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What do we know about C-section?

When medically necessary, a caesarean section is an effective mean to prevent maternal and newborn mortality.



WHO recommends, C-section should not exceed 10-15% of all births.¹



Infants born by C-section birth might have an increased risk of childhood infections and non-communicable diseases²⁻⁴, like asthma⁵, obesity⁶⁻⁷ and type 2 diabetes later in life.⁸

Infants born by C-section have a delayed gut microbiota colonization, especially by Bifidobacterium and Bacteroides, compared to healthy vaginally born infants.9-15

Bifidobacteria is the most abundant bacteria species in the gut of healthy breastfed infants.20



This delayed colonization, resulting in a compromised microbiota, can last several weeks after birth, up to 1 to 2 years of age.¹¹⁻¹²

Bifidobacterium and Bacteroides are keystone colonizers that have the capability to metabolize Human Milk Oligosaccharides (HMOs) and play a pivotal role in immune function.¹⁷⁻¹⁹

Nutritional strategies offer a great opportunity to **rebalance the compromised microbiota** in early life.

Bifidobacteria levels in C-section born babies





Log 10 cells/g of feces 10 Day 1 7 days 1 **3** years 3 6 month months months Vaginal delivery (n=80) C-section (n=28)

Bifidobacteria levels up to 3 years

Bifidobacteria delay is statistically significant in the first 3 months of life¹¹

However, it can take several months before the gap is closed completely¹²

Breast milk is the ideal food for infants. WHO recommends exclusive breastfeeding for the first 6 months of life,

continuing up to 2 years and beyond with gradual introduction of safe and suitable complementary feeding.

C-section surgery remain essential medical treatments saving millions of lives each year.

Synbiotics support immunity in C-section born babies

Definition

Synbiotics are defined as "a mixture comprising live microorganisms and substrate(s) selectively utilized by host microorganisms that confers a health benefit on the host".21



Synbiotic for C-section born babies

The combination of Bifidobacteria breve M-16V and scGOS/IcFOS (9:1) has been shown to restore delayed colonization by Bifidobacteria in babies born by C-section²²



Probiotic Bifidobacterium breve M-16V (B. breve M-16V)²³

Bifidobacterium breve is a species commonly isolated from the gut of healthy breastfed infants and from human milk. The specific strain B. breve M-16V was selected, because of its well-established clinical data on safety and efficacy in positively modulating the gut microbiome of infants.





Prebiotic scGOS/IcFOS (9:1)

The prebiotic mixture of scGOS/IcFOS (9:1) is designed to closely reflect the quantity, diversity and functionality of HMOs* in breast milk. In more than 40 clinical studies (>90 publications), scGOS/lcFOS has been shown to support a healthy gut and immune system development by:

- Stimulating growth of beneficial bacteria²⁴⁻²⁵
- Suppressing growth of pathogens²⁶⁻²⁷
- Modulation of the gut microbiome and the immune system^{24,28-29}
- Stool softening and frequency closer to that of healthy breast-fed infants^{24,29}
- Reducing the risk of infections^{26,30}

Synbiotics in action The Julius study²²

Randomised, Double-Blind, **Multicentre Study**

(N=153 Infants Born by C-Section)







Restored bifidobacterial gap closer to vaginally

born healthy infants

Reduced

skin symptoms (AE)

- 1. Betran et al., BJOG. 2016;123:667-70.
- 2. Miller JE et al., PLoS medicine, 2020; 17.
- 4. Sevelsted et al. Pediatrics. 2015; 135:e92-8.
- 5. Stokholm et al. Sci Trans Med. 2020; 12:eaax9929
- 6. Zhou et al. Int J Environ Res Public Health. 2020; 17:2003.
- 7. Chojnacki et al. Early Hum dev. 2019; 129:52-59.
- 9. Dominguez-Bello et al. PNAS. 2020; 107:
- 11. Nagpal et al. Sci rep. 2017; 7:10097.
- 12. Korpela-de Vos Curr Opin Microbiol. 2018; 44:70-78.
- 13. Jakobsson et al. Gut. 2014; 63:559-66.
- 14. Martin, et al. PLOS one. 2016; 11:e0158498.
- 15. Lay C et al., BMC microbiology, 2021. 21: 191.

- 18. Troy EB and DL Kasper, Front Biosci (Landmark Ed),

- 22. Chua MC et al., J Pediatr Gastroenterol Nutr. 2017. 65:102-106.
- 23. Moro G et al., J Pediatr Gastroenterol Nutr, 2002. 34: p. 291-5.
- 24. Knol J et al., 2005. 40: p. 36-42.
- 25. Bruzzese E et al., Clin Nutr, 2009. 28: p. 156-61.
- 26. Knol J et al., J Pediatr Gastroenterol Nutr, 2003. 36: p. 566.
- 27. Miqdady M et al., Pediatric gastroenterology, hepatology & nutrition, 2020. 23: p. 1-14.
- 28. Boehm G and G Moro, J Nutr, 2008. 138,1818s-1828s.

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- 29. Arslanoglu S et al., J Nutr, 2007. 1372420-4.
- 30. Chatchatee P et al., J Pediatr Gastroenterol Nutr, 2014. 58 428-437.