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(19) **HU****MAGYARORSZÁG**
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N.V. Nutricia, 2712 HM Zoetermeer (NL) |
| (72) Feltaláló(k):
VAN GOUDOEVER, Johannes Bernard, NL-2275 CJ Voorburg (NL)
VAN DER BEEK, Eline Marleen, NL-6705 DB Wageningen (NL)
ABRAHAMSE-BERKEVELD, Marieke, NL-6666 HS Heteren (NL)
BOEHM, Günther, DE-04107 Leipzig (DE) | (74) Képviselő:
dr. Fehérvári Flóra, DANUBIA Szabadalmi és Jogi Iroda Kft., Budapest |

- (54) **Alacsony fehérjetartalmú csecsemőgyógyászati készítmény megemelt esszenciális aminosavtartalommal**

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(54) **LOW PROTEIN INFANT FORMULA WITH INCREASED ESSENTIAL AMINO ACIDS**

PROTEIN-ARMES UND REICH AN ESSENTIELLEN AMINOSÄUREN NAHRUNGSMITTEL FÜR
SÄUGLINGE

FORMULE INFANTILE A TENEUR REDUITE EN PROTEINES ET AUGMENTEE EN ACIDES AMINES
ESSENTIELS

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- **VAN DER BEEK, Eline Marleen**
NL-6705 DB Wageningen (NL)
- **ABRAHAMSE-BERKEVELD, Marieke**
NL-6666 HS Heteren (NL)
- **BOEHM, Günther**
DE-04107 Leipzig (DE)

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(74) Representative: **Nederlandsch Octrooibureau**
P.O. Box 29720
2502 LS The Hague (NL)

(73) Proprietor: **N.V. Nutricia**
2712 HM Zoetermeer (NL)

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(72) Inventors:
• **VAN GOUDOEVER, Johannes Bernard**
NL-2275 CJ Voorburg (NL)

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Description**Field of the invention**

5 [0001] The present invention is in the field of infant nutrition and the content of essential branched chain amino acids thereof.

Background of the invention

10 [0002] Knowledge of essential amino acid requirement in (preterm) infants is important since excessive or deficient intake might lead to long term morbidity such as obesity (Singhal et al. (2002) Am J Clin Nutr 75:993-9) or suboptimal growth and impaired neurodevelopment outcome (Stephens et al (2009) Pediatrics 2009;123:1337-43).

[0003] WO2008054200A2 relates to a method for preventing and/or treating visceral adiposity and/or to alter body fat distribution by administering a certain nutritional composition to a human subject with the age between 0 and 48 months, and preventing the occurrence of diseases later in life. The role of BCAA is not discussed in this document.

15 [0004] In EP0418593A2 protein, peptide and amino acid mixtures of optimized amino acid composition are provided which can be used as the protein base for the preparation of additives for women's milk and as a protein base for producing milk foods for prematurely born babies and other infants. The mixtures according to the invention contain 18 essential amino acids in a defined ratio, this ratio being calculated and selected on the basis of the nutritive protein content of women's milk.

20 [0005] The branched-chain amino acids (BCAA) leucine (Leu), isoleucine (Ile) and valine (Val) account for 35-40 % of the dietary indispensable amino acids in body protein and 14% of the total amino acids in skeletal muscle. Infant nutrition has been formulated using mother's milk, breast milk, as the most ideal composition. However, due to the different amino acid composition of protein sources used for production of infant nutrition, protein levels are most often higher compared to those in breast milk to ensure sufficient intake of BCAA.

25 [0006] It now becomes increasingly clear that early nutrition that is consumed by the infant has metabolic effects on the infant, also later in life, i.e. metabolic imprinting. For example WO 2008/054200 discloses that metabolic imprinting is an important mechanism in programming future health and food is specifically designed to prevent obesity later in life by giving low protein diets. Hence, diets too high or too low in protein may therefore be harmful to the infants. However, the exact requirement of essential amino acids for term and preterm infants is not known.

Summary of the invention

35 [0007] Historically, descriptive or gross measures like growth and nitrogen balance have been used for studying the amino acid requirements. No studies have been performed using stable isotope techniques to measure essential amino acid requirements for enteral fed infants properly. The inventors therefore investigated the essential amino acid requirement by using this preferred technique in enteral fed infants in order to determine the optimal ratio of the amino acids in a nutritional composition and to determine the absolute requirements of the three essential BCAA valine, isoleucine and leucine. The results of these studies now radically change the known concepts of the amino acid requirement. This is particularly relevant for formulas with a low concentration of protein that aim to prevent adverse metabolic imprinting effects in the infant.

40 [0008] The present inventors surprisingly found that the valine requirement, that is currently set on 87 mg/kg/day, should be between 100 and 175 mg/kg/day, with a preferred range between 110 and 160 mg/kg/day. Also for isoleucine and leucine increased requirements have been established by the inventors. For isoleucine it has been found to be between 100 and 160 mg/kg/day, preferably between 105 and 150 mg/kg/day; the current recommendation is 88 mg/kg/day. As the leucine requirement appeared to be comparable to the currently recommended values, the most prominent consequence of the present finding is that the ratios of these three essential branched chain amino acids has to be adapted, to a weight ratio Leu:Ile:Val between (1.1-1.5):(0.9-1.1):1.0. Instead of the currently recommended weight ratio Leu:Ile:Val of 1.9:1.0:1.0. Thus in one embodiment, the invention concerns an infant formula comprising protein, digestible carbohydrates and fat, wherein the protein comprises the amino acids leucine, isoleucine and valine in a weight ratio leucine:isoleucine:valine between (1.1-1.5):(0.9-1.1):1.0, and wherein the total protein content is between 1.3 and 1.9 g protein/100 kcal. Preferably, the weight ratio Leu:Ile:Val is between (1.3-1.5):(0.9-1.1):1.0, preferably between (1.3-1.5):(0.9-1.0):1.0, preferably between 1.3:1.0:1.0 and 1.5:0.9:1.0.

50 [0009] A preferred composition according to the invention therefore comprises per 100 ml between 40 and 120 mg valine, preferably between 55 and 120 mg valine, preferably between 70 and 110 mg valine and between 40 and 120 mg isoleucine, preferably between 55 and 110 mg isoleucine, preferably between 70 and 100 mg isoleucine and between 70 and 180 mg leucine, preferably between 90 and 170 mg leucine. The composition according to the invention is intended for an infant between 0-36 months. With a preferred energy content of between 60 and 70 kcal per 100 ml, a

preferred composition according to the invention comprises between 90 and 180 mg valine per 100 kcal in the total composition, preferably between 100 and 150 mg valine per 100 kcal, preferably between 105 and 121 mg valine per 100 kcal, and between 90 and 180 mg isoleucine per 100 kcal in the total composition, preferably between 95 and 170 mg, preferably between 100 and 150 mg isoleucine per 100 kcal and between 120 and 260 mg leucine per 100 kcal in the total composition, preferably between 120 and 180 mg leucine per 100 kcal, preferably between 130 and 160 mg leucine per 100 kcal. Preferably the ratio leucine:isoleucine:valine is between (1.3-1.5):(0.9-1.1):1.0

[0010] Infant formulas comprising the above mentioned ranges of amino acids are preferably used for enteral or parental feeding of an infant. In particular the infant formulas are used for preventing obesity during infancy or later in their life, while maintaining optimal growth in the period the infant is depending mainly on the composition for the protein intake.

Detailed description of the invention

[0011] *Protein content:* The term 'protein content' as used in the present document can be calculated from the nitrogen content using the formula: Nitrogen content x 6.25. The nitrogen content can be measured according to standard procedures known to the man skilled in the art.

[0012] *Infant:* The term 'infant' according to the present invention means a human with an age between 0 and 36 months, preferably between 0 and 18 months and even more preferably between 0 and 6 months. The younger the infant the more dependent the infant is on the infant formula for the protein intake. Therefore, the formula is preferred for the age group of less than 12 months or infants that due to digestive problems or allergies depend on the infant formula for their intake of protein.

[0013] *Premature and/or small for gestational age infants:* A premature infant relates to an infant born before the standard period of pregnancy is completed before or on 37 weeks pregnancy of the mother, i.e. before or on 37 weeks from the beginning of the last menstrual period of the mother. SGA infants are those whose birth weight lies below the 10th percentile for that gestational age. They have usually been the subject of intrauterine growth restriction (IUGR). Premature and/or SGA infants include low birth weight infants (LBW infants), very low birth weight infants (VLBW infants), and extremely low birth weight infants (ELBW infants). LBW infants are defined as infants with a weight less than 2500 g. VLBW infants as infants with a weight which is less than 1500 g, and ELBW infants as infants with a weight less than 1000 g.

[0014] *Infant formula* is a nutritionally complete formula comprising protein, fat, carbohydrates, and micronutrients such as vitamins and minerals. Preferably the infant formula comprises dietary fiber, nucleotides and the fatty acids arachidonic acid (AA) and docosahexaenoic acid (DHA). The term infant formula excludes human breast milk.

[0015] Good nutrition is essential for optimal growth and development in the preterm born and term born infant. Protein is an important component of adequate nutrition as it provides essential amino acids required for critical protein synthesis and growth. Nutrition is especially important during the early phase of life since protein intake in the first 4 weeks of life can have a major influence on later cognitive function and blood pressure.

[0016] The alarming increase in both prevalence and severity of obesity in children has renewed the interest of feeding pattern in infancy. High early weight gain in the first 1-2 year of life is associated with later adverse health outcomes, such as increased blood pressure, increased overweight and body fat deposition, and increased risk of diabetes. The higher protein intake in infants fed infant formula compared to breastfed children, may play a role since formula-fed children reach a higher body weight and weight for length at one year of age. Lowering the protein content of infant food might be one strategy that could contribute to decreasing these adverse effects. A good protein quality, i.e. the right amounts of nutritionally available amino acids becomes than more critical.

[0017] Classically nine amino acids are regarded as dietary essential; if these amino acids are not administered in the right proportions, protein synthesis will be reduced. The requirements of the indispensable amino acids have been determined by a number of different methods. Historically, descriptive or gross measures like growth and nitrogen balance have been used. No studies have been performed using stable isotope techniques to measure essential amino acid requirements for enteral fed infants properly.

[0018] For the vulnerable populations such as neonates, it is unacceptable to maintain a deficient diet for a long period.

[0019] Brunton et al. (1998) Curr Opin Clin Nutr Metab Care, 1(5): p. 449-53 validated a minimal invasive protocol to use the indicator amino acid oxidation (IAAO) technique in infants and children. This protocol has recently been used to determine total branched chain amino acid requirements in healthy school aged children. Requirement estimates in children were similar to the estimates in adult humans, which suggest that the experimentally derived values predominantly reflect maintenance requirements and do not take in account all the growth needs.

[0020] Current recommendations of infants 0-6 months are based the amino acid content of human milk which might be inadequate since breastfed infants have quite variable milk intakes and the breastfed infant themselves largely regulate the intake they require. The European Food and Safety Authority (EFSA) directives require infant formula to contain 88 mg valine, 92 mg isoleucine and 167 mg leucine per 100 kcal with a minimum of 60 and a maximum of 70

kcal/100ml formula, i.e. between 52.5 and 62.0 mg valine per 100ml, 55 and 64.9 mg isoleucine and between 100 and 118 mg leucine per 100 ml formula. According to the present invention these ranges are not adequate.

[0021] According to the present invention the total protein content in infant formula is between 1.3 and 1.9 g protein/100 kcal, even more preferably between 1.3 and 1.8 g protein/100 kcal, resulting in an infant formula comprising between 5.2 and 7.6 % protein based on the total calories of the composition. A preferred composition comprises between 90 and 180 mg valine and between 90 and 180 mg isoleucine per 100 kcal and between 130 and 260 mg leucine per 100 kcal. For an amino acid based or protein hydrolysates based composition the preferred ranges according to the present inventions are between 105 and 120 mg valine and between 100 and 120 mg isoleucine per 100 kcal and between 130 and 160 mg isoleucine per 100 kcal since these are the ranges that come closest to the experimentally determined average amino acid requirement (see examples 1-2). For non-hydrolysed (intact) protein based compositions, these latter narrow ranges are preferably about 10% lower, preferably between 10 and 20% lower. Thus in one embodiment, for non-hydrolysed (intact) protein based compositions the preferred ranges according to the present invention are between 105 and 120 mg valine and between 100 and 120 mg isoleucine per 100 kcal and between 130 and 160 mg isoleucine per 100 kcal.

[0022] In another preferred embodiment of the present invention an infant formula comprises a lipid, protein and digestible carbohydrate component wherein the protein component provides between 5.0 and 7.6% of the total calories, the lipid component provides 35 to 55% of the total calories and the digestible carbohydrate component provides 30 to 60% of the total calories for the manufacture of a nutritional composition, and wherein the composition comprises per 100 ml between 40 and 110 mg valine, preferably between 40 and 70 mg valine, and between 40 and 110 mg isoleucine. Preferably the weight ratio leucine:valine is in the range from 1.1:1.0 to 1.5:1.0, more preferably in the range from 1.3:1.0 to 1.5:1.0 and the weight ratio isoleucine:valine is in the range from 0.9:1.0 to 1.1:1.0, preferably is about 1.0:1.0 which means that the amounts of Ileu and Val do not necessarily have to be exactly the same, but lie within the rounded off margin of the second decimal. Thus the weight ratio leucine:isoleucine:valine is in the range from (1.1-1.5):(0.9-1.1):1.0, preferably in the range from (1.1-1.5):1.0:1.0.

[0023] Preferably the composition according to the invention is used for feeding infants that largely depend on their protein intake on the nutritional composition according to the present invention. These infants preferably have with an age range between 0 and 36 months preferably between 0 and 18 months and even more preferably between 0 and 6 months. The composition can also be used for infants suffering from (multiple) food allergies that would normally restrict their intact protein intake by using hydrolyzed protein formulas or amino acid based formulas. In addition the composition can be used for the dietary management of diseases like Phenylketonuria (PKU), maple syrup urine disease (MSUD) or Tyrosinaemia.

[0024] Normally infants are fed 150 ml of an infant milk formula per kg body weight per day. Preferably with this feeding regimen an infant ingests the amounts of leucine, isoleucine and valine now established. Thus in one embodiment, the invention concerns a non medical use of an infant formula for feeding an infant, or a non-medical method for feeding an infant comprising administering an infant formula, wherein the infant formula comprises protein that provides leucine, isoleucine and valine in a ratio of (1.1-1.5):(0.9-1.1):1.0, and the infant formula provides between 100 and 175 mg valine per kg body weight per day, and between 100 and 160 mg isoleucine per kg body weight per day when the infant is fed 150 ml of the nutritional composition per kg body weight per day.

Protein

[0025] Protein is preferably present in the composition below 8 % based on total calories of the composition. Preferably the nutritional composition comprises between 5.0 and 8.0 % protein based on total calories, more preferably between 5.5 and 8.0 %, and even more preferably between 5.7 and 7.6 % protein based on total calories. As total calories of the composition the sum of calories delivered by the fats, proteins and digestible carbohydrates of the composition is taken. A low protein concentration ensures a lower insulin response, thereby preventing proliferation of adipocytes, especially visceral adipocytes in infants. The protein concentration in a nutritional composition is determined by the sum of protein, peptides and free amino acids. The protein concentration is determined by determining the amount of nitrogen, multiplying this with a factor 6.25. One gram of protein equals 4 kcal. Based on dry weight the composition preferably comprises less than 12 wt.% protein, more preferably between 6 to 11 wt.%, even more preferably 7 to 10 wt.%. Based on a ready-to-drink or reconstituted powder liquid product the composition preferably comprises less than 1.5 g protein per 100 ml, more preferably between 0.8 and 1.35 g per 100 ml.

[0026] The source of the protein is preferably selected in such a way that the minimum requirements for essential amino acid content are met and satisfactory growth is ensured. Hence protein sources based on cows' milk proteins such as whey, casein and mixtures thereof and proteins based on soy are preferred. In case whey proteins are used, the protein source is preferably based on acid whey or sweet whey, whey protein isolate or mixtures thereof and may include α -lactalbumin and β -lactoglobulin. More preferably, the protein source is based on acid whey or sweet whey from which the caseino-glyco-macropptide (CGMP) has been removed. Removal of CGMP from sweet whey protein or the

use of acid whey advantageously reduces the threonine content. Preferably •-lactalbumin enriched whey protein is used in order to optimize the amino acid profile. Using protein sources with an optimized amino acid profile closer to that of human breast milk enables all essential amino acids to be provided at reduced protein concentration, below 8 % based on based on total energy content, preferably between 5.5 and 8.0% based on total energy content provided by the protein, fat and digestible carbohydrate and still ensure a satisfactory growth.

[0027] To ensure that low levels of total protein can be reached the nutritional compositions according to the present invention preferably comprise a protein source wherein the sum of leucine, isoleucine and valine provides at least 20 wt% of the total amino acid content.

[0028] If modified sweet whey is used as the protein source, it is preferably supplemented by free arginine in an amount from 0.1 to 3 wt.% and/or free histidine in an amount from 0.1 to 1.5 wt.% based on total protein.

[0029] The proteins may be intact or hydrolysed or a mixture of intact and hydrolysed proteins although intact proteins are generally preferred. Preferably the composition comprises hydrolyzed casein and/or hydrolyzed whey protein. It was found that administration of a composition wherein the protein comprises hydrolyzed casein and hydrolyzed whey results in reduced postprandial levels of both insulin and glucose compared to the administration of a composition comprising intact casein and intact whey protein. Increased levels of both insulin and glucose indicate a form of insulin insensitivity and/or resistance in formula fed infants. The present composition preferably comprises at least 25 wt.% peptides with a chain length of 2 to 30 amino acids based on dry weight of protein. The amount of peptides with a chain length between 2 and 30 amino acids can for example be determined as described by de Freitas et al, (1993), J. Agric. Food Chem. 41:1432-1438. The present composition may include casein hydrolysate or the present composition may include whey protein hydrolysate or both. The present composition preferably includes both casein hydrolysate and whey protein hydrolysate because the amino acid composition of bovine casein is more similar to the amino acid composition found in human milk protein and whey protein is easier to digest and found in greater ratios in human milk. The composition preferably comprises at least 50 wt.%, preferably at least 80 wt.%, most preferably about 100 wt.% of a protein hydrolysate, based on total weight of the protein. The present composition preferably comprises a protein with a degree of hydrolysis of the protein between 5 and 25%, more preferably between 7.5 and 21%, most preferably between 10 and 20%. The degree of hydrolysis is defined as the percentage of peptide bonds which have been broken down by enzymatic hydrolysis, with 100% being the total potential peptide bonds present. A suitable way to prepare a hydrolysate is described in WO 01/41581.

[0030] When using amino acid based protein source it will be better predictable how much of the amino acids will actually become available to the infant. Therefore, a preferred composition according to the invention comprises amino acids as protein source. In one embodiment of the composition according to the present invention the protein consist essentially of free amino acids. A preferred embodiment is shown in example 3.

Dietary fibers

[0031] Early differences in fecal nucleobacteria composition in children may predict overweight (Kalliomaki et al. (2008) Am J Clin Nutr 87(3):534-538). They showed that *Bifidobacterium* spp affecting both the quantity and quality of the microbiota during the first year of life was higher in number in children exhibiting a normal weight at 7 years than in children developing overweight. Without being bound by theory, the inventors believe that in addition to a low protein intake, the use of dietary fiber capable of stimulating the bifido flora will have an additional effect on preventing or treatment of obesity in infants. The dietary fibers are selected from the group consisting of fibers that can stimulate the growth of *Bifidobacterium* spp.

[0032] Preferably the non-digestible oligosaccharides have a DP between 2 and 60. The non-digestible oligosaccharide is preferably selected from the group consisting of fructo-oligosaccharides (including inulin), galacto-oligosaccharides (including transgalacto-oligosaccharides), gluco-oligosaccharides (including gentio-, nigero- and cyclodextrin-oligosaccharides), arabino-oligosaccharides, mannan-oligosaccharides, xylo-oligosaccharides, fuco-oligosaccharides, arabinogalacto-oligosaccharides, glucomanno-oligosaccharides, galactomanno-oligosaccharides, sialic acid comprising oligosaccharides and uronic acid oligosaccharides. Preferably the present composition comprises fructo-oligosaccharides, galacto-oligosaccharides and/or galacturonic acid oligosaccharides, more preferably galacto-oligosaccharides, most preferably beta-linked galacto-oligosaccharides. In a preferred embodiment the composition comprises a mixture of β-linked galacto-oligosaccharides and fructo-oligosaccharides, more preferably in a weight ratio of 20-2 : 1 more preferably 12-7 : 1. Preferably the present composition comprises galacto-oligosaccharides with a DP of 2-10 and/or fructooligosaccharides with a DP of 2-60. The galacto-oligosaccharide is preferably selected from the group consisting of beta-linked galacto-oligosaccharides, transgalacto-oligosaccharides, galacto-oligosaccharides, lacto-N-tetraose (LNT), lacto-N-neotetraose (neo-LNT), fucosyl-lactose, fucosylated LNT and fucosylated neo-LNT. β-linked galacto-oligosaccharides are for example sold under the trademark Vivinal™ (Borculo Domo Ingredients, Netherlands). Preferably the saccharides of the galacto-oligosaccharides are •-linked, since this is also the case in human milk galacto-oligosaccharides. Fructo-oligosaccharide is a NDO comprising a chain of beta-linked fructose units with a DP or average DP of 2 to 250, more

preferably 10 to 100. Fructo-oligosaccharide includes inulin, levan and/or a mixed type of polyfructan. An especially preferred fructo-oligosaccharide is inulin. Fructo-oligosaccharide suitable for use in the compositions is also already commercially available, e.g. Raftiline®HP (Orafti). Uronic acid oligosaccharides are preferably obtained from pectin degradation, more preferably apple pectin, beet pectin and/or citrus pectin. Preferably the composition comprises β -linked galacto-oligosaccharide: fructo-oligosaccharide: uronic acid oligosaccharide in a weight ratio of 20-2 : 1 : 1-3 more preferably 12-7 : 1 : 1-2.

[0033] Preferably, the composition comprises of 80 mg to 2 g non-digestible oligosaccharides per 100 ml, more preferably 150 mg to 1.50 g, even more preferably 300 mg to 1 g. Based on dry weight, the composition preferably comprises 0.25 wt.% to 5.5 wt.%, more preferably 0.5 wt.% to 4 wt.%, even more preferably 1.5 wt.% to 3 wt.%. A lower amount of non-digestible oligosaccharides will be less effective in stimulating the beneficial bacteria in the microbiota, whereas a too high amount will result in side-effects of bloating and abdominal discomfort.

[0034] A study has been done determining the stimulation of the bifidoflora in formula fed infants showing an improved bifidogenic effect of galacto-oligosaccharides and long chain inulin when they are given in combination, compared to a formula with only one dietary fiber.

[0035] Infants of 9-10 months of age where given 500 ml per day of a milk formula comprising 1.2 g beta-galacto-oligosaccharides plus long chain inulin for 1 month (group A). A control group received milk formula without beta-galacto-oligosaccharides and long chain inulin (group B). In total 138 children entered the study. The faecal flora was examined before and after this period by FISH analysis.

[0036] After the intervention time, the ratio of Bifidobacteria/total bacteria was significantly higher in group A than in group B. A preferred composition according to the invention therefore comprises a combination of beta-galacto-oligosaccharides and long chain inulin.

Fat

[0037] As explained above, the protein content of infant formula has effects on the lipid metabolism and fat deposition in infants. Also, the lipid composition of the fat in infant formula has been shown to have an important impact on the prevention of obesity, and in particular on central obesity or visceral adiposity, see for example WO2008/054208. The term 'visceral adiposity' refers to a condition with increased visceral fat mass. The term visceral adiposity is also referred to as central obesity. Visceral adiposity is typically caused by (accumulation of) excessive visceral fat mass. Visceral fat, also known as organ fat, intra-abdominal fat, peritoneal fat or central fat is normally located inside the peritoneal cavity as opposed to subcutaneous fat which is found underneath the skin and intramuscular fat which is found interspersed in skeletal muscles. Visceral fat includes mesenteric fat, perirenal fat and retroperitoneal fat.

[0038] Previous research, disclosed in WO2008/054208, showed that Medium chain fatty acids (MCFA) contribute to a reduced fat mass later in life. Therefore, in addition to the specific protein and amino acid content according to the present claims, the composition advantageously comprises MCFA.

[0039] The effect on visceral fat deposition were specifically shown when using a nutritional composition comprising (i) a LA/ ALA ratio between 2 and 6 and (ii) a low LA content (<14.5 wt.% based on total fatty acids) and optionally LC-PUFA (particularly DHA). Such composition resulted in a decrease in visceral adiposity later in life.

[0040] A preferred nutritional composition therefore comprises protein, digestible carbohydrates and fat, wherein the protein comprises the amino acids leucine, isoleucine and valine in a ratio leucine:isoleucine:valine in the range from (1.1-1.5):(0.9-1.1):1.0, and the fat comprises linoleic acid and alpha linoleic acid in a ratio between 2 and 6 and the linoleic acid content is less than 14.5 wt.% based on total fatty acids. This composition could preferably be used for enteral feeding of an infant for prevention of obesity later in life.

Applications

[0041] The present infant formula is advantageously to be ingested by or administered to premature infants or small for gestational age infants and hence is intended for the enteral or parenteral treatment of premature infants or small for gestational age infants. Also the present infant formula, and preferably the form wherein the protein consists essentially of free amino acids, is advantageously used for the treatment of infants with a metabolic disease selected from the group PKU, MSUD and tyrosinaemia, or for the treatment of infants with a food allergy. Preferably the present infant formula is advantageously to be ingested by or administered to an infant with an age between 0 and 36 months, preferably between 0 and 24 months.

[0042] The invention also concerns the use of the infant formula according to the present invention for the manufacture of a medicament for the treatment of a) infants with an age between 0 and 24 months and/or b) premature infants or c) small for gestational age infants or d) infants with a metabolic disease selected from the group consisting of PKU, MSUD and tyrosinaemia or e) infants with a food allergy.

[0043] In a preferred embodiment, the present invention concerns the use of the infant formula according to the present

invention for the manufacture of a medicament for the prevention of obesity later in life.

[0044] Also the present invention concerns the non medical use of the infant formula according to the present invention for the manufacture of a composition for feeding of an infant with an age in the range between 0 and 36 months.

[0045] Also the present invention concerns the non medical use of an infant formula for feeding an infant, wherein the infant formula comprises protein that provides leucine, isoleucine and valine in a ratio of (1.1-1.5):(0.9-1.1):1.0, and the infant formula provides between 100 and 175 mg valine per kg body weight per day, and between 100 and 160 mg isoleucine per kg body weight per day when the infant is fed 150 ml of the nutritional composition per kg body weight per day.

Examples

Example 1. Valine and isoleucine requirements in infant

Experimental design

[0046] Term male infants (n=28) were enrolled in this study. They had a gestational age of 37-43 weeks, a birth weight of more than 2500 gram and their postnatal age was $\geq$ 28 days.

[0047] The IAAO technique (Zello et al. (1993) Am J Physiol 264:E677-85) was used to determine the requirement of valine and in a separate experiment to determine the requirement of isoleucine. This method uses an indicator that is oxidized when one essential amino acid is limiting, since there is no storage of amino acids and amino acids must be partitioned between incorporation into protein or oxidation. If the tested amino acid is deficient in the diet, this will limit protein synthesis and the indicator amino acid will be oxidized. If the dietary intake of the test amino acid increases, the oxidation of the indicator will decrease until requirement of the test amino acid is met. When intake meets the requirement then protein synthesis occurs at optimum capacity and the oxidative degradation of all other essential amino acids plateau. The requirement of the test amino acid is identified by this breakpoint.

[0048] During the study period, subjects were randomly assigned to receive graded intakes of valine ranging from 5 to 236 mg/kg day or graded intakes of isoleucine ranging from 5 to 216 mg/kg day. After adaptation to the study diet for 24 hours, baseline breath samples were obtained and a tracer protocol was started. Subjects were weighted daily, before and at the end of the tracer protocol and a head circumference was measured at the study day.

Study Formula

[0049] We used a study formula identical to regular Neocate, an amino acid based formula designed to fulfil the amino acid requirements of infants (SHS, Liverpool, UK) but without the test amino acid and with a decreased amount of phenylalanine. The amount of valine was adjusted separately as L-valine. The amount of isoleucine was adjusted separately as L-isoleucine. L-phenylalanine was supplied during the adaptation time and during the infusion of [^{13}C] bicarbonate to obtain a stable total intake of 166 mg/kg/d during the entire study. To make the formula isonitrogenous we added L-alanine separately. Since phenylalanine was used as an indicator and phenylalanine is hydroxylated to tyrosine before oxidation can occur, we made sure that tyrosine intake was high above present requirements. A too limited tyrosine intake might reduce recovery of ^{13}C label in expiratory air. To minimize the effect of feeding on the [^{13}C] bicarbonate plateau continuous dripfeeding was given during the [^{13}C]bicarbonate infusion. To minimize the discomfort for the subjects they could drink a bottle every hour during the [^{1-13}C]phenylalanine infusion..

Tracer protocol

[0050] On the study day subjects received a primed (15 $\mu\text{mol}/(\text{kg})$) continuous (10 $\mu\text{mol}/(\text{kg}\cdot\text{h})$) enteral infusion of [^{13}C] bicarbonate (sterile pyrogen free, 99% ^{13}C Atom Percent Excess (APE); Cambridge Isotopes, Woburn, MA) for 3 h to quantify individual CO_2 production. The labeled sodium bicarbonate infusion was directly followed by a primed (30 $\mu\text{mol}/(\text{kg})$, continuous (30 $\mu\text{mol}/(\text{kg}\cdot\text{h})$) enteral infusion of [^{1-13}C]phenylalanine (99% ^{13}C APE; Cambridge Isotopes, Woburn, MA) for five hours by an infusion pump via the nasogastric tube. The syringes with tracers were weighted before and after infusion to determine the exact amount of tracer given during the study.

Sample collection

[0051] Breath samples were collected at the adaptation day in the first 8 patients to determine the time needed obtain a stable background enrichment, using the direct sampling method described by Van der Schoor et al. (2004) Pediatr Res 55:50-4. At the study day baseline samples were obtained 15 and 5 minutes before starting tracer infusion. During the experiment duplicate ^{13}C -enriched breath samples were collected every 10 minutes during the isotopic steady state

of the [^{13}C]bicarbonate infusion starting after 1.75 hours, and every 15 minutes during the isotopic steady state of the [$1\text{-}^{13}\text{C}$]phenylalanine infusion starting after 3 hours.

Analysis and Calculations

[0052] $^{13}\text{CO}_2$ isotopic enrichment in expired air was measured by isotope ratio mass spectrometry (ABCA; Europe Scientific, Van Loenen Instruments, Leiden, the Netherlands) and expressed as APE above baseline. Steady state was defined as three or more consecutive points with a slope not different from zero ($p < 0.05$). Estimated body CO_2 production ($\text{mmol}/(\text{kg}\cdot\text{h})$) was calculated for each infant as described previously (Riedijk et al. (2005) *Pediatr Res* 58:861-4). The rate of fractional [$1\text{-}^{13}\text{C}$]phenylalanine oxidation was calculated using the following equation:

$$\text{fractional phenylalanine oxidation (\%)} = [\text{IE}_{\text{PHE}} \times i_{\text{B}}] / [i_{\text{PHE}} \times \text{IE}_{\text{B}}] \times 100\%$$

where IE_{PHE} is the ^{13}C isotopic enrichment in expired air during [$1\text{-}^{13}\text{C}$]phenylalanine infusion (APE), i_{B} is the infusion rate of [^{13}C]bicarbonate ($\mu\text{mol}/(\text{kg}\cdot\text{h})$), i_{PHE} is the infusion rate of [$1\text{-}^{13}\text{C}$]phenylalanine ($\mu\text{mol}/(\text{kg}\cdot\text{h})$) and IE_{B} is the ^{13}C isotopic enrichment in expired air during [^{13}C]bicarbonate infusion (van der Schoor et al. (2004) *Gut* 53: 38-43).

Statistical analysis

[0053] Descriptive data were expressed as mean $\pm$ SD. Steady state of $^{13}\text{CO}_2$ in expired breath during the [$1\text{-}^{13}\text{C}$]phenylalanine was achieved when the linear factor of the slope was found to be not significantly different from zero ($p < 0.05$). The valine requirement was determined by applying a two-phase linear regression crossover model (Ball and Bayley (1984) *J Nutr* 114: 1741-6; Seber GAF. *Linear Regression Analysis*. New York: John Wiley, 1977) on the fractional oxidation rates. The safe level of intake (upper 95% CI) was determined using the Fieller's theorem (Seber GAF. *Ibid.*). All statistical analyses were done using SPSS (SPSS version 15.0, Chicago, IL, USA).

Results

[0054] All subjects achieved isotopic steady state (plateau) at both [^{13}C]bicarbonate and [$1\text{-}^{13}\text{C}$]phenylalanine infusion defined by the absence of a significant slope between the data points at either plateau.

[0055] The Spearman's rank correlation coefficient between valine intake and the fractional oxidation was 0.63 ($p=0.000$). Using the two-phase regression analysis with the valine intake as the independent variable and the fractional oxidation of the [$1\text{-}^{13}\text{C}$]phenylalanine tracer as the dependant variables, the breakpoint was determined to be 110 mg/kg/d. The safe population intake determined by the upper 95% CI was 164.6 mg/kg/d.

[0056] The Spearman's rank correlation coefficient between isoleucine intake and the fractional oxidation was 0.74 ($p=0.000$). From the two-phase regression analysis with the isoleucine intake as the independent variable and the fractional oxidation of the [$1\text{-}^{13}\text{C}$]phenylalanine tracer as the dependant variable, the breakpoint was determined to be 105 mg/kg/d. The population-safe intake determined by the upper 95% CI was 152 mg/kg/d.

Example 2. Leucine requirement of infants

[0057] Following the same procedure as in example 1, the leucine requirement of infants was determined.

[0058] Term male infants ($n=33$) were enrolled in this study. They had a gestational age of 37-43 weeks, a birth weight of more than 2500 gram and their postnatal age was ≥ 28 days.

Results

[0059] All subjects achieved isotopic steady state (plateau) at both [^{13}C]bicarbonate and [$1\text{-}^{13}\text{C}$]phenylalanine infusion defined by the absence of a significant slope between the data points at either plateau.

[0060] Using the two-phase regression analysis with the leucine intake as the independent variable and the fractional oxidation of the [$1\text{-}^{13}\text{C}$]phenylalanine tracer as the dependant variables, the breakpoint was determined to be 140 mg/kg/d. The safe population intake determined by the upper 95% CI was 245 mg/kg/d.

Example 3. Amino acid based nutrition suitable for treating allergic infants

[0061]

EP 2 552 242 B1

	NUTRIENT NAME	PER 100 kcal	
	PROTEIN EQUIVALENT (N x 6.25)	1.36	g
5	NITROGEN	0.22	g
	CARBOHYDRATE	12.94	g
	FAT (TOTAL)	4.75	g
	(MCT)	0.19	g
10	(LCT)	4.28	g

MINERALS

	SODIUM	24.98	mg
15	POTASSIUM	138.45	mg
	CHLORIDE	106.96	mg
	CALCIUM	96.47	mg
	PHOSPHORUS	70.20	mg
20	MAGNESIUM	5.11	mg
	Ca: P RATIO	1.37	

TRACE ELEMENTS

25	IRON	1.48	mg
	ZINC	1.03	mg
	IODINE	14.22	µg
	MANGANESE	0.08	mg
30	COPPER	0.08	mg
	MOLYBDENUM	2.97	µg
	SELENIUM	2.28	µg
	CHROMIUM	2.06	µg

VITAMINS

	VITAMIN A	151.24	µg
	VITAMIN E	1.12	IU
40	L-ASCORBIC ACID	21.90	mg
	THIAMIN	0.11	mg
	RIBOFLAVIN	0.15	mg
	PYRIDOXINE	0.13	mg
45	NIACIN	1.16	mg
	PANTOTHENIC ACID	0.60	mg
	MYO-INOSITOL	25.05	mg
	CHOLINE	12.42	mg
	VITAMIN D3	1.94	µg
50	CYANOCOBALAMIN	0.31	µg
	FOLACIN	10.33	µg
	d-BIOTIN	6.82	µg
	VITAMIN K1	7.23	µg
55	NIACIN (equivalent)	1.64	mg

AMINO ACIDS

	L-ALANINE	55.4	mg
5	L-ARGININE	98.3	mg
	L-ASPARTIC ACID	91.7	mg
	L-CYSTINE	36.3	mg
	L-GLUTAMIC ACID	0	mg
	GLYCINE	86.5	mg
10	L-HISTIDINE	56.2	mg
	L-ISO LEUCINE	108.5	mg
	L-LEUCINE	148.4	mg
	L-LYSINE	100.9	mg
15	L-METHIONINE	23.7	mg
	L-PROLINE	105.2	mg
	L-PHENYLALANINE	66.1	mg
	L-SERINE	64.7	mg
	L-THREONINE	72.8	mg
20	L-TRYPTOPHAN	29.1	mg
	L-TYROSINE	66.1	mg
	L-VALINE	109.7	mg
	L-ASPARAGINE	0	mg
25	L-CITRULLINE	0	mg
	L-CARNITINE	1,2	mg

Example 4. Infant formula comprising intact milk proteins amino acids according to the invention.

30 **[0062]**

	COMPONENT	UNIT	Per 100 kcal
	1. Protein (equivalent)	g	1,4
	2. Carbohydrate	g	11,7
35	- Sugars	g	11,5
	3. Fat	g	5,12
	- Saturates	g	2,22
	4. Fibre, dietary	g	0,856
40	5. Sodium	g	25,8

D. NUTRITIONAL PROPERTIES

Energy percentages:

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[0063]

	1. Protein (equivalent)	En%	5,6
	2. Carbohydrate	En%	46,4 *
50	3. Fat	En%	46,3
	4. Fibre	En%	1,7
	Total	En%	100
55	* Including organic acids and polyols		

E. COMPOSITION**[0064]**

5	<u>1. Protein (equivalent), total</u>	g	1,4		
	- Nitrogen (Protein)	g	0,224		
	- Animal protein	g	1,428		
10	Whey protein	g	0,861		
	· Casein	g	0,574		
	· isoleucine and valine each	mg	25		
			g	11,7	
15	<u>2. Carbohydrate</u>				
	- Sugars	g	11,5		
	glucose	g	0,4		
20	· Fructose	g			
	· Galactose	g	0,026		
	· Lactose	g	11,1		
25	<u>3. Fat</u> g 5,12				
	- Vegetable	g	5,02		
	- Animal	g	0,11		
	· of which milk	g	0,06		
30	- Saturates	g	2,22		
	· of which MCT	g			
35	- Monounsaturates	g	2,06		
	- Polyunsaturates	g	0,84		
	<u>4. Fibre, dietary</u>	g	0,856		
40	- Soluble	g	0,86		
	- Insoluble	g			
	<u>5. Moisture/ water</u>	g	136		

F. AMINO ACID COMPOSITION**[0065]**

50	L-Alanine	mg	65,8	
	L-Arginine	mg	46,9	
	L-Aspartic acid / L-Asparagine	mg	116,2	
	L-Cyst(e)ine	mg	25,2	
	L-Glutamic acid / L-Glutamine	mg	263,2	
55	Glycine	mg	29,4	
	L-Histidine	mg	33,6	
	L-Isoleucine	mg	109,7	(84,7 from protein)
	L-Leucine	mg	158,2	

(continued)

L-Lysine	mg	136,5
L-Methionine	mg	37,1
L-Phenylalanine	mg	63,7
L-Proline	mg	110,6
L-Serine	mg	86,8
L-Threonine	mg	86,8
L-Tryptophan	mg	23,1
L-Tyrosine	mg	59,5
L-Valine	mg	114,6

 (89,6 from protein) ratio leucine:isoleucine:valine=1.4:1.0:1.0

Claims

1. An infant formula comprising protein, digestible carbohydrates and fat, wherein the protein comprises the amino acids leucine, isoleucine and valine in a weight ratio leucine:isoleucine:valine between (1.1-1.5):(0.9-1.1):1.0, and wherein the total protein content is between 1.3 and 1.9 g protein/100 kcal.
2. The infant formula according to claim 1 wherein the weight ratio leucine:isoleucine:valine is between (1.3-1.5):(0.9-1.1):1.0.
3. The infant formula according to claim 1 or 2 wherein the sum of leucine, isoleucine and valine provides at least 20 wt% of the total amino acid content
4. The infant formula according to any of the preceding claims, wherein the protein comprises between 120 and 180 mg leucine per 100 kcal in the total composition, between 90 and 180 mg isoleucine per 100 kcal in the total composition, and between 90 and 180 mg valine per 100 kcal in the total composition.
5. The infant formula according to claim 4, wherein the protein comprises between 130 and 160 mg leucine per 100 kcal in the total composition, between 100 and 120 mg isoleucine per 100 kcal in the total composition and between 105 and 121 mg valine per 100 kcal in the total composition.
6. The infant formula according to any of the preceding claims, wherein the protein is present between 5 and 8 % based on total calories of the composition.
7. The infant formula according to any of the preceding claims further comprising at least one bifidogenic dietary fiber selected from the group consisting of beta-linked galacto -oligosaccharides, transgalacto-oligosaccharides, galacto-oligosaccharides, lacto-N-tetraose (LNT), lacto-N-neotetraose (neo-LNT), fucosyl-lactose, fucosylated LNT, fucosylated neo-LNT, and xylooligosaccharides.
8. The infant formula according to to any of the preceding claims, further comprising at least one PUFA selected from the group consisting of ARA, DHA and EPA.
9. The infant formula according to any of the preceding claims wherein the protein consists essentially of free amino acids.
10. The infant formula according to any of the preceding claims wherein the total protein content is between 1.3 and 1.8 g protein/100 kcal.
11. The infant formula according to any of the preceding claims, for use in the enteral or parenteral treatment of premature infants or small for gestational age infants.
12. The infant formula according to claim 9 for use in the treatment of infants with a metabolic disease selected from the group PKU, MSUD and tyrosinaemia, or for the treatment of infants with a food allergy.

13. Infant formula according to any of claims 1-11 for use in the treatment of;

- a. premature infants; or
- b. small for gestational age infants; or
- c. infants with a metabolic disease selected from the group consisting of PKU, MSUD and tyrosinaemia; or
- d. infants with a food allergy.

14. Infant formula according to any of claims 1-11 for use in the prevention of obesity later in life.

15. Non-therapeutic use of the infant formula according to any of claims 1-10 for feeding of an infant with an age in the range between 0 and 36 months.

Patentansprüche

1. Nahrungsmittel für Säuglinge, das Protein, verdauliche Kohlenhydrate und Fett umfasst, wobei das Protein die Aminosäuren Leucin, Isoleucin und Valin in einem Gewichtsverhältnis Leucin : Isoleucin : Valin zwischen (1,1-1,5) : (0,9-1,1) : 1,0 umfasst und wobei der Gesamtproteingehalt zwischen 1,3 und 1,9 g Protein/100 kcal liegt.
2. Nahrungsmittel für Säuglinge nach Anspruch 1, wobei das Gewichtsverhältnis Leucin : Isoleucin : Valin zwischen (1,3-1,5) : (0,9-1,1) : 1,0 liegt.
3. Nahrungsmittel für Säuglinge nach Anspruch 1 oder 2, wobei die Summe von Leucin, Isoleucin und Valin wenigstens 20 Gew.-% des Gesamtaminosäuregehalts bereitstellt.
4. Nahrungsmittel für Säuglinge nach einem der vorangehenden Ansprüche, wobei das Protein zwischen 120 und 180 mg Leucin pro 100 kcal in der Gesamtzusammensetzung, zwischen 90 und 180 mg Isoleucin pro 100 kcal in der Gesamtzusammensetzung und zwischen 90 und 180 mg Valin pro 100 kcal in der Gesamtzusammensetzung umfasst.
5. Nahrungsmittel für Säuglinge nach Anspruch 4, wobei das Protein zwischen 130 und 160 mg Leucin pro 100 kcal in der Gesamtzusammensetzung, zwischen 100 und 120 mg Isoleucin pro 100 kcal in der Gesamtzusammensetzung und zwischen 105 und 121 mg Valin pro 100 kcal in der Gesamtzusammensetzung umfasst.
6. Nahrungsmittel für Säuglinge nach einem der vorangehenden Ansprüche, wobei das Protein zwischen 5 und 8 %, basierend auf den Gesamtkalorien der Zusammensetzung, vorhanden ist.
7. Nahrungsmittel für Säuglinge nach einem der vorangehenden Ansprüche, das weiterhin wenigstens eine bifidogene Diätfaser, ausgewählt aus der Gruppe bestehend aus beta-verknüpften Galaktooligosacchariden, Trans-Galaktooligosacchariden, Galaktooligosacchariden, Lacto-N-tetraose (LNT), Lacto-N-neotetraose (neo-LNT), Fucosyllactose, fucosylierter LNT, fucosylierter neo-LNT und Xylooligosacchariden, umfasst.
8. Nahrungsmittel für Säuglinge nach einem der vorangehenden Ansprüche, das weiterhin wenigstens eine PUFA, ausgewählt aus der Gruppe bestehend aus ARA, DHA und EPA, umfasst.
9. Nahrungsmittel für Säuglinge nach einem der vorangehenden Ansprüche, wobei das Protein im Wesentlichen aus freien Aminosäuren besteht.
10. Nahrungsmittel für Säuglinge nach einem der vorangehenden Ansprüche, wobei der Gesamtproteingehalt zwischen 1,3 und 1,8 g Protein/100 kcal liegt.
11. Nahrungsmittel für Säuglinge nach einem der vorangehenden Ansprüche zur Verwendung bei der enteralen oder parenteralen Behandlung von Frühgeborenen oder bezogen auf das Reifealter kleinen Säuglingen.
12. Nahrungsmittel für Säuglinge nach Anspruch 9 zur Verwendung bei der Behandlung von Säuglingen mit einer Stoffwechselerkrankung, ausgewählt aus der Gruppe PKU, MSUD und Tyrosinämie, oder für die Behandlung von Säuglingen mit einer Nahrungsmittelallergie.

13. Nahrungsmittel für Säuglinge nach einem der Ansprüche 1-11 zur Verwendung bei der Behandlung von:

- a. - Frühgeborenen; oder
- b. - bezogen auf das Reifealter kleinen Säuglingen; oder
- c. - Säuglingen mit einer Stoffwechselerkrankung, ausgewählt aus der Gruppe bestehend aus PKU, MSUD und Tyrosinämie; oder
- d. - Säuglingen mit einer Nahrungsmittelallergie.

14. Nahrungsmittel für Säuglinge nach einem der Ansprüche 1-11 zur Verwendung bei der Prävention von Adipositas im späteren Leben.

15. Nichttherapeutische Verwendung des Nahrungsmittels für Säuglinge nach einem der Ansprüche 1-10 zur Ernährung eines Säuglings mit einem Alter im Bereich zwischen 0 und 36 Monaten.

Revendications

1. Lait infantile comprenant des protéines, des glucides digestibles et des graisses, dans lequel les protéines comprennent les acides aminés leucine, isoleucine et valine dans un rapport pondéral leucine/isoleucine/valine situé entre (1,1 à 1,5)/(0,9 à 1,1)/1,0, et dans lequel la teneur totale en protéines se situe entre 1,3 et 1,9 g de protéines/100 kcal.
2. Lait infantile selon la revendication 1, dans lequel le rapport pondéral leucine/isoleucine/valine se situe entre (1,3 à 1,5)/(0,9 à 1,1)/1,0.
3. Lait infantile selon la revendication 1 ou 2, dans lequel la somme de la leucine, de l'isoleucine et de la valine fournit au moins 20 % en poids de la teneur totale en acides aminés.
4. Lait infantile selon l'une quelconque des revendications précédentes, dans lequel les protéines comprennent entre 120 et 180 mg de leucine pour 100 kcal dans la composition totale, entre 90 et 180 mg d'isoleucine pour 100 kcal dans la composition totale, et entre 90 et 180 mg de valine pour 100 kcal dans la composition totale.
5. Lait infantile selon la revendication 4, dans lequel les protéines comprennent entre 130 et 160 mg de leucine pour 100 kcal dans la composition totale, entre 100 et 120 mg d'isoleucine pour 100 kcal dans la composition totale, et entre 105 et 121 mg de valine pour 100 kcal dans la composition totale.
6. Lait infantile selon l'une quelconque des revendications précédentes, dans lequel les protéines sont présentes à entre 5 et 8 % en se basant sur les calories totales de la composition.
7. Lait infantile selon l'une quelconque des revendications précédentes, comprenant en outre au moins une fibre alimentaire bifidogène choisie dans le groupe constitué de galacto-oligosaccharides à liaisons bêta, transgalacto-oligosaccharides, galacto-oligosaccharides, lacto-N-tétraose (LNT), lacto-N-néotétraose (néo-LNT), fucosyl-lactose, LNT fucosylé, néo-LNT fucosylé, et xylo-oligosaccharides.
8. Lait infantile selon l'une quelconque des revendications précédentes, comprenant en outre au moins un AGPI choisi dans le groupe constitué d'ARA, DHA et EPA.
9. Lait infantile selon l'une quelconque des revendications précédentes, dans lequel les protéines sont constituées essentiellement d'acides aminés libres.
10. Lait infantile selon l'une quelconque des revendications précédentes, dans lequel la teneur totale en protéines se situe entre 1,3 et 1,8 g de protéines/100 kcal.
11. Lait infantile selon l'une quelconque des revendications précédentes, pour une utilisation dans le traitement entéral ou parentéral des nourrissons prématurés ou des nourrissons petits pour l'âge gestationnel.
12. Lait infantile selon la revendication 9, pour une utilisation dans le traitement des nourrissons souffrant d'une maladie métabolique choisie dans le groupe de PCU, MSUD et tyrosinémie, ou pour le traitement de nourrissons souffrant

d'une allergie alimentaire.

13. Lait infantile selon l'une quelconque des revendications 1 à 11 pour une utilisation dans le traitement de :

- 5 a. nourrissons prématurés ; ou
 b. nourrissons petits pour l'âge gestationnel ; ou
 c. nourrissons souffrant d'une maladie métabolique choisie dans le groupe constitué de PCU, MSUD et tyrosinémie ; ou
10 d. nourrissons souffrant d'une allergie alimentaire.

14. Lait infantile selon l'une quelconque des revendications 1 à 11 pour une utilisation dans la prévention de l'obésité ultérieurement dans la vie.

15. Utilisation non thérapeutique du lait infantile selon l'une quelconque des revendications 1 à 10 pour l'alimentation
15 des nourrissons d'un âge dans la gamme comprise entre 0 et 36 mois.

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REFERENCES CITED IN THE DESCRIPTION

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Szabadalmi igénypontok

1. Csecsemőgyógyászati készítmény, amely tartalmaz fehérjét, emészthető szénhidrátokat és zsírt, ahol a fehérje leucin, izoleucin és valin aminosavakat tartalmaz $(1,1-1,5):(0,9-1,1):1,0$ közötti leucin:izoleucin:valin tömegarányban és ahol a teljes fehérjetartalom $1,3-1,9$ g fehérje /100 kcal között van.

2. Az 1. igénypont szerinti csecsemőgyógyászati készítmény, ahol a leucin:izoleucin:valin tömegarány $(1,3-1,5):(0,9-1,1):1,0$.

3. Az 1. vagy 2. igénypont szerinti csecsemőgyógyászati készítmény, ahol a leucin, az izoleucin és a valin össz mennyisége a teljes aminosavtartalom legalább 20 tömeg%-át teszi ki.

4. Az előző igénypontok bármelyike szerinti csecsemőgyógyászati készítmény, ahol a fehérje $120-180$ mg leucin per 100 kcal között van a teljes készítményben, $90-180$ mg izoleucin per 100 kcal között van a teljes készítményben és $90-180$ mg valin per 100 kcal között van a teljes készítményben.

5. A 4. igénypont szerinti csecsemőgyógyászati készítmény, ahol a fehérje $130-160$ mg leucin per 100 kcal között van a teljes készítményben, $100-120$ mg izoleucin per 100 kcal között van a teljes készítményben és $105-121$ mg valin per 100 kcal között van a teljes készítményben.

6. Az előző igénypontok bármelyike szerinti csecsemőgyógyászati készítmény, ahol a fehérje a készítmény összes kalóriájára vonatkoztatva 5% és 8% között van jelen.

7. Az előző igénypontok bármelyike szerinti csecsemőgyógyászati készítmény, amely tartalmaz továbbá legalább egy hasított diétás rostot béta típusú kötésekkel tartalmazó galakto-oligoszacharidok, transzgalakto-oligoszacharidok, galakto-oligoszacharidok, lakto-N-tetralóz (LNT), lakto-N-neotetralóz (neo-LNT), fukozil-laktóz, fukozilezett LNT, fukozilezett neo-LNT és xilooligoszacharidok csoportjából kiválasztva.

8. Az előző igénypontok bármelyike szerinti csecsemőgyógyászati készítmény, amely tartalmaz továbbá legalább egy PUFA-t ARA, DHA és EPA csoportjából kiválasztva.

9. Az előző igénypontok bármelyike szerinti csecsemőgyógyászati készítmény, ahol a fehérje lényegében szabad aminosavakból áll.

10. Az előző igénypontok bármelyike szerinti csecsemőgyógyászati készítmény, ahol a teljes fehérjetartalom $1,3-1,8$ g fehérje/100 kcal között van.

11. Az előző igénypontok bármelyike szerinti csecsemőgyógyászati készítmény koraszülött csecsemők vagy a terhességi időhöz képes kis súllyal született csecsemők enterális vagy parenterális kezelésében történő alkalmazásra.

12. A 9. igénypont szerinti csecsemőgyógyászati készítmény PKU, MSUD és tirozinémia közül kiválasztott metabolikus betegségben szenvedő csecsemők kezelésében történő alkalmazásra vagy ételallergiás csecsemők kezelésére.

13. Az 1-11. igénypontok bármelyike szerinti csecsemőgyógyászati készítmény a következők kezelésében történő alkalmazásra:

- a) koraszülött csecsemők; vagy
- b) terhességi időhöz képest kis súllyal született csecsemők; vagy
- c) PKU, MSUD és tirozinémia csoportjába tartozó metabolikus betegségben szenvedő csecsemők; vagy
- d) ételallergiás csecsemők.

14. Az 1-11. igénypontok bármelyike szerinti csecsemőgyógyászati készítmény későbbi életkorban bekövetkező elhízás megelőzésében történő alkalmazásra.

15. Az 1-10. igénypontok bármelyike szerinti csecsemőgyógyászati készítmény alkalmazása 0 és 36 hónap tartományba eső életkorú csecsemő táplálására.