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The Role of Nutritional Aspects in Food Allergy: Prevention and Management

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1 Review

2 The role of nutritional aspects in food allergy: prevention and management

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11 **Abstract:** The prevalence of food allergy in childhood appears to be increasing in both developed
12 and transitional countries. The aim of this paper is to review and summarise key findings in the
13 prevention and management of food allergy focusing on the role of dietary components and
14 nutritional habits in the development and optimal functioning of the immune system. Essential fatty
15 acids, zinc and vitamin D are likely to enhance the anti-inflammatory and antioxidative barrier and
16 promote immunologic tolerance. Additionally nutritional components such as pre and probiotics
17 represent a novel research approach in the attempt to induce a tolerogenic immune environment. For
18 all these reasons, the traditional avoidance diet has been in recent years completely reconsidered.
19 New findings on the protective effect of an increased diversity of food introduced in the first year
20 of life on allergic diseases are consistent with the hypothesis that exposure to a variety of food
21 antigens during early life might play a role in the development of immune tolerance. Accordingly,
22 therapeutic (and even preventive) interventions should be planned on an individual basis.

23 **Keywords:** food allergy; children; diet diversity; adequate nutrition

24

25 1. INTRODUCTION

26 Food allergy (FA) represents a substantial health problem in childhood. The prevalence appears
27 to be increasing in both developed and transitional countries, however a true increase has been
28 difficult to demonstrate [1]. Over 90% of food allergies are caused by eight common allergens;
29 namely: eggs, peanuts, cows' milk, soy, nuts, shellfish, fish, or wheat [2]. On the whole, food allergy
30 affects approximately 6% of infants younger than three years [2], and prevalence decreases over the
31 first decade. The cumulative incidence of food hypersensitivity over a 10-year period is 6.7% (95%
32 CI: 5.20 to 8.4); 3.0% (95% CI: 1.8–4.2%) had IgE-mediated food allergy and 0.6% (95% CI: 0.07–
33 1.3%) had non-IgE-mediated food allergy/food intolerance [3]. A systematic review from the
34 European Academy of Allergy and Clinical Immunology concluded that food allergy prevalence in
35 Europe range between 0.1 to 6.0% [4]. The Institute of Medicine report states that the prevalence of
36 food allergies in children range between 1.1 – 10.4% [1]. Food allergic infants commonly present
37 with symptoms and signs of atopic eczema, gastrointestinal symptoms and/or recurrent wheezing [5].
38 Diet plays a crucial role in both the prevention and management of food allergy. A number of factors
39 including the maternal diet, the microbiome and early life feeding have been investigated for the
40 prevention of allergic diseases [6]. The aim of this paper is to review and summarise key findings in
41 the prevention and management of food allergy, with particular reference to nutrients of concern (fats,
42 micronutrients), gut flora (including the role of pre- and probiotics), early life feeding and formula
43 choice in cows' milk allergy.

44

45 2. PREVENTION OF FOOD ALLERGY: THE ROLE OF NUTRITION IN THE 46 DEVELOPMENT AND OPTIMAL FUNCTIONING OF THE IMMUNE SYSTEM

47 Allergy results when there is a breakdown in normal “tolerance” mechanisms, which leads to
48 inappropriate and detrimental immune responses to normally harmless substances, including food
49 allergens such as cow’s milk protein, eggs, nuts, or shellfish [7]. At birth, the immune system is
50 immature, but it develops with age, antigen stimulation, and appropriate nutrition [8]. In addition,
51 bacterial colonization occurs during the first weeks of life, and interactions between intestinal flora
52 and the developing mucosa result in further development of immune responses and oral tolerance [7].

53 Nutrition plays a key role in the development, maintenance, and optimal functioning of immune
54 cells. Nutrients, such as zinc and vitamin D and nutritional factors, such as pre and probiotics, can
55 influence the nature of an immune response and are important in ensuring appropriate functioning of
56 the immune system, as described in the paragraphs below.

57

58 2.1 FAT

59 Appropriate fat intake may become seriously compromised in allergen-restricted diets and may
60 be further influenced by the “westernized” dietary practices. The role of fat on the immune system
61 can be divided into the role of saturated vs. unsaturated fats and the particular role of the essential
62 fatty acids.

63

64 Saturated vs. unsaturated fats

65 It has been reported that typical western diets rich in protein and saturated fat and low in
66 carbohydrates may negatively effect the diversity of the gut microbiome [9]. This was supported by
67 David et al. [10], showing that an animal based diet high in protein and fat, with very little fibre
68 intake, resulted in increased abundance of bile-tolerant microorganisms (*Alistipes*, *Bilophila*, and
69 *Bacteroides*) and decreased levels of Firmicutes that metabolize dietary plant polysaccharides
70 (*Roseburia*, *Eubacterium rectale*, and *Ruminococcus bromii*) within a five day period. A recent
71 review also concluded that the amount, type (e.g., unsaturated vs saturated), and mixture of dietary
72 fats can dramatically shift gut microbial community membership and function [11]. In addition high
73 fat, high sugar diets also affect the gut barrier function in mice as demonstrated by high horseradish
74 peroxidase (HRP) influx, lower portal vein endotoxin levels and decreased goblet cell numbers [12].
75 The gut barrier function may be permanently affected in non-IgE mediated food allergies and
76 temporarily affected during allergen exposure in IgE mediated food allergies [13, 14].

77 Essential fatty acids (EFAs)

78 EFAs are important immune regulators. Linoleic acid (LA), the parental n-6 polyunsaturated
79 fatty acid (PUFA), is converted into arachidonic acid (AA) by fatty acid elongase and desaturase, and
80 subsequently may give origin to pro-inflammatory and pro-allergic lipid mediators, whose cumulative
81 name is eicosanoids [15]. In contrast, α -linolenic acid (ALA), an n-3 PUFA, is converted in
82 mammalian body to eicosapentaenoic acid (EPA) and docosahexaenoic acid (DHA), which are
83 subsequently converted into anti-inflammatory and/or pro-resolving lipid mediators (such as resolvins
84 and protectins). EPA forms the precursors of the 3 series of prostaglandins and the 5 series of
85 leukotrienes, which are biologically less powerful than the corresponding derivatives which form the
86 n-6 compounds. Because n-3 and n-6 PUFAs compete for the same metabolic pathways, an increase
87 of n-3 PUFA, parallel to a decrease of n-6 PUFA intake, might theoretically reduce the onset of human
88 immunologic conditions, including allergies thanks to the replacement of EPA instead AA in the
89 membranes of inflammatory cells. EFA, including long-chain PUFAs, may be consumed as part of
90 the normal diet through breast milk, formula and food, or as supplements at any stage in the life cycle
91 [15].

92 The fatty acid status is of particular concern in infants and children. Essential fatty acids (EFA)
93 promote the renewal of the protective hydrolipidic film layer of the skin and, accordingly, an altered

94 EFA metabolism has been associated with the pathogenesis of atopic dermatitis (AD). Moreover the
95 clinical spectrum of EFA deficiency may range from mild skin irritation to life-threatening conditions
96 [16].

97 In spite of intensive research in the field, a recent systematic review [17] concerning the role of
98 dietary PUFAs in the development of allergy shows that PUFA supplementation in infancy seems not
99 to affect infant incidence, childhood incidence or childhood prevalence of food allergy (GRADE level
100 of evidence: very low) even taking into account a moderate heterogeneity between studies that
101 reported infant incidence of food allergy (3 studies; 915 infants; RR 0.81, 95% CI 0.56 to 1.19, I²=
102 63%; RD -0.02, 95% CI -0.06 to 0.02, I² = 74%). However, well documented immunomodulatory
103 effects of n-3 PUFAs (both in vitro and in vivo) highlight the potential role in preventing and treating
104 allergic disease but larger longitudinal intervention studies are clearly warranted to confirm this
105 observation [18].
106

107 2.2 ZINC

108 Children with food hypersensitivity have increased amounts of mastocytes, eosinophils and
109 neutrophils in the digestive tract. Persistent exposure to allergen can lead to chronic inflammatory
110 changes of mucous membrane and increased production of reactive oxygen species (ROS) [19].
111 Excess ROS should be neutralized by components of the antioxidative barrier. Therefore all
112 disturbances of enzymatic and non-enzymatic mechanisms of this barrier lead to many unfavourable
113 reactions including oxidation of cell membrane lipids. Zinc is an essential trace element and it is
114 needed for various cellular functions, specifically it is a cofactor of many enzymes including
115 superoxide dismutase (SOD) that play an important role in maintaining the oxidative-antioxidative
116 balance. A study performed in 134 children with food allergy, aged 1 to 36 months, showed that
117 children with food allergy had significantly lower concentrations of zinc and therefore a weakened
118 antioxidative barrier [19]. To our knowledge there are no RCTs investigating zinc supplementation
119 and allergic outcomes.
120

121 2.3 VITAMIN D

122
123 The classical role of Vitamin D is in fact related to calcium homeostasis and bone health.
124 However, over the last decade, the effects of vitamin D on the innate and adaptive immune system
125 have been investigated and expanded [20]. The active form of the vitamin, i.e. 1,25(OH)₂D, has
126 effects on epithelial cells, T cells, B cells, macrophages and dendritic cells. It stimulates innate
127 immune responses by enhancing the chemotactic and phagocytotic responses of macrophages as well
128 as the production of antimicrobial proteins such as cathelicidin. This action plays a role in maintaining
129 the mucosal integrity by stimulating junction genes. Nevertheless, the potential effect of vitamin D
130 on Th1/Th2 adaptive immune response is of interest and related to food allergy [21, 22, 23]. Almost
131 all cells of the adaptive immune system express the vitamin D receptor, making them also capable of
132 being vitamin responsive. When specifically considering a potential role for vitamins in food allergy,
133 vitamin D has been shown to affect several mechanisms that promote immunologic tolerance,
134 including T regulatory cell function and the induction of tolerogenic dendritic cells. However clinical
135 trials on vitamin D supplementation in children and the possible role in preventing food allergy are
136 lacking. A systematic review of vitamin D supplementation for the prevention of allergic diseases
137 found no evidences about the protective role of this nutrient in children, but the currently available
138 data are poor [24].
139

140 2.4 THE ROLE OF PREBIOTICS, PROBIOTICS AND MICROBIOTA IN THE PREVENTION OF 141 FOOD ALLERGY

142 The innate immune system has the ability to modulate adaptive immune responses to food
143 proteins. Therefore, the type of gastrointestinal microbiota of the newborn and the preservation of

144 intestinal permeability is crucial for preventing the development of food allergies. The dietary
145 modulation of nutritional factors through pre, pro- and synbiotic preparations represent a novel
146 research hypothesis and a challenge for dietitians and pediatric allergists. The modulation of the
147 immune system using functional foods is a promising research hypothesis in the attempt to induce a
148 tolerogenic immune environment [16].
149

150 Prebiotics

151 Prebiotics have been defined as “non-digestible food components that beneficially affect the host
152 by selectively stimulating the growth and/or activity of one or a limited number of bacteria in the
153 colon and thereby improving host health” and recently redefined as “a selectively fermented
154 ingredient that allows specific changes, both in the composition and/or activity in the gastrointestinal
155 microbiota that confers benefits” [25]. In December 2016 the panel of expert convened by the
156 International Scientific Association for Probiotics and Prebiotics (ISAPP) suggested a new definition,
157 i.e. “a substrate that is selectively utilized by host microorganisms conferring a health benefit” [26].
158 Based on the body of available evidence, the Guidelines for Atopic Disease Prevention (GLAD-p)
159 panel concluded that it is likely that prebiotic supplementation in infants reduces the risk of
160 developing recurrent wheezing and possibly also the development of food allergy. However there is
161 very low certainty that there is an effect of prebiotics on other outcomes, other than an indirect effect
162 due to it’s effect on the microbiome. In fact, their activity can be affected by many individual factors,
163 (e.g. host’s microbiota or the genetic predisposition to diseases). Also environmental factors such as
164 diet or antibiotics can influence the use of prebiotics [26].
165

166 Probiotics

167 Probiotics are living microorganisms that have been proposed as immune-modulators of the
168 allergic response by affecting phagocytosis and production of pro-inflammatory cytokines, and thus
169 are being advocated as therapeutic and preventive interventions for allergic diseases [27]. They are
170 present in everyday food (not only in yoghurt or fermented milk but also in cheese- either hard or
171 soft, and also in less expected sources such as kefir, miso soup or tempeh) and they are a common
172 exposure in almost everyone's life [27]. The probiotic effects of complex oligosaccharides in human
173 milk promote the establishment of a bifidogenic microbiota which, in turn, induces a milieu of
174 tolerogenic immune responses to foods. Earlier studies suggested a positive effect of probiotic
175 interventions on atopic dermatitis, but meta-analyses have failed to confirm it.

176 The new World Allergy Organization (WAO) guidelines determined that it is likely that probiotic
177 supplementation in infants reduces the risk of developing eczema and suggest that probiotics should
178 be recommended in mothers of high-risk infants and in infants at high risk of allergic disease, where
179 “high risk for allergy in a child” is defined as biological parent or sibling with existing or history of
180 allergic rhinitis, asthma, eczema, or food allergy [27]. The recommendations are conditional and
181 based on very low quality evidence, with no specific recommendation regarding strains, dose,
182 treatment duration etc .

183 In terms of tolerance development in those with established food allergy, one study from
184 Australia performed oral immunotherapy (OIT) to peanut in combination with *Lactabillus GG*,
185 showing that 89.7% of the study participants in this arm were desensitized to peanut. The authors
186 speculate that this protective effect may be seen because of the possible effect of the probiotic on T-
187 regulatory cells [28]. Further scientific confirmations are required to include probiotics and prebiotics
188 in the therapeutic plans. Practical implications and how this should be incorporated in advising food
189 allergy sufferers are also unclear in terms of advising regular intake of foods high in short chain
190 fructo-oligo saccharides, fermented foods and yoghurts.

191 3. THE ROLE OF ALLERGEN INTAKE AND DIETARY DIVERSITY IN PREVENTION 192 OF FOOD ALLERGY

193

194 3.1 ALLERGEN INTAKE

195 Measures to prevent allergy and food allergy have traditionally included maternal allergen
196 avoidance during pregnancy and/or lactation, periods of exclusive breast feeding and avoidance of
197 potential allergens including food and environmental antigens during the first year of life and beyond
198 [29]. The value and significance of food avoidance for preventive purposes has been in recent years
199 completely reconsidered.

200 On the contrary, an ideal age to introduce potentially allergenic foods into an infant's diet has
201 been debated for the past 2 decades, particularly in high-income countries where allergic disease has
202 become highly prevalent. Initial approaches to primary prevention of food allergy largely focused on
203 "avoidance" strategies. In 2000 [30], practice guidelines generally recommended that allergenic
204 foods (such as egg, cow's milk, and peanut) be avoided during the first 1 to 3 years of life. As data
205 accumulated from both observational studies and experimental models, it became apparent that
206 avoidance practices may not be beneficial.

207 Given the increasing interest on the role of time of introduction of allergic food into the infant
208 diet (the so called "window of opportunity") and the risk of allergic diseases, intervention trials
209 evaluating the intake of food, as milk, egg, peanuts etc, during the first year of life have been
210 performed.

211 For instance, a recent RCT found no evidence that regular egg intake from age 4 to 6.5 months
212 substantially alters the risk of egg allergy by age 1 year in infants who are at hereditary risk of allergic
213 disease and had no eczema symptoms at study entry [31]. These findings are generally supportive
214 of other data in high-risk patients showing a risk-reducing benefit of early egg introduction, and risk-
215 reducing benefit for early peanut introduction [32]. The EAT study [33] also showed a reduced risk
216 in the general population using the per protocol analysis but not intention to treat analysis. For peanut,
217 clinical practice guidelines in the US have incorporated these findings and do recommend early
218 peanut introduction in the first year of life for high and standard risk children [34]. However, despite
219 some evidence for early introduction of egg, the US guidelines only made recommendations regarding
220 peanut intake and concluded that there was not enough evidence to suggest early introduction of egg.
221 Surprisingly, the UK COT report [35] published very recently, suggested that all foods should be
222 introduced after a period of exclusive breast feeding from 6 months and that there is no need to
223 introduce peanut or egg differently from other foods. It seems as if despite the data from recent RCTs
224 on peanut and egg the weaning debate will continue, as there is still no consensus about the age of
225 introduction of these foods. The only consistent messages are: Start weaning once the infant is
226 developmentally ready; Don't delay introduction of allergens: once they are introduced into the diet,
227 continue to feed them.

228

229 3.2 DIET DIVERSITY AND OTHER RELATED FACTORS

230

231 Dietary diversity

232 Recent findings on the protective effect of an increased diversity of food introduced in the first
233 year of life on allergic diseases (asthma, atopic dermatitis, food allergy and atopic sensitization) are
234 consistent with the hypothesis that exposure to a variety of food antigens during early life might be
235 important for the development of immune tolerance [36-38].

236 The microbiome plays an important role in ensuring the gut wall integrity and regulation of the
237 immune system. Diet diversity has been shown to reduce allergic diseases [39, 40]. This may well be
238 that the more diverse diet leads to a more diverse microbiome [41] and that natural microbial load of
239 food enhances this process [42]. This in turn may improve the gut wall integrity and regulation of the
240 immune system, but human trials are needed to confirm this theory.

241

242 Food production

243 Food production and cooking methods may also affect allergy the immune system (perhaps) via
244 its effect on the microbiome. Lang et al. [43] reported that the microbial load of different diets (e.g.
245 USA diet vs. vegan diet) differs due to the foods excluded and cooking methods used. Chaturvedi, et
246 al. [44] reported that the natural microbial load of fruits and vegetables differ between groups from a
247 different socio-economic status. In addition Venter and Maslin reported an association seen between
248 increase in baby food sales and allergic diseases [45], underlining that commercial baby foods are
249 sterile and that the diversity of ingredients and nutrient content is variable. All these factors highlight
250 that the foods we eat (irrespective of their nutrient content) may affect the immune system and perhaps
251 development and management of allergic diseases.

252 Healthy diet

253 It is unclear at present what a “healthy diet” in terms of allergy prevention and management means
254 and if a healthy diet as we know it (20% protein, 50% carbohydrate, 30% fat) has any relevance in
255 allergy prevention. Currently either the healthy eating index [46] or a mediterranean style diet [47] is
256 being used as a proxy measure for healthy eating. Research using the healthy eating index tool,
257 specific to the pregnancy diet, found no association between overall healthy eating score and recurrent
258 wheeze in infants at the age of 3 years [46] and this was confirmed in a another study by Moonesinghe
259 et al. focusing on eating patterns in pregnancy and allergic diseases [48]. In addition to these two
260 studies, two review papers addressed the issue of the mediterranean diet on allergy prevention. Venter
261 et al. summarised studies during pregnancy [49]. Three observational studies have investigated the
262 role of the Mediterranean diet on allergy outcomes. One study showed a possible increased risk for
263 the infant to develop allergic disease [50], one showed a reduction in wheeze [51], and another study
264 showed no effect on allergy prevention [52]. Mediterranean style eating patterns shows more
265 promising effects with reduction in asthma/wheezing symptoms seen but no effect on other allergic
266 symptoms [47]. More studies are therefore needed with well-defined criteria for healthy eating to
267 study its effect on allergy prevention.

268

269 Other factors

270 More recently the role of advanced glycosylated end products in food and the direct effect on the
271 Th2 immune system and the microbiome has been described [53]. One mouse model study also
272 questioned the role of emulsifiers on the gut microbiome. This study showed that a diet high in
273 emulsifiers destroyed the epithelial mucous layer in the gut, altered gut microbial composition and
274 promoted inflammation [54].

275 4. THE ROLE OF DIET IN THE MANAGEMENT OF FOOD ALLERGY

276 The cornerstone of the nutritional management of food allergies is an individualized allergen
277 avoidance management plan. In children, the main goals are to prevent the occurrence of acute and
278 chronic symptoms by avoiding the offending food(s), whilst providing an adequate, healthy and
279 nutritionally balanced diet and maintaining optimal growth; ideally under the guidance of a trained
280 dietitian [55]. Complete avoidance of the allergen is still required by some, but latest developments
281 in food allergy have indicated that some individuals with food allergies tolerate baked forms of milk
282 and egg [56]. Additionally, complete avoidance of all nuts is not necessarily recommended any more,
283 and only those nuts reactive to, should be eliminated from the diet [57]. In addition to nutritional
284 consequences of food allergy, it is known that children and families with food allergies experience a
285 decreased quality of life across a number of domains, which can create anxiety and lead to avoidance
286 of social situations [58-61]. Hence it is suggested that liberalization of the diet, when appropriate and
287 safe, will increase both quality of life and nutritional intake.

288

289 4.1 COW'S MILK ALLERGY

290 Exclusion of any food group can result in a nutritionally deficient diet, but the elimination of
291 milk and products in infancy is particularly likely to cause nutritional deficiencies [62] and deserves
292 special emphasis. Cow's milk proteins (CM) are among the first foods introduced into an infant's diet
293 and accordingly they represent one of the first and most common causes of food allergy in early
294 childhood. Cows' milk allergy generally requires a strict exclusion diet usually for the first year of
295 life. This exclusion of a main food group occurs at a critical time in the development of food
296 preferences and eating habits. The management of CMA in infants and young children requires
297 individualized advice regarding avoidance of cows' milk, including advice to breastfeeding mothers
298 and/or guidance on the most appropriate specialized formula or milk substitute [63]. In many cases
299 micronutrient supplements will also be required, however their usage is not always intuitive with both
300 under and over supplementation occurring [64].

301 Cow's milk proteins could induce an allergic reaction: in particular beta-lactoglobulin (BLG),
302 included in the whey fraction, is not present in human milk and therefore is considered the principal
303 component involved in the etiology of the disease. During the production of infant formula, only the
304 processes of extensive hydrolysis, ultrafiltration or an enzymatic cleavage result in a truly
305 hypoallergenic formulas [16].
306

307 Choice of formula in CMA

308 The nutritional value of a milk substitute must be taken into account at ages lower than 2 years
309 of life when such a type of food is needed [16]. As breast milk composition differs both in component
310 ratios and structure from other milks, the composition of infant formula should serve to meet the
311 particular nutritional requirements and to promote normal growth and development of the infants for
312 whom they are intended [65, 66]. When a replacement formula is needed, allergologists can avail
313 themselves with different types of formula [67]. The alternative formulas considered for CMA are
314 extensively hydrolyzed whey or casein formula (eHWF or eHCF) and amino acid-based formula
315 (AAF), which are considered of low antigenic potential and are therefore preferred in highly allergic
316 children. The unpalatable taste of hydrolyzed formulas has often been associated with reduced intakes
317 and a consequent growth faltering in infants fed these types of formula, particularly in the first year
318 of life [62].

319 In recent years, an alternative explanation has been proposed based on the content of free amino
320 acids (FAAs) in hydrolyzed formulas, added to complete their biologic value. Glutamic acid, in
321 particular, has been suggested to downregulate appetite during feeding by interacting with specific
322 receptors in the oral cavity and gastrointestinal tract. However recent studies have shown no negative
323 effect of feeding AA formulas in infants, in contrary, they may be beneficial for growth [68].
324

325 Other studies have demonstrated that dietary management with extensively hydrolysed casein-
326 based formula (EHCF) supplemented with the probiotic *Lactobacillus rhamnosus* GG (LGG) results
327 in a higher rate of tolerance acquisition in infants with CMA than in those treated with EHCF without
328 supplementation or with other noncasein-based formulas. The mechanistic basis for this effect could
329 be the possible influence of EHCF+LGG on the strain-level bacterial community structure of the
330 infant gut [69]. However, randomised controlled trials to date have not yielded sufficient evidence to
331 recommend probiotics for the primary prevention of allergic disorders. Indeed, the Nutrition
332 Committee of the European Society for Paediatric Gastroenterology Hepatology and Nutrition
333 (ESPGHAN) does not support routine supplementing with probiotics in infant formulas [70].
334

335 Soy protein-based formula may be an option in infants older than 6 months who do not accept
336 the bitter taste of an eHF, or in cases in which the higher cost of an eHF is a limiting factor [71].
337 However, soy formulae have nutritional disadvantages because their absorption of minerals and trace
338 elements may be lower because of their phytate content, and they contain appreciable amounts of
339 isoflavones with a potentially weak estrogenic action that can lead to high serum concentrations in
340 infants. Also the possible derivation from genetic modified soy should be considered. Hence, the
341 European Society of Pediatric Gastroenterology, Hepatology, and Nutrition (ESPGHAN) and the
342 American Academy of Pediatrics (AAP) recommend that cow's-milk-based formulae should be

343 preferred over soy formula in healthy infants, and soy protein– based formulae should not usually be
344 used during the first 6 months of life [71].

345

346 Other mammal's milks, those of, goats, ewe's, mare, donkey, or camel have been proposed as
347 substitutes in the management of CMA in infants and children, but are NOT recommended due to
348 either nutritional issues, cross-reactions or both. The DRACMA guidelines state that milk allergens
349 of various mammalian species cross-react [16]. The greatest homology is found between a cow's,
350 buffalo's, sheep's and goat's milks protein. Proteins in their milks have less structural similarity with
351 pig, horse, donkey, camel and dromedary. Goats, buffalo and ewe's milk are particularly not
352 recommended by the World Allergy Organization due to cross-reactivity with cow's milk [16]. The
353 tolerance of other mammalian milks needs to be further investigated in clinical trials and there are
354 some concerns about their chemical composition and sanitation. In conclusion, either amino acid-
355 based formulas or extensively HF represent the most available solutions for allergic infants no longer
356 breast-fed. The therapeutic interventions should be therefore indicated on an individual basis.

357 5. CONCLUSION

358 Food allergy represents a significant health burden on an individual and population level
359 worldwide. Recent guidelines for the prevention of food allergies advocate that there is no need to
360 delay the introduction of allergenic foods once weaning has commenced. In terms of food allergy
361 management (end even prevention), individualized strategies should be implemented. These
362 strategies will include development readiness to be weaned, prevalence of particular food allergies in
363 certain countries, family eating patterns and availability of physician and dietetic care.

364 Care should be taken to ensure adequate intake of nutrients, particularly in relation to cow's milk
365 allergy, when selecting a suitable hypoallergenic formula. There is emerging evidence regarding the
366 role of fats (particularly AGEs), pre/probiotics, commercial foods, healthy eating and micronutrients
367 on food allergy. A better understanding of how nutrients and other aspects of food, food patterns and
368 food preparation may affect the immune system and allergy outcomes is required to best advise those
369 at risk of developing food allergies and those with current food allergies.

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371 **Conflicts of Interest:** The authors declare no conflict of interest.

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