

Maturing the immune system

probiotics for disease prevention

Medicine<sup>fx</sup>

90%

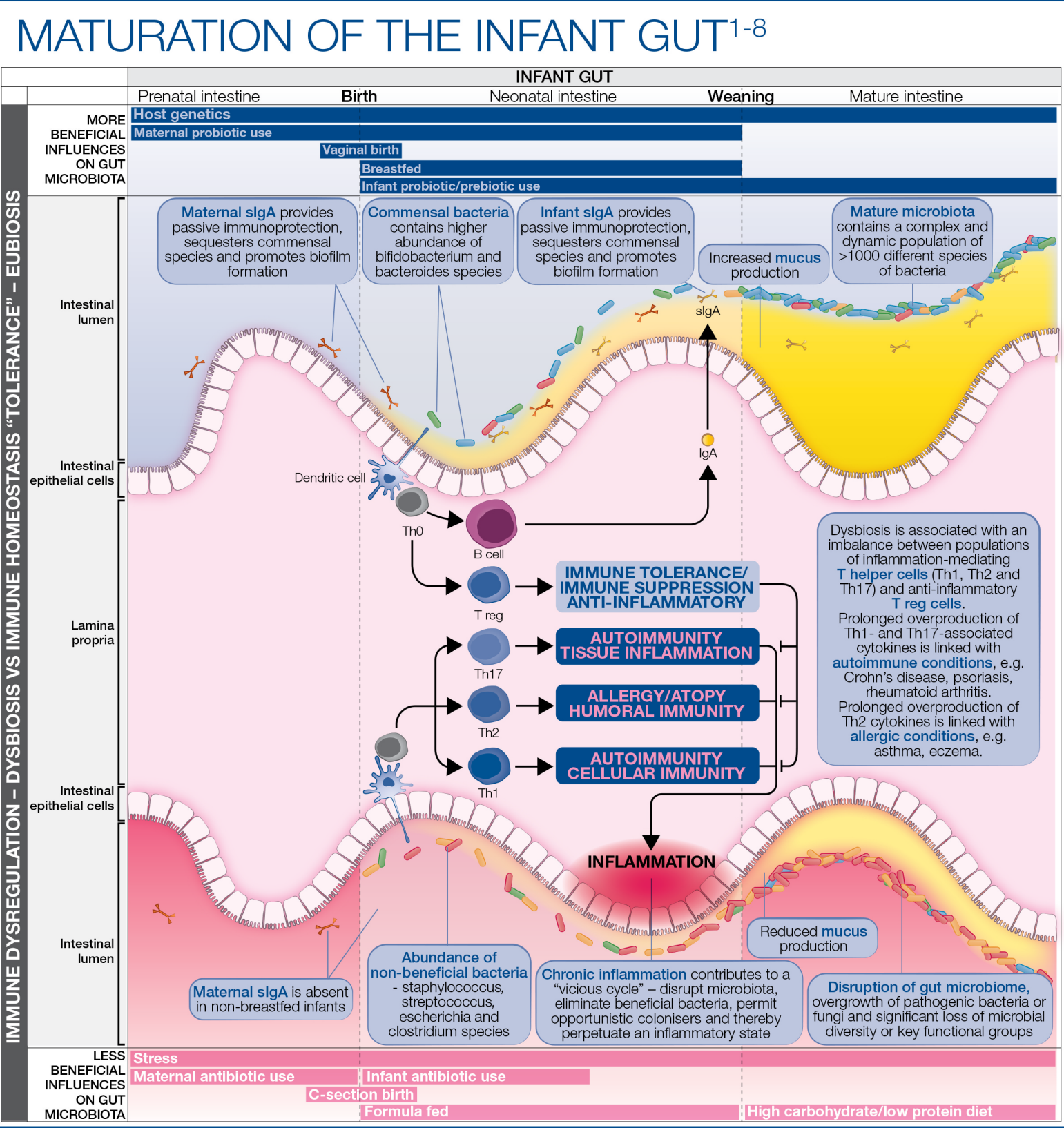
of the cells in the HUMAN BODY are BACTERIAL<sup>1</sup>

The MICROBIOME has a PROFOUND INFLUENCE on HOST HEALTH and the development of the IMMUNE RESPONSE<sup>1</sup>

IMMUNE DISORDERS: FAILURE in the development of a BALANCED IMMUNE RESPONSE<sup>1</sup>

Over the last 50 years there has been a STEEP INCREASE in the incidence of ATOPIC AND AUTOIMMUNE DISEASE<sup>2</sup>

PREDISPOSITION TO DISEASE may, in part, be determined IN UTERO<sup>1</sup>



### MATERNAL SOURCES OF BACTERIA<sup>7</sup>

**Mouth**  
Bacteria of oral origin found in amniotic fluid.

**External breast**  
Skin of the breast is an external source of bacteria for breast milk.

**Internal breast**  
Hundreds of microbes identified in breast milk. Abundant genera include: staphylococcus, streptococcus, serratia, pseudomonas, corynebacterium, ralstonia, propionibacterium, sphingomonas, bradyrhizobiaceae. The bacterial composition of breast milk changes over time – milk produced immediately after labour contains more lactic acid bacteria (staphylococcus, streptococcus and lactococcus). Milk after 6 months of lactation shows an increase in bacteria typically found in the oral cavity (veillonella, leptotrichia and prevotella, typically found in the oral cavity). An entero-mammary pathway exists that transports bacteria from the maternal gut to the mammary gland via dendritic cells, lymph and blood circulation.

**Uterus**  
Bacteria found in umbilical cord blood, the placenta, amniotic fluid, foetal membranes and meconium of healthy infants. Meconium harbours microbes commonly found in gastrointestinal tract: enterococcus and escherichia.

**Vagina**  
Delivery mode (vaginal delivery versus C-section) has been shown to influence postnatal gut microbiome. Microbiota of vaginally born infants resembles that of their mother’s vagina, while that of C-section infants is representative of skin microbes.

### CONSEQUENCES OF IMMUNE IMBALANCES<sup>1-8</sup>

**Balanced immune system**

**HOMEOSTASIS**

**T helper cells (inflammatory)** vs **T reg cells (anti-inflammatory)**

**AUTOIMMUNE** vs **ALLERGY**

**Imbalanced immune system**

**Th1 (IFN-γ, TNF-β)** vs **Th2 (IL-5, IL-5, IL-10)**

**Th1 (IFN-γ, TNF-β)** vs **Th2 (IL-5, IL-5, IL-10)**

**Th17 (IL-17, IL-21, IL-22)** vs **T reg (IL-10, TGF-β)**

**Autoimmune conditions e.g. Crohn’s disease**

**Allergic conditions e.g. asthma**

**Autoimmune conditions e.g. rheumatoid arthritis**

**Legend:**

- IgA: immunoglobulin A; sIgA: secretory immunoglobulin A; Th0: naive T cell; T reg: regulatory T cell; Th1: T-helper 1; Th2: T helper 2; Th17: T helper 17; IFN-γ: interferon-gamma; TNF-β: tumour necrosis factor-beta; IL-4: interleukin 4; IL-5: interleukin 5; IL-10: interleukin 10; IL-17: interleukin 17; IL-21: interleukin 21, IL-22: interleukin 22; TGF-β: transforming growth factor-beta