revealed that only dietary fiber and sodium content seems to differ between the two.

## Appendix B. Fiber Intake and Risk Ratio Estimates

Fiber intake was categorized in four different classes. Table S2 summarizes median fiber intakes in different classes and scenarios, and the dose–response data (risk ratios) for DM2 and CVD for each of those classes. From the lowest to the highest classes, fiber intake increased considerably, reaching almost double in the highest class when compared to the lowest. Table S2 also presented risk ratio estimates for DM2 and CVD according to different classes of fiber intake. For DM2 and CVD the RR estimates decreased from the lowest to the highest classes, as fiber exposure increased. For DM2, the lowest intake of fiber (Class 1) for the Worst IC scenario presented the highest RR estimated (0.891, 95% CI: 0.866–0.917). On the other hand, the lowest RR was estimated for the highest-class exposure (Class 4) of the Best BC scenario (0.732, 95% CI: 0.677–0.790). The same trend was observed for CVD, with a highest RR of 0.840 (95% CI: 0.761–0.924) and a lowest of 0.626 (95% CI: 0.476–0.807).

## References

1. Amezdroz, E.; Carpenter, L.; O’Callaghan, E.; Johnson, S.; Waters, E. Transition from milks to the introduction of solid foods across the first 2 years of life: Findings from an Australian birth cohort study. *J. Hum. Nutr. Diet.* **2015**, *28*, 375–383. [\[CrossRef\]](#)
2. Rodrigues, S.S.P.; Lopes, C.; Naska, A.; Trichopoulou, A.; de Almeida, M.D.V. Comparison of national food supply, household food availability and individual food consumption data in Portugal. *J. Public Health* **2007**, *15*, 447–455. [\[CrossRef\]](#)
3. Collins, H.M.; Burton, R.A.; Topping, D.L.; Liao, M.-L.; Bacic, A.; Fincher, G.B. REVIEW: Variability in Fine Structures of Noncellulosic Cell Wall Polysaccharides from Cereal Grains: Potential Importance in Human Health and Nutrition. *Cereal Chem.* **2010**, *87*, 272–282. [\[CrossRef\]](#)
4. Schwartz, M.B.; Vartanian, L.R.; Wharton, C.M.; Brownell, K.D. Examining the nutritional quality of breakfast cereals marketed to children. *J. Am. Diet. Assoc.* **2008**, *108*, 702–705. [\[CrossRef\]](#) [\[PubMed\]](#)
5. Assunção, R.; Vasco, E.; Nunes, B.; Loureiro, S.; Martins, C.; Alvito, P. Single-compound and cumulative risk assessment of mycotoxins present in breakfast cereals consumed by children from Lisbon region, Portugal. *Food Chem. Toxicol.* **2015**, *86*, 274–281. [\[CrossRef\]](#) [\[PubMed\]](#)
6. Assunção, R.; Martins, C.; Vasco, E.; Jager, A.; Oliveira, C.; Cunha, S.C.; Fernandes, J.O.; Nunes, B.; Loureiro, S.; Alvito, P. Portuguese children dietary exposure to multiple mycotoxins—An overview of risk assessment under MYCOMIX project. *Food Chem. Toxicol.* **2018**, *118*, 399–408. [\[CrossRef\]](#)
7. Martins, C.; Assunção, R.; Cunha, S.C.; Fernandes, J.O.; Jager, A.; Petta, T.; Oliveira, C.A.; Alvito, P. Assessment of multiple mycotoxins in breakfast cereals available in the Portuguese market. *Food Chem.* **2018**, *239*, 132–140. [\[CrossRef\]](#)
8. Zhang, K.; Flannery, B.M.; Oles, C.J.; Adeuya, A. Mycotoxins in infant/toddler foods and breakfast cereals in the U.S. retail market. *Food Addit. Contam. Part B* **2018**, *11*, 183–190. [\[CrossRef\]](#)



9. Gardener, H.; Bowen, J.; Callan, S.P. Lead and cadmium contamination in a large sample of United States infant formulas and baby foods. *Sci. Total Environ.* **2019**, *651*, 822–827. [[CrossRef](#)]
10. Jean, J.; Sirot, V.; Hulin, M.; Le Calvez, E.; Zinck, J.; Noël, L.; Vasseur, P.; Nesslany, F.; Gorecki, S.; Guérin, T.; et al. Dietary exposure to cadmium and health risk assessment in children—Results of the French infant total diet study. *Food Chem. Toxicol.* **2018**, *115*, 358–364. [[CrossRef](#)]
11. Guérin, T.; Chekri, R.; Chafey, C.; Testu, C.; Hulin, M.; Noël, L. Mercury in foods from the first French total diet study on infants and toddlers. *Food Chem.* **2018**, *239*, 920–925. [[CrossRef](#)]
12. Tinggi, U.; Schoendorfer, N. Analysis of lead and cadmium in cereal products and duplicate diets of a small group of selected Brisbane children for estimation of daily metal exposure. *J. Trace Elem. Med. Biol.* **2018**, *50*, 671–675. [[CrossRef](#)]
13. Martins, C.; Vasco, E.; Paixão, E.; Alvito, P. Total mercury in infant food, occurrence and exposure assessment in Portugal. *Food Addit. Contam. Part B* **2013**, *6*, 151–157. [[CrossRef](#)] [[PubMed](#)]
14. Lambert, M.; Inthavong, C.; Hommet, F.; Leblanc, J.-C.; Hulin, M.; Guérin, T. Levels of acrylamide in foods included in ‘the first French total diet study on infants and toddlers. *Food Chem.* **2018**, *240*, 997–1004. [[CrossRef](#)] [[PubMed](#)]
15. Lambert, M.; Inthavong, C.; Desbourdes, C.; Hommet, F.; Sirot, V.; Leblanc, J.-C.; Hulin, M.; Guérin, T. Levels of furan in foods from the first French Total Diet Study on infants and toddlers. *Food Chem.* **2018**, *266*, 381–388. [[CrossRef](#)] [[PubMed](#)]
16. Kacmaz, S.; Zelinkova, Z.; Wenzl, T. Rapid and sensitive method for the determination of four EU marker polycyclic aromatic hydrocarbons in cereal-based foods using isotope-dilution GC/MS. *Food Addit. Contam.—Part A Chem. Anal. Control. Expo. Risk Assess.* **2016**, *33*, 631–638. [[CrossRef](#)]
17. Food and Agriculture Organization of the United Nations (FAO); World Health Organization (WHO). *Ranking of Low Moisture Foods in Support of Microbiological Risk Management—Preliminary Report*; FAO: Rome, Italy, 2014.
18. World Health Organization (WHO). *Guideline: Sodium Intake for Adults and Children*; World Health Organization (WHO): Geneva, Switzerland, 2012; ISBN 9789241504836.
19. World Health Organization (WHO). *Guideline: Sugars Intake for Adults and Children*; World Health Organization (WHO): Geneva, Switzerland, 2015; ISBN 9789241549028.
20. Lopes, C.; Torres, D.; Oliveira, A.; Severo, M.; Alarcão, V.; Guiomar, S.; Mota, J.; Teixeira, P.; Rodrigues, S.; Lobato, L.; et al. *Inquérito Alimentar Nacional e de Atividade Física, IAN-AF 2015–2016: Relatório de Resultados*; University of Porto: Porto, Portugal, 2017; ISBN 978989746181.
21. Lopes, C.; Torres, D.; Oliveira, A.; Severo, M.; Guiomar, S.; Alarcão, V.; Vilela, S.; Ramos, E.; Rodrigues, S.; Oliveira, L.; et al. National Food, Nutrition and Physical Activity Survey of the Portuguese general population. *EFSA Support. Publ.* **2017**, *14*. [[CrossRef](#)]
22. Lopes, C.; Torres, D.; Oliveira, A.; Severo, M.; Guiomar, S.; Alarcão, V.; Ramos, E.; Rodrigues, S.; Vilela, S.; Oliveira, L.; et al. National Food, Nutrition, and Physical Activity Survey of the Portuguese General Population (2015–2016): Protocol for Design and Development. *JMIR Res. Protoc.* **2018**, *7*, e42. [[CrossRef](#)]
23. Alvito, P.C.; Sizoo, E.A.; Almeida, C.M.M.; van Egmond, H.P. Occurrence of aflatoxins and ochratoxin A in baby foods in Portugal. *Food Anal. Methods* **2010**, *3*, 22–30. [[CrossRef](#)]
24. Assunção, R.; Martins, C.; Dupont, D.; Alvito, P. Patulin and ochratoxin A co-occurrence and their bioaccessibility in processed cereal-based foods: A contribution for Portuguese children risk assessment. *Food Chem. Toxicol.* **2016**, *96*, 205–214. [[CrossRef](#)]
25. Food and Agriculture Organization of the United Nations (FAO); World Health Organization (WHO). *Energy and Protein Requirements*; WHO: Geneva, Switzerland, 1985; ISBN 9241207248.
26. Assunção, R.; Alvito, P.; Brazão, R.; Carmona, P.; Fernandes, P.; Jakobsen, L.S.; Lopes, C.; Martins, C.; Membré, J.M.; Monteiro, S.; et al. Building capacity in risk-benefit assessment of foods: Lessons learned from the RB4EU project. *Trends Food Sci. Technol.* **2019**, *91*, 541–548. [[CrossRef](#)]
27. Nauta, M.J.; Andersen, R.; Pilegaard, K.; Pires, S.M.; Ravn-Haren, G.; Tetens, I.; Poulsen, M. Meeting the challenges in the development of risk-benefit assessment of foods. *Trends Food Sci. Technol.* **2018**, *76*, 90–100. [[CrossRef](#)]
28. Pires, S.M.; Boué, G.; Boobis, A.; Eneroth, H.; Hoekstra, J.; Membré, J.M.; Persson, I.M.; Poulsen, M.; Ruzante, J.; van Klaveren, J.; et al. Risk Benefit Assessment of foods: Key findings from an international workshop. *Food Res. Int.* **2019**, *116*, 859–869. [[CrossRef](#)]
29. EFSA Scientific Committee Guidance on human health risk-benefit assessment of foods. *EFSA J.* **2010**, *8*, 1673. [[CrossRef](#)]
30. Verhagen, H.; Tijhuis, M.J.; Gunnlaugsdóttir, H.; Kalogeras, N.; Leino, O.; Luteijn, J.M.; Magnússon, S.H.; Odekerken, G.; Pohjola, M.V.; Tuomisto, J.T.; et al. State of the art in benefit–risk analysis: Introduction. *Food Chem. Toxicol.* **2012**, *50*, 2–4. [[CrossRef](#)]
31. Alvito, P.; Brazão, R.; Carmona, P.; Carvalho, C.; Correia, D.; Fernandes, P.; Jakobsen, L.S.; Lopes, C.; Martins, C.; Membré, J.; et al. RiskBenefit4EU—Partnering to strengthen Risk-Benefit Assessment within the EU using a holistic approach. *EFSA Support. Publ.* **2019**, *16*, 1768E. [[CrossRef](#)]
32. Tijhuis, M.J.; de Jong, N.; Pohjola, M.V.; Gunnlaugsdóttir, H.; Hendriksen, M.; Hoekstra, J.; Holm, F.; Kalogeras, N.; Leino, O.; van Leeuwen, F.X.R.; et al. State of the art in benefit—Risk analysis: Food and nutrition. *Food Chem. Toxicol.* **2012**, *50*, 5–25. [[CrossRef](#)] [[PubMed](#)]
33. Wu, F.; Groopman, J.D.; Pestka, J.J. Public Health Impacts of Foodborne Mycotoxins. *Annu. Rev. Food Sci. Technol.* **2014**, *5*, 351–372. [[CrossRef](#)] [[PubMed](#)]
34. International Agency for Research on Cancer (IARC). *A Review of Human Carcinogens: Chemical Agents and Related Occupations*; IARC Publications: Lyon, France, 2012; Volume 100F, ISBN 978 92 832 1323 9.

35. International Agency for Research on Cancer (IARC). Cancer Fact Sheets: Liver Cancer. Available online: <https://gco.iarc.fr/today/data/factsheets/cancers/11-Liver-fact-sheet.pdf> (accessed on 22 June 2020).
36. Duc, L.H.; Dong, T.C.; Logan, N.A.; Sutherland, A.D.; Taylor, J.; Cutting, S.M. Cases of emesis associated with bacterial contamination of an infant breakfast cereal product. *Int. J. Food Microbiol.* **2005**, *102*, 245–251. [[CrossRef](#)] [[PubMed](#)]
37. Furtado, R.; Belo Correia, C.; Alvito, P. Contaminantes de Origem Microbiológica em Alimentação Infantil. *Bol. EPI-DEMIOLÓGICO Obs.* 2013, Volume 2, pp. 6–7. Available online: [http://www2.insa.pt/sites/INSA/Portugues/PublicacoesRepositorio/Documents/observacoesN62013\\_artigo2.pdf](http://www2.insa.pt/sites/INSA/Portugues/PublicacoesRepositorio/Documents/observacoesN62013_artigo2.pdf) (accessed on 1 September 2021).
38. Lesley, M.B.; Velnetti, L.; Yousr, A.N.; Kasing, A.; Samuel, L. Presence of *Bacillus cereus* s.l. From ready-to-eat cereals (RTE) products in Sarawak. *Int. Food Res. J.* **2013**, *20*, 1031–1034.
39. Reynolds, A.; Mann, J.; Cummings, J.; Winter, N.; Mete, E.; Te Morenga, L. Carbohydrate quality and human health: A series of systematic reviews and meta-analyses. *Lancet* **2019**, *393*, 434–445. [[CrossRef](#)]
40. European Food Safety Authority (EFSA). Panel on Dietetic Products Nutrition and Allergies Scientific Opinion on Dietary Reference Values for carbohydrates and dietary fibre. *EFSA J.* **2010**, *8*, 1462. [[CrossRef](#)]
41. Scientific Advisory Committee on Nutrition (SACN). *Carbohydrates and Health*; TSO: London, UK, 2015; ISBN 978 0 11 708284 7.
42. World Cancer Research Fund and the American Institute for Cancer Research. Diet, Nutrition, Physical Activity and Colorectal Cancer. Continuous Update Project Expert Report 2018. Available online: <https://www.wcrf.org/diet-and-cancer/> (accessed on 22 June 2020).
43. Hauner, H.; Bechthold, A.; Boeing, H.; Brönstrup, A.; Buyken, A.; Leschik-Bonnet, E.; Linseisen, J.; Schulze, M.; Strohm, D.; Wolfram, G. Evidence-Based Guideline of the German Nutrition Society: Carbohydrate Intake and Prevention of Nutrition-Related Diseases. *Ann. Nutr. Metab.* **2012**, *60*, 1–58. [[CrossRef](#)] [[PubMed](#)]
44. WHO. Joint WHO/FAO Expert Consultation on Diet Nutrition and the Prevention of Chronic Diseases. In *Diet, Nutrition and Prevention of Chronic Diseases: Report of a Joint WHO/FAO Expert Consultation*; WHO: Geneva, Switzerland, 2003.
45. Anderson, J.W.; Baird, P.; Davis, R.H., Jr.; Ferreri, S.; Knudtson, M.; Koraym, A.; Waters, V.; Williams, C.L. Health benefits of dietary fiber. *Nutr. Rev.* **2009**, *67*, 188–205. [[CrossRef](#)]
46. Turck, D.; Castenmiller, J.; de Henauw, S.; Hirsch-Ernst, K.; Kearney, J.; Knutsen, H.K.; Maciuk, A.; Mangelsdorf, I.; McArdle, H.J.; Pelaez, C.; et al. Dietary reference values for sodium. *EFSA J.* **2019**, *17*, e05778. [[CrossRef](#)] [[PubMed](#)]
47. European Food Safety Authority. Guidance on the EU Menu methodology. *EFSA J.* **2014**, *12*, 3944. [[CrossRef](#)]
48. Instituto Nacional de Saúde Doutor Ricardo Jorge Tabela da Composição dos Alimentos (Foodcomposition Database). Available online: <http://www2.insa.pt/sites/INSA/Portugues/AreasCientificas/AlimentNutricao/AplicacoesOnline/TabelaAlimentos/Paginas/TabelaAlimentos.aspx> (accessed on 15 June 2020).
49. European Commission. Commission Regulation (EC) No 1881/2006 of 19 December 2006 setting maximum levels for certain contaminants in foodstuffs. *Off. J. Eur. Union* **2006**, *364*, 5–24.
50. ASAE Plano Nacional de Colheita de Amostras. Available online: <https://www.asae.gov.pt/> (accessed on 22 June 2020).
51. Devleesschauwer, B.; Haagsma, J.A.; Angulo, F.; Bellinger, D.C.; Cole, D.; Dopfer, D.; Aamir, F.; Fevre, E.M.; Gibb, H.; Hald, T.; et al. Methodological Framework for World Health Organization Estimates of the Global Burden of Foodborne Disease. *PLoS ONE* **2015**, *10*, e0142498. [[CrossRef](#)]
52. Thomsen, S.T.; Pires, S.M.; Devleesschauwer, B.; Poulsen, M.; Fagt, S.; Ygil, K.H.; Andersen, R. Investigating the risk-benefit balance of substituting red and processed meat with fish in a Danish diet. *Food Chem. Toxicol.* **2018**, *120*, 50–63. [[CrossRef](#)] [[PubMed](#)]
53. Institute for Health Metrics and Evaluation (IHME). GBD Results Tool 2017. Available online: <http://ghdx.healthdata.org/gbd-results-tool> (accessed on 22 June 2020).
54. FAO/WHO Expert Committee on Food Additives. *Evaluation of Certain Contaminants in Food: Eighty-Third Report of the Joint FAO/WHO Expert Committee on Food Additives*; WHO: Geneva, Switzerland, 2017.
55. Dekkers, A.L.; Verkaik-Kloosterman, J.; van Rossum, C.T.; Ocké, M.C. SPADE, a New Statistical Program to Estimate Habitual Dietary Intake from Multiple Food Sources and Dietary Supplements. *J. Nutr.* **2014**, *144*, 2083–2091. [[CrossRef](#)]
56. Berlin, J.A.; Longnecker, M.P.; Epidemiology, S.; May, N.; Berlin, J.A.; Longnecker, M.P.; Greenland, S. Meta-analysis of Epidemiologic Dose-Response Data. *Epidemiology* **1993**, *4*, 218–228. [[CrossRef](#)] [[PubMed](#)]
57. Barendregt, J.J.; Veerman, J.L. Categorical versus continuous risk factors and the calculation of potential impact fractions. *J. Epidemiol. Community Health* **2010**, *64*, 209–212. [[CrossRef](#)] [[PubMed](#)]
58. National Academies of Sciences. *Dietary Reference Intakes for Sodium and Potassium*; The National Academies Press: Washington, DC, USA, 2019; ISBN 9780309488341.
59. European Food Safety Authority (EFSA). Opinion of the scientific panel on contaminants in the food chain [CONTAM] related to the potential increase of consumer health risk by a possible increase of the existing maximum levels for aflatoxins in almonds, hazelnuts and pistachios and derived prod. *EFSA J.* **2007**, *5*, 446. [[CrossRef](#)]
60. European Food Safety Authority (EFSA). Management of left-censored data in dietary exposure assessment of chemical substances. *EFSA J.* **2010**, *8*, 1–96. [[CrossRef](#)]
61. European Food Safety Authority (EFSA). Scientific opinion on the risks for public health related to the presence of *Bacillus cereus* and other *Bacillus* spp. including *Bacillus thuringiensis* in foodstuffs. *EFSA J.* **2016**, *14*. [[CrossRef](#)]

62. Farakos, S.S.; Pouillot, R.; Spungen, J.; Flannery, B.; Dolan, L.; Doren, J. Van Implementing a Framework to Evaluate the Impact of Food Intake Shifts on Risk of Illness Using a Case Study with Infant Cereal (P11-118-19). *Curr. Dev. Nutr.* **2019**, *3*, 1096. [CrossRef]
63. Fraeyman, S.; Croubels, S.; Devreese, M.; Antonissen, G. Emerging Fusarium and Alternaria Mycotoxins: Occurrence, Toxicity and Toxicokinetics. *Toxins* **2017**, *9*, 228. [CrossRef]
64. Eskola, M.; Kos, G.; Elliott, C.T.; Hajšlová, J.; Mayar, S.; Krska, R. Worldwide contamination of food-crops with mycotoxins: Validity of the widely cited 'FAO estimate' of 25%. *Crit. Rev. Food Sci. Nutr.* **2020**, *60*, 2773–2789. [CrossRef]
65. Angelino, D.; Rosi, A.; Dall'Asta, M.; Pellegrini, N.; Martini, D. Evaluation of the Nutritional Quality of Breakfast Cereals Sold on the Italian Market: The Food Labelling of Italian Products (FLIP) Study. *Nutrients* **2019**, *11*, 2827. [CrossRef]
66. Italian Ministry of Health. Common Objectives for the Improvement of the Nutritional Characteristics of Food Products with a Special Focus on Children (Ages 3–12) 2017. Monitoring Report. Available online: [https://www.salute.gov.it/imgs/C\\_17\\_opuscoliPoster\\_376\\_0\\_alleg.pdf](https://www.salute.gov.it/imgs/C_17_opuscoliPoster_376_0_alleg.pdf) (accessed on 15 June 2020).
67. Ministério das Finanças; Ministério da Administração Interna; Ministério da Educação; Ministério da Saúde; Ministério da Economia; Ministério da Agricultura Florestas e Desenvolvimento Rural; Ministério do Mar. Despacho N° 11418/2017 Estratégia integrada para a promoção da alimentação saudável. *Diário da República* **2017**, *249*, 29595–29598.
68. Freitas-Silva, O.; Venâncio, A. Brazil nuts: Benefits and risks associated with contamination by fungi and mycotoxins. *Food Res. Int.* **2011**, *44*, 1434–1440. [CrossRef]
69. Nugent, A.P.; Thielecke, F. Wholegrains and health: Many benefits but do contaminants pose any risk? *Nutr. Bull.* **2019**, *44*, 107–115. [CrossRef]
70. European Food Safety Authority (EFSA). Panel on Dietetic Products Nutrition and Allergies Scientific Opinion on Dietary Reference Values for fats, including saturated fatty acids, polyunsaturated fatty acids, monounsaturated fatty acids, trans fatty acids, and cholesterol. *EFSA J.* **2016**, *8*, 1461. [CrossRef]
71. Abushelaibi, A.A.; Sofos, J.N.; Samelis, J.; Kendall, P.A. Survival and growth of Salmonella in reconstituted infant cereal hydrated with water, milk or apple juice and stored at 4 °C, 15 °C and 25 °C. *Food Microbiol.* **2003**, *20*, 17–25. [CrossRef]
72. World Health Organization (WHO). *Enterobacter Sakazakii and Salmonella in Powdered Infant Formula: Meeting Report*; WHO: Geneva, Switzerland, 2006; ISBN 9241563311.
73. Lee, H.-Y.; Chai, L.-C.; Tang, S.-Y.; Jinap, S.; Ghazali, F.M.; Nakaguchi, Y.; Nishibuchi, M.; Son, R. Application of MPN-PCR in biosafety of Bacillus cereus s.l. for ready-to-eat cereals. *Food Control* **2009**, *20*, 1068–1071. [CrossRef]
74. Young, I.; Waddell, L.; Cahill, S.; Kojima, M.; Clarke, R.; Rajic, A. Application of a Rapid Knowledge Synthesis and Transfer Approach To Assess the Microbial Safety of Low-Moisture Foods. *J. Food Prot.* **2015**, *78*, 2264–2278. [CrossRef] [PubMed]
75. Sirot, V.; Traore, T.; Guérin, T.; Noël, L.; Bachelot, M.; Cravedi, J.-P.; Mazur, A.; Glorennec, P.; Vasseur, P.; Jean, J.; et al. French infant total diet study: Exposure to selected trace elements and associated health risks. *Food Chem. Toxicol.* **2018**, *120*, 625–633. [CrossRef] [PubMed]
76. Vos, M.B.; Kaar, J.L.; Welsh, J.A.; Van Horn, L.V.; Feig, D.I.; Anderson, C.A.M.; Patel, M.J.; Cruz Munos, J.; Krebs, N.F.; Xanthakos, S.A.; et al. Added Sugars and Cardiovascular Disease Risk in Children: A Scientific Statement From the American Heart Association. *Circulation* **2017**, *135*, e1017–e1034. [CrossRef]
77. He, F.J.; Marrero, N.M.; MacGregor, G.A. Salt and blood pressure in children and adolescents. *J. Hum. Hypertens.* **2008**, *22*, 4–11. [CrossRef]
78. Kranz, S.; Brauchla, M.; Slavin, J.L.; Miller, K.B. What Do We Know about Dietary Fiber Intake in Children and Health? The Effects of Fiber Intake on Constipation, Obesity, and Diabetes in Children. *Adv. Nutr.* **2012**, *3*, 47–53. [CrossRef]
79. Ma, Y.; He, F.J.; Macgregor, G.A. High salt intake: Independent risk factor for obesity? *Hypertension* **2015**, *66*, 843–849. [CrossRef]
80. Te Morenga, L.; Montez, J.M. Health effects of saturated and trans-fatty acid intake in children and adolescents: Systematic review and meta-analysis. *PLoS ONE* **2017**, *12*, e0186672. [CrossRef] [PubMed]
81. D'Elia, L.; Rossi, G.; Ippolito, R.; Cappuccio, F.P.; Strazzullo, P. Habitual salt intake and risk of gastric cancer: A meta-analysis of prospective studies. *Clin. Nutr.* **2012**, *31*, 489–498. [CrossRef] [PubMed]
82. Aburto, N.J.; Ziolkovska, A.; Hooper, L.; Elliott, P.; Cappuccio, F.P.; Meerpohl, J.J. Effect of lower sodium intake on health: Systematic review and meta-analyses. *BMJ* **2013**, *346*, f1326. [CrossRef] [PubMed]
83. Leirisalo-Repo, M.; Helenius, P.; Hannu, T.; Lehtinen, A.; Kreula, J.; Taavitsainen, M.; Koskimies, S. Long term prognosis of reactive salmonella arthritis. *Ann. Rheum. Dis.* **1997**, *56*, 516–520. [CrossRef]
84. Ehling-Schulz, M.; Fricker, M.; Scherer, S. Bacillus cereus, the causative agent of an emetic type of food-borne illness. *Mol. Nutr. Food Res.* **2004**, *48*, 479–487. [CrossRef]
85. Stenfors Arnesen, L.P.; Fagerlund, A.; Granum, P.E. From soil to gut: Bacillus cereus and its food poisoning toxins. *FEMS Microbiol. Rev.* **2008**, *32*, 579–606. [CrossRef]
86. Mahler, H.; Pasi, A.; Kramer, J.M.; Schulte, P.; Scoging, A.C.; Bär, W.; Krähenbühl, S. Fulminant Liver Failure in Association with the Emetic Toxin of Bacillus cereus. *N. Engl. J. Med.* **1997**, *336*, 1142–1148. [CrossRef]