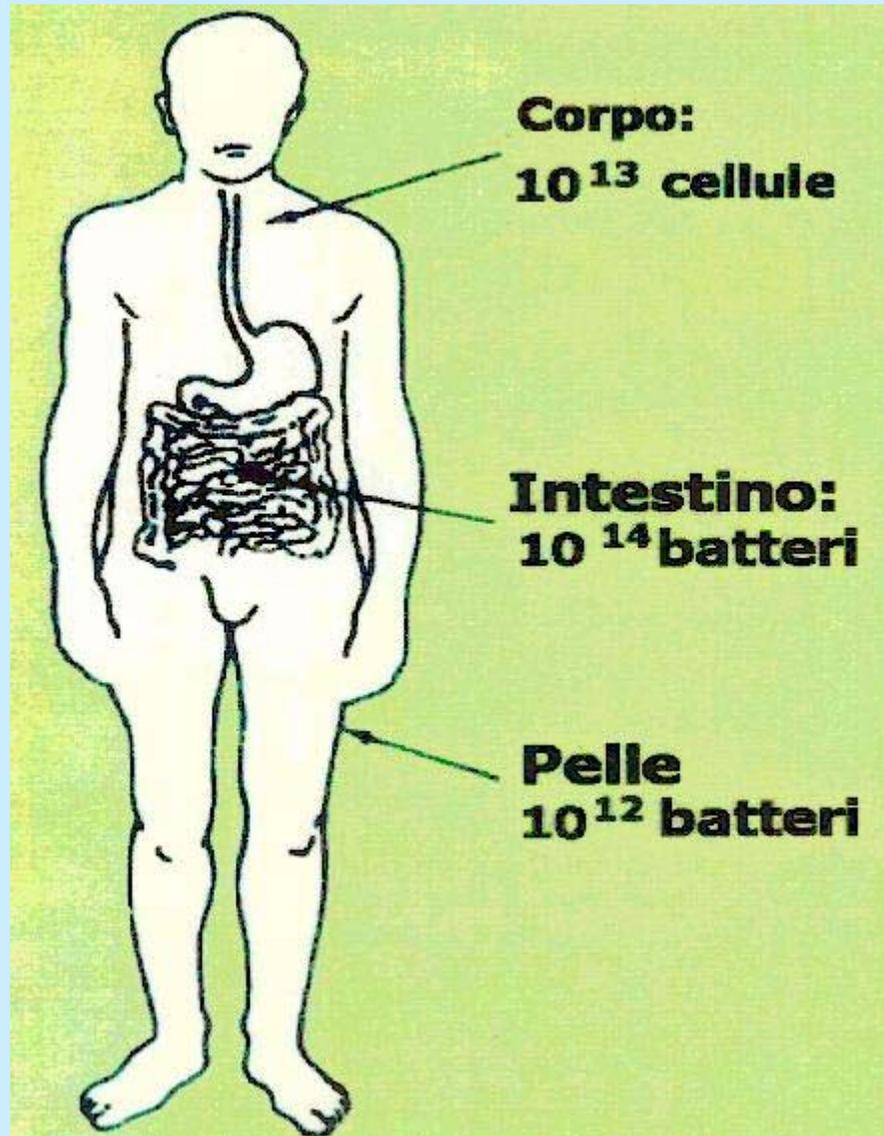


# **L'effetto barriera della flora batterica intestinale-**

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Piacenza**



# Microflora del neonato...

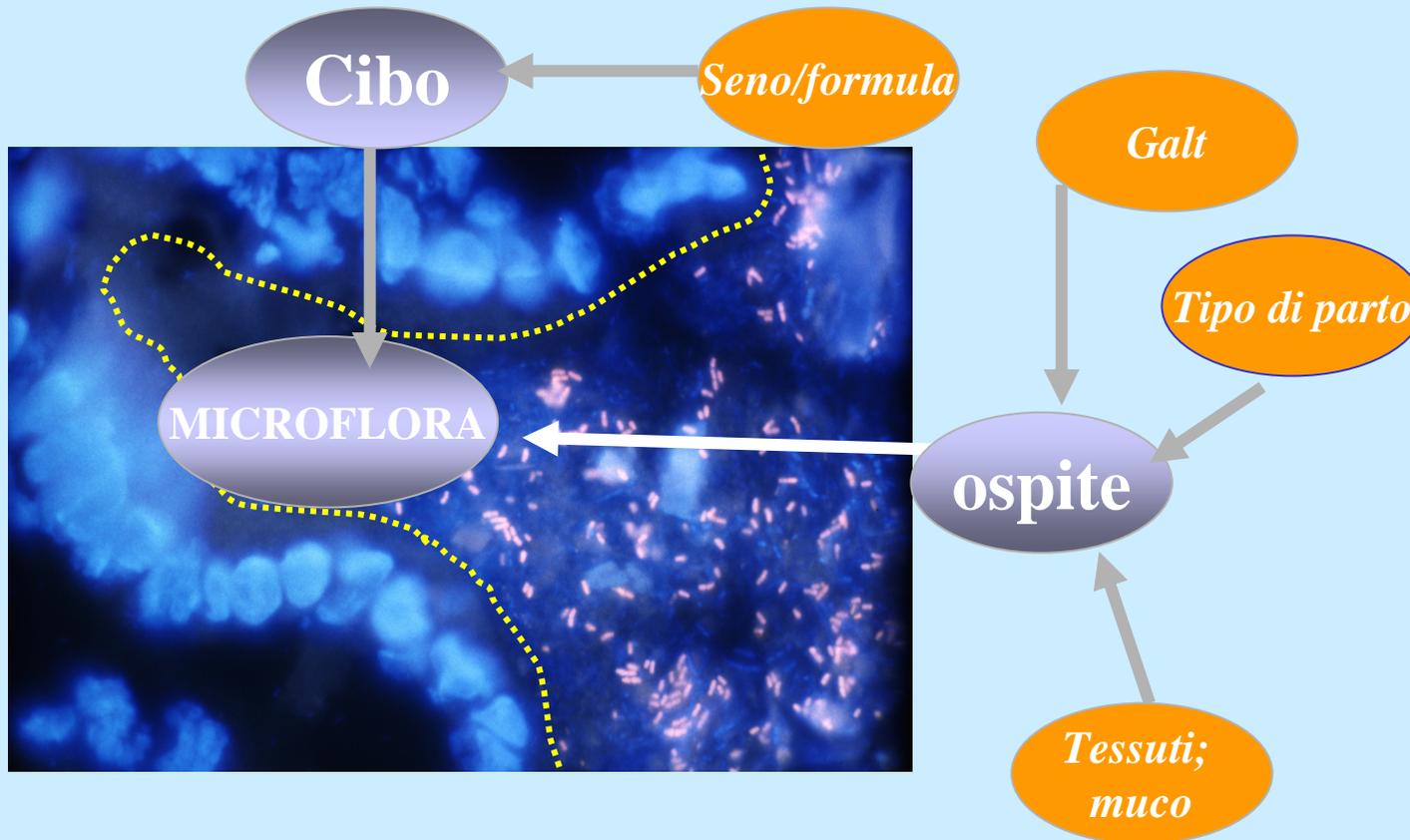


Dal deserto...

alla jungla



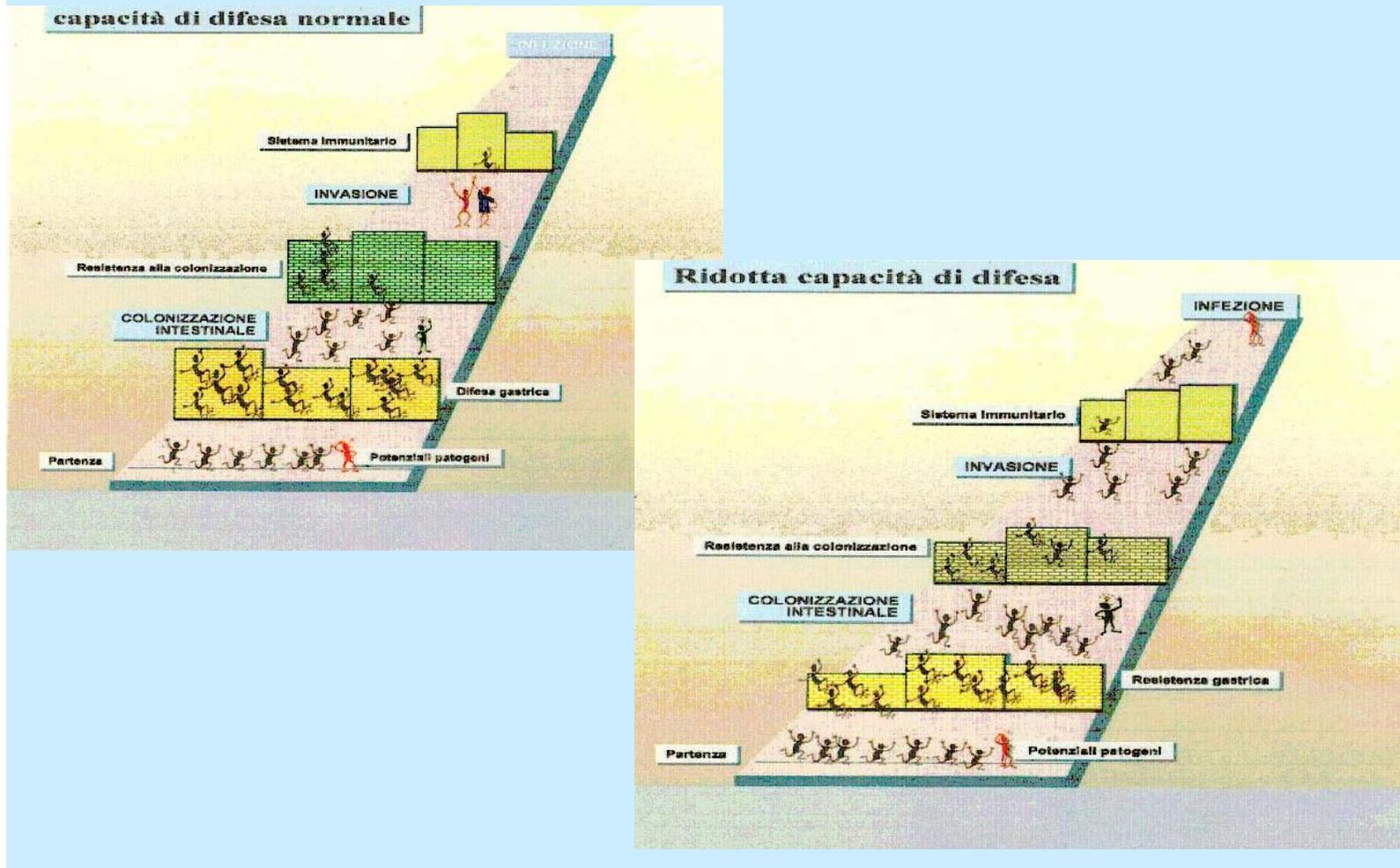
# “Driving forces”



## logN/g di batteri nel tratto intestinale

	Uomo	Ratto	Suini	Topo	Conigli
<b>Cavità orale</b>	<b>7-8</b>	<b>7-8</b>	<b>7-8</b>	<b>7-8</b>	<b>7-8</b>
<b>Stomaco</b>	<b>&lt;3</b>	<b>7</b>	<b>6-7</b>	<b>7</b>	<b>4</b>
<b>Digiuno</b>	<b>&lt;3</b>	<b>7</b>	<b>8</b>	<b>7</b>	<b>4</b>
<b>Ileo</b>	<b>5</b>	<b>8</b>	<b>10</b>	<b>8</b>	<b>8</b>
<b>Colon</b>	<b>11</b>	<b>11</b>	<b>11</b>	<b>11</b>	<b>11</b>

# La microflora ha un effetto barriera



Competizione fisica (adesione-probiotici)

Competizione nutrizionale (prebiotici)

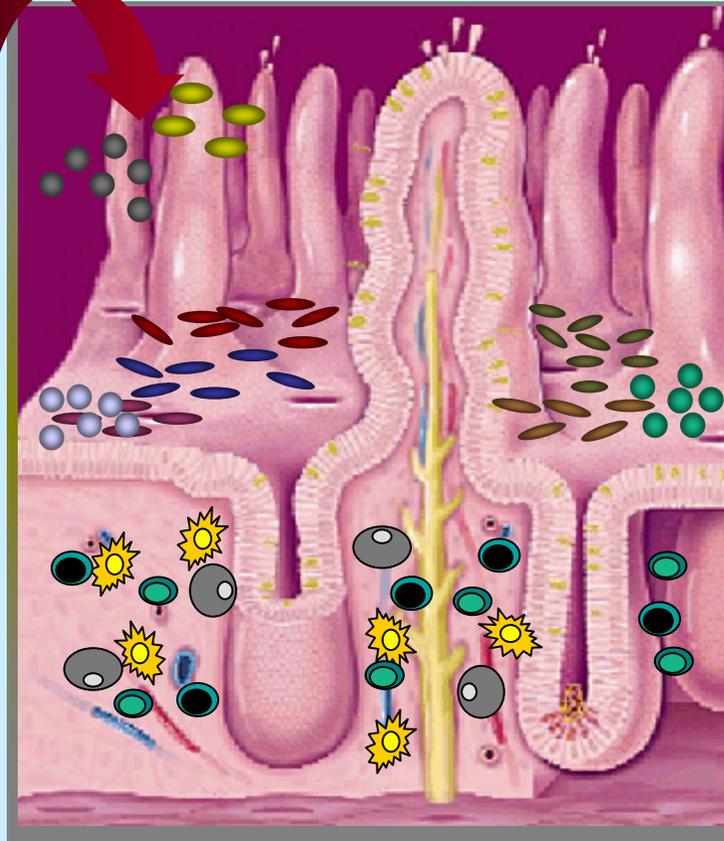
*Inibizione (probiotici)*



## Intestinal ecosystem

Flora transiente

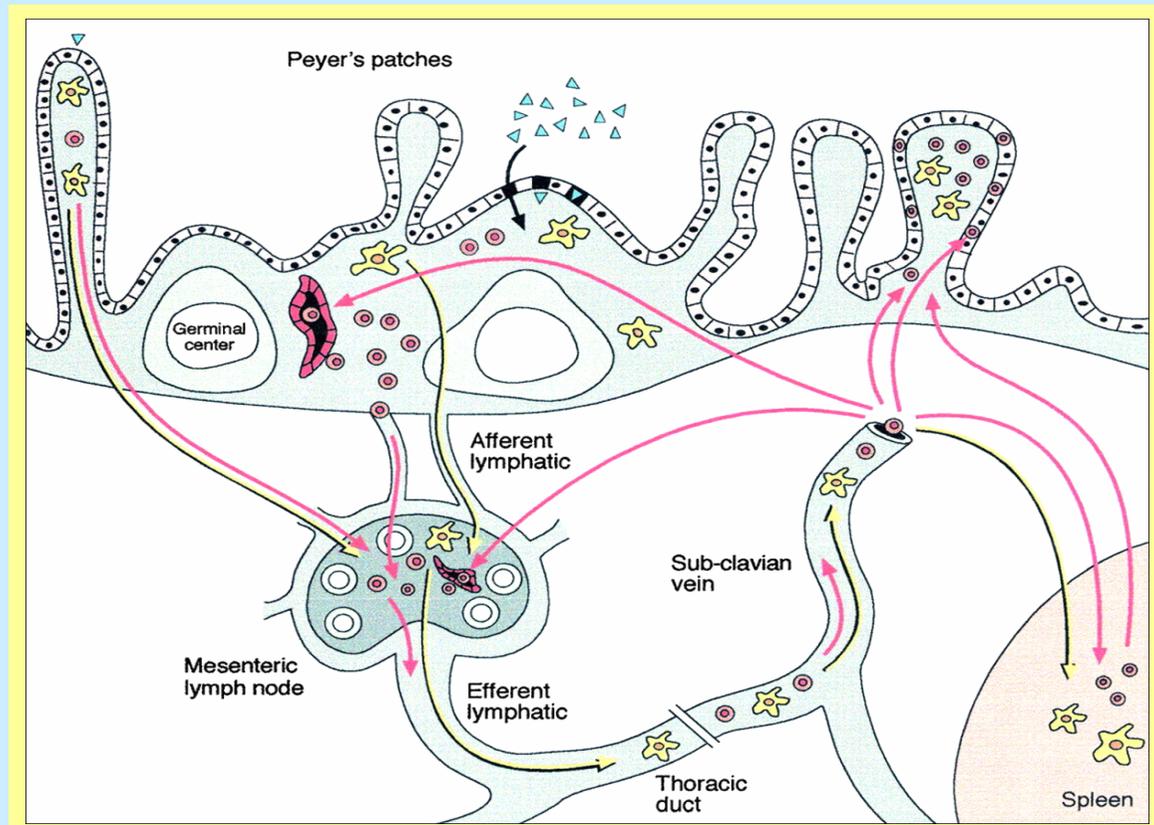
Cellule del S.I.  
Macrophages  
Dendritic cells  
B-lymphocytes  
T-lymphocytes



Flora residente

*Bacteroides*  
*Bifidobacterium*  
*Clostridium*  
*Escherichia coli*  
*Enterococcus*  
*Lactobacillus*

## Induzione della risposta immune



## Association of *Lactobacillus* spp. with Peyer's Patches in Mice

LAURA PLANT AND PATRICIA CONWAY\*

*School of Microbiology and Immunology, University of New South Wales, Sydney, Australia, 2052*

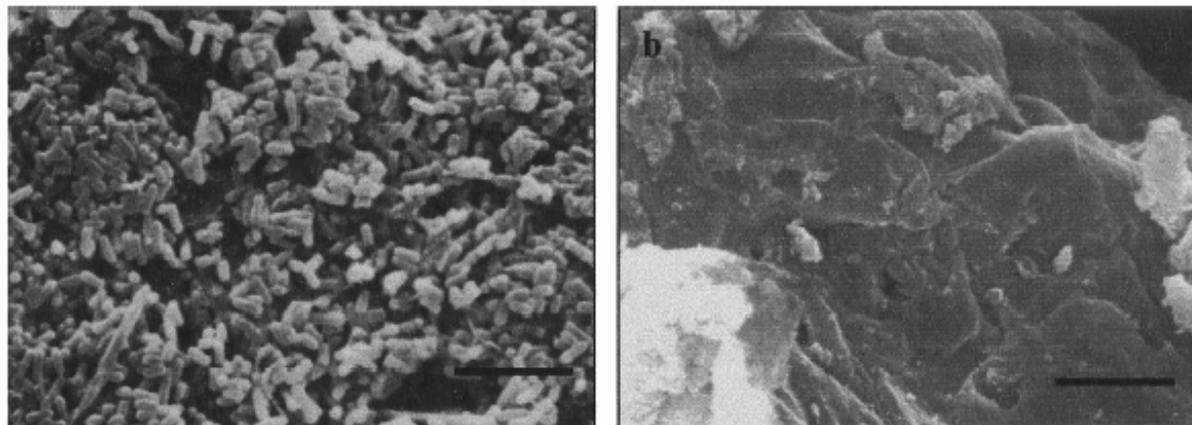


FIG. 1. Scanning electron micrograph of mouse tissue after incubation in *L. fermentum* KLD cell suspension. (a) Typical aggregates of *L. fermentum* KLD-like cells on the FAE of the Peyer's patch; (b) absence of cells on nonlymphoid villous intestine. Bars, 10  $\mu$ m.

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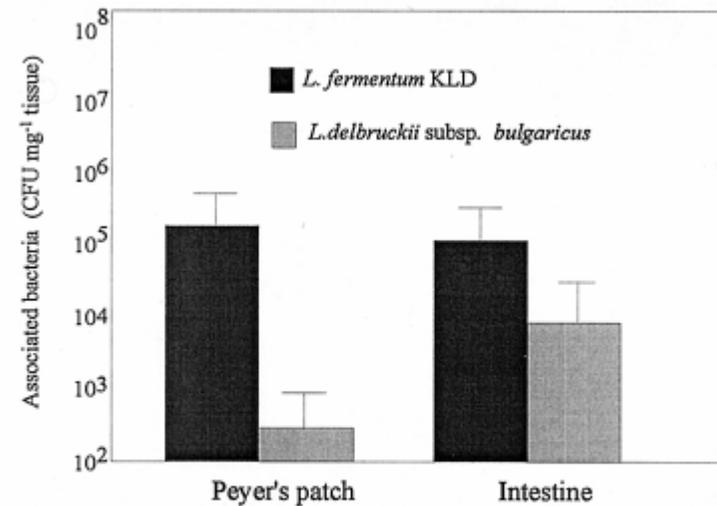


FIG. 4. Association of *L. fermentum* and *L. delbrueckii* subsp. *bulgaricus* with the Peyer's patches and nonlymphoid villous intestine 2 h following administration of an oral dose of  $10^9$  viable *Lactobacillus* cells. Results are expressed as numbers of CFU per milligram (wet weight) of tissue ( $n = 35$  per treatment) as the mean value. Error bars indicate standard deviations.

## Probiotici e GALT

TABLE 2. Association of *Lactobacillus* strains with mouse Peyer's patches and nonlymphoid intestinal tissue

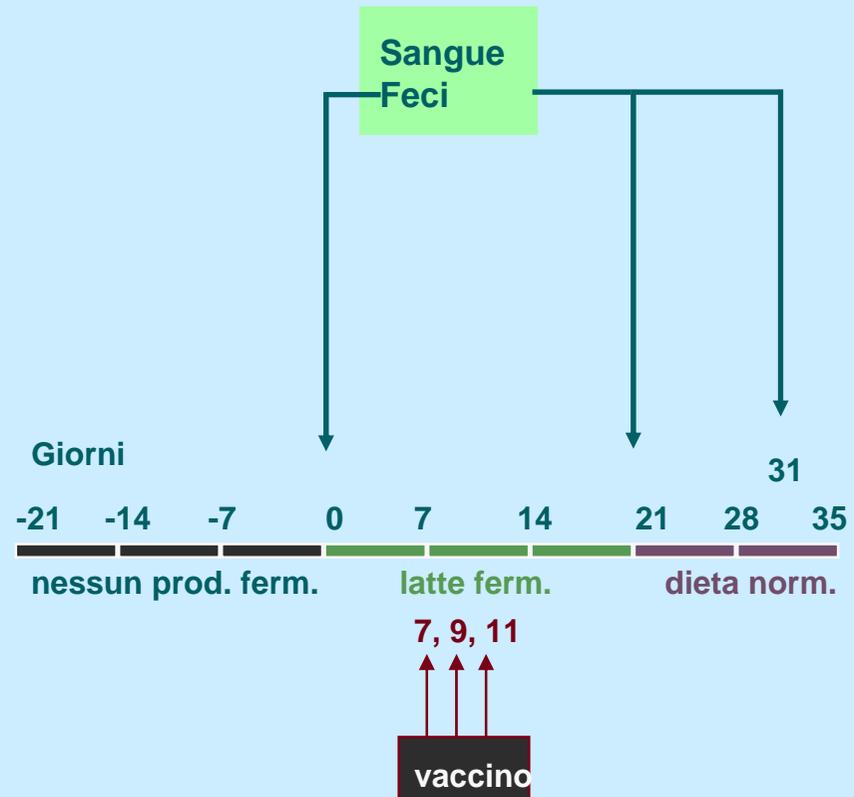
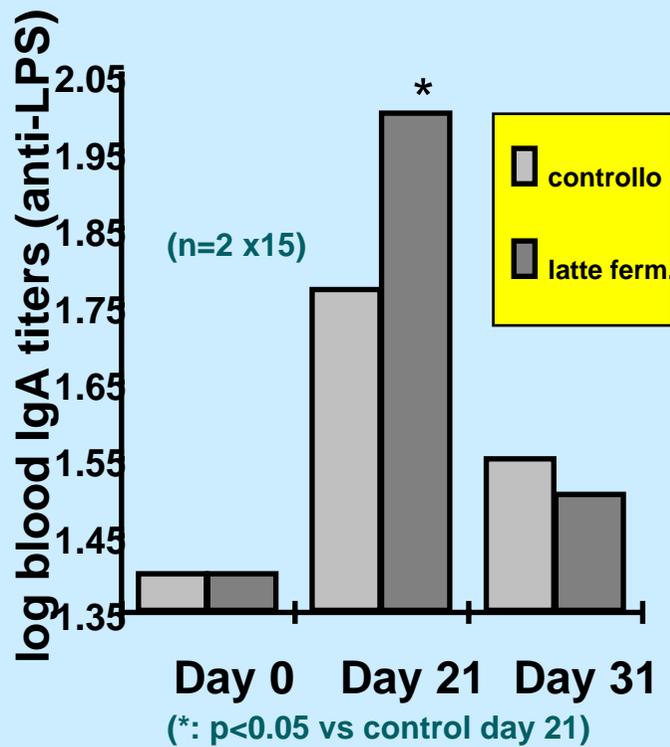
Strain	Peyer's patch association <sup>a</sup>	Nonlymphoid intestine association <sup>a</sup>
<i>Lactobacillus</i> sp. strain 003	-	-
<i>Lactobacillus</i> sp. strain 004	-	-
<i>Lactobacillus</i> sp. strain 005	++	+
<i>Lactobacillus</i> sp. strain 006	++	+
<i>Lactobacillus</i> sp. strain 008	-	+
<i>L. fermentum</i> KLD	++	-
<i>L. acidophilus</i>	+	-
<i>L. fermentum</i> LMN	-	-
<i>L. fermentum</i> 104S	+	-
<i>Lactobacillus</i> sp. strain HBL	+	++
<i>L. paracasei</i> 43338	++	++
<i>L. salivarius</i> 43321	-	-
<i>L. paracasei</i> 42319	+	+
<i>Lactobacillus</i> sp. strain 433121	-	-
<i>L. bulgaricus</i>	-	-
<i>L. salivarius</i> subsp. <i>salivarius</i>	+	-

<sup>a</sup> The association of *Lactobacillus* spp. to small intestinal tissue, as visualized by scanning electron microscopy: -, no association; +, low level of association (5 to 50 bacteria/field); ++, high level of association (>50 bacteria/field).

Plant e Conway, 2001



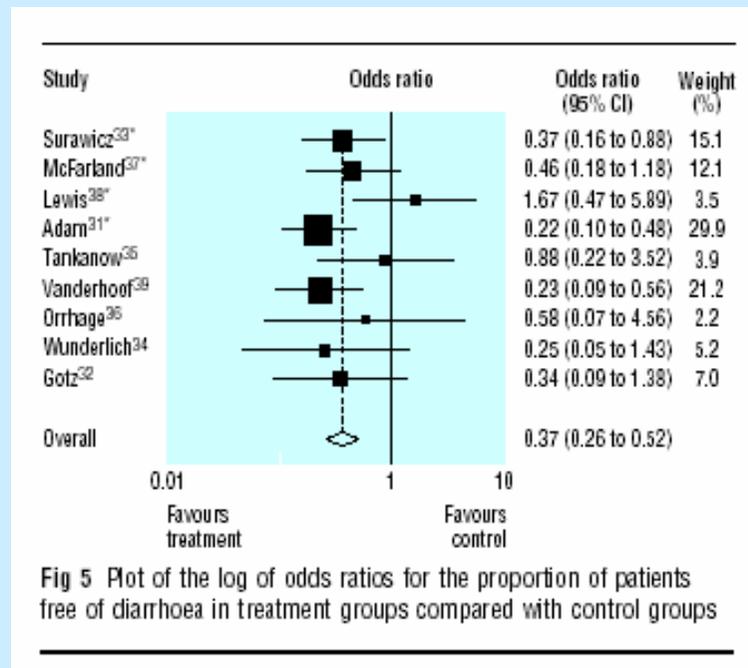
## IgA serica contro *Salmonella typhi* Ty21a a seguito di ingestione di latte fermentato contenente La1



# Effetto barriera nelle meta-analisi

BMJ VOLUME 324 8 giugno 2002

Importanza della dose





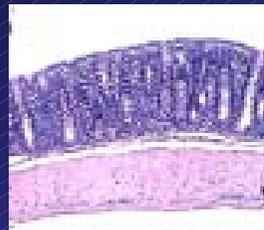
# Lack of effect of *L. crispatus* MU5 on DSS colitis Importanza del ceppo



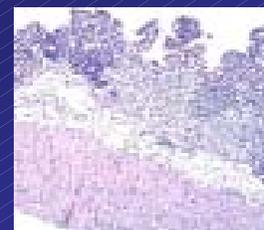
CONT



DSS



DSS + M247



DSS + MU5

# Bacterial ecology and allergy?

[www.fems-microbiology.com](http://www.fems-microbiology.com)

Characterizing the composition of intestinal microflora as a prospective treatment target in infant allergic disease

Pirkka V. Kirjavainen <sup>a,\*</sup>, Effie Apostolou <sup>a</sup>, Taina Arvola <sup>c</sup>, Seppo J. Salminen <sup>a</sup>,  
Glenn R. Gibson <sup>d</sup>, Erika Isolauri <sup>b</sup>

FEMS Immunology and Medical Microbiology 32 (2001) 1–7

# Bacterial ecology and allergy?

4

*P. V. Kirjavainen et al. / FEMS Immunology and Medical Microbiology 32 (2001) 1–7*

**Table 1**  
The presence of selected predominant gut genera in the fecal flora of atopic and healthy infants as assessed by FISH during exclusive breast-feeding and after cessation of breast-feeding

Bacterial group	Time of detection	Number of cells per gram of feces (the log mean <sup>a</sup> of each group with 95% CI in parentheses)						
		Healthy infants (n = 10)	Atopic infants				Positive CM challenge (n = 6)	Negative CM challenge (n = 9)
			All <sup>b</sup> (n = 15)	GI symptoms during BF <sup>c</sup> (n = 5)	No GI symptoms during BF <sup>c</sup> (n = 10)			
Bifidobacteria	During BF	<b>10.18 (9.73–10.47)</b>	9.71 (8.89–10.53)	<b>9.43 (8.73–10.13)<sup>cd</sup></b>	<b>10.19 (9.72–10.67)</b>	10.00 (8.97–11.02)	9.52 (8.15–10.89)	
	After BF	9.40 (9.03–9.78) <sup>f</sup>	8.92 (8.41–9.43) <sup>f</sup>					
Bacteroides	During BF	<b>6.99 (5.37–8.60)</b>	8.23 (7.11–9.34)	7.96 (5.56–10.36)	8.36 (6.82–9.90)	<b>9.57 (7.59–11.54)<sup>d</sup></b>	7.70 (6.07–9.33)	
	After BF	8.02 (6.78–9.26)	8.27 (7.60–8.94)					
Lactobacilli/enterococci	During BF	6.51 (4.46–8.55)	7.72 (6.84–8.59)	7.93 (7.26–8.60)	7.61 (6.23–8.99)	7.34 (4.81–9.87)	7.96 (7.38–8.55)	
	After BF	6.47 (4.84–8.10)	7.33 (6.59–8.07)					
<i>C. histolyticum</i>	During BF	6.99 (5.88–8.11)	6.47 (5.29–7.65)	7.08 (5.64–8.53)	6.16 (4.39–7.93)	6.62 (5.16–8.08)	6.37 (4.38–8.36)	
	After BF	7.46 (6.19–8.73)	6.95 (6.10–7.79)					
Total cell counts	During BF	<b>10.65 (10.44–10.87)</b>	<b>10.38 (10.21–10.55)<sup>d</sup></b>	10.23 (9.85–10.61)	10.56 (10.29–10.83)	10.51 (10.28–10.73)	10.41 (10.05–10.77)	
	After BF	10.22 (10.04–10.41) <sup>f</sup>	10.27 (10.12–10.41)					
Total IgE in serum (kU l <sup>-1</sup> )	During BF	ND	79 (10–46)	55 (13–43)	96 (0–48)	<b>113 (12–60)<sup>c</sup></b>	<b>12 (0–19)</b>	
SCORAD score	During BF	ND	27 (9–38)	29 (15–45)	25 (6–33)	<b>32 (18–44)<sup>c</sup></b>	<b>17 (4–28)</b>	

# From ecology to applications?

Gut, 2002

- **Aberrant composition of gut microbiota of allergic infants: a target of bifidobacterial therapy at weaning?**

- P V Kirjavainen<sup>1</sup>, T Arvola<sup>2</sup>, S J Salminen<sup>1</sup> and E Isolauri<sup>3</sup>

**Background:** Recent data have outlined a relationship between the composition of the intestinal microflora and allergic inflammation, and demonstrated the competence of probiotics in downregulation of such inflammation.

- **Aims:** Our aims were to characterise the relationship between gut microbes and the extent of allergic sensitisation and to assess whether the efficacy of bifidobacterial supplementation in the treatment of allergy could relate to modulation of the intestinal microbiota.
- **Methods:** This randomised study included 21 infants with early onset atopic eczema of whom eight were intolerant (highly sensitised group (HSG)) and 13 tolerant (sensitised group (SG)) to extensively hydrolysed whey formula (EHF). In the SG, six were weaned to EHF without (placebo group (PG)) and seven to EHF with *Bifidobacterium lactis* Bb-12 supplementation (bifidobacteria treated group (BbG)). The faecal microflora of infants in the HSG was analysed only before weaning whereas in the SG the faecal microflora was analysed both before and after weaning.
- **Results:** Infants in the HSG had greater numbers of lactobacilli/enterococci than those in the SG. Serum total IgE concentration correlated directly with *Escherichia coli* counts in all infants and with bacteroides counts in the HSG, indicating that the presence of these bacteria is associated with the extent of atopic sensitisation. The effect of supplementation was characterised as a decrease in the numbers of *Escherichia coli* and protection against an increase in bacteroides numbers during weaning.
- **Conclusions:** These data indicate that bifidobacterial supplementation appears to modify the gut microbiota in a manner that may alleviate allergic inflammation. Further studies are needed to confirm this conclusion.

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# Conclusions

**The experts agreed that adequate scientific evidence exists to indicate that there is potential for the derivation of health benefits from consuming food containing probiotics. However, it was felt that additional research data are needed to confirm a number of these health benefits in humans, applying a systematic approach and following the guidelines for the assessment of probiotics suggested in this report. (FAO/WHO, 2001)**