



河南师范大学
NENAN NORMAL UNIVERSITY

水产动物肠道菌群研究进展

2017. 4. 28



厚德博学·止于至善



主要内容

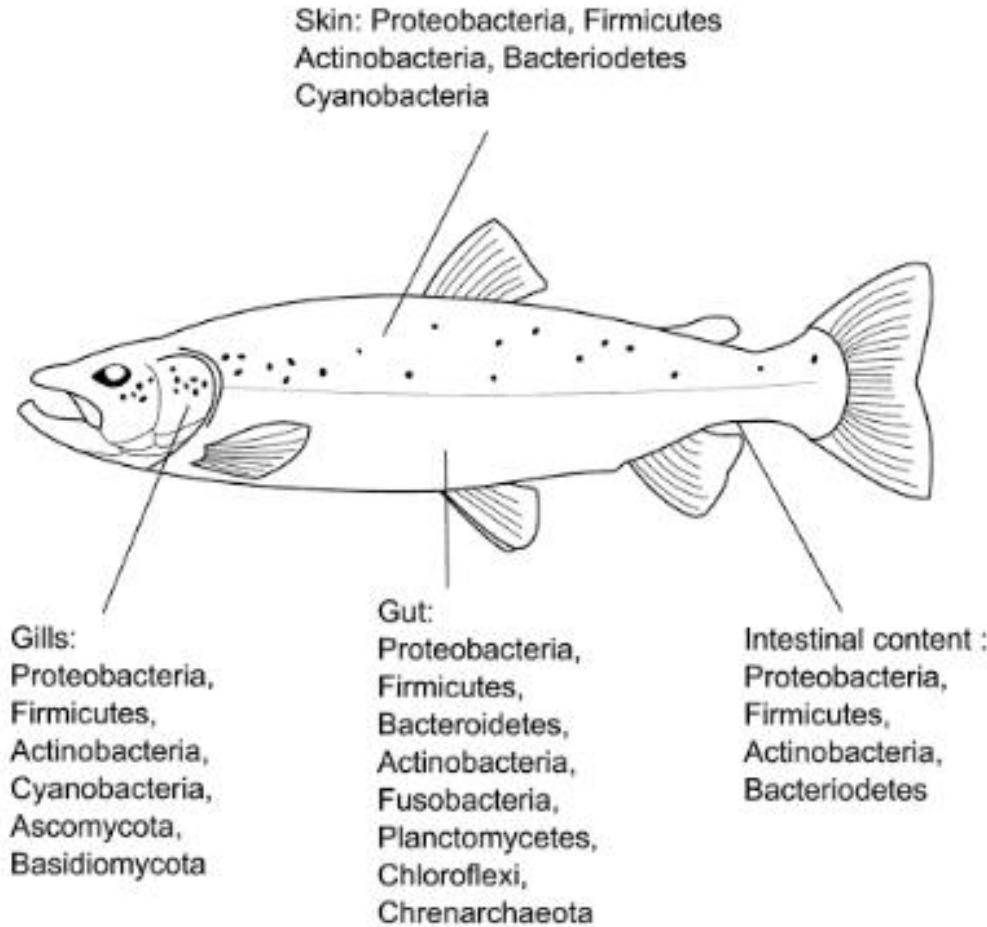
Part I 鱼类肠道微生物的结构

Part II 鱼类肠道微生物功能研究

Part III 评价微生物群落的方法



Part I 鱼类肠道微生物的结构



鱼类肠道微生物组成

Proteobacteria
Bacteroidetes
Actinobacteria
Firmicutes
Fusobacterium





Part I 鱼类肠道微生物的结构

Environment and diet



marine

freshwater





Part I 鱼类肠道微生物的结构



Diet

Carnivorous
Omnivorous
Herbivorous



Part I 鱼类肠道微生物的结构

Marine fish

兼性厌氧菌

Vibrio, (弧菌属)

Pseudomonas, (假单胞菌属)

Acinetobacter, (不动杆菌属)

Corynebacterium (棒状杆菌属)

Alteromonas (单胞菌属)

Flavobacterium (黄杆菌属)

Micrococcus (微球菌属)

Freshwater fish

Aeromonas, (其单胞菌属)

Pseudomonas, (假单胞菌属)

Bacteroides, (拟杆菌属)

Plesiomonas, (邻单胞菌属)

Enterobacteriaceae (肠杆菌属)

Micrococcus (微球菌属)

Acinetobacter (不动杆菌属)

Clostridium (梭菌属)

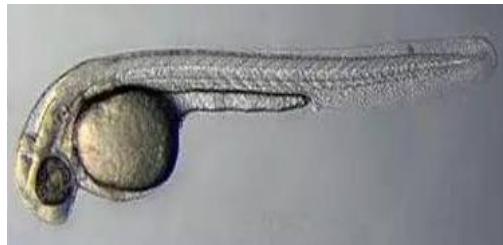
Fusarium (镰刀菌属)





Part I 鱼类肠道微生物的结构

Establishment of GI microbiota

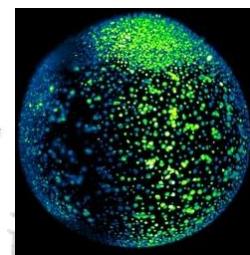
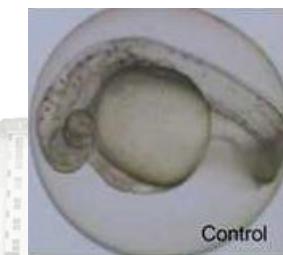


Proteobacteria
Actinobacteria

Proteobacteria
Actinobacteria
Firmicutes
Bacteroidetes

2-5 years old
Similar to adult

- 1. Eggs
- 2. Larval rearing water
- 3. Live feed

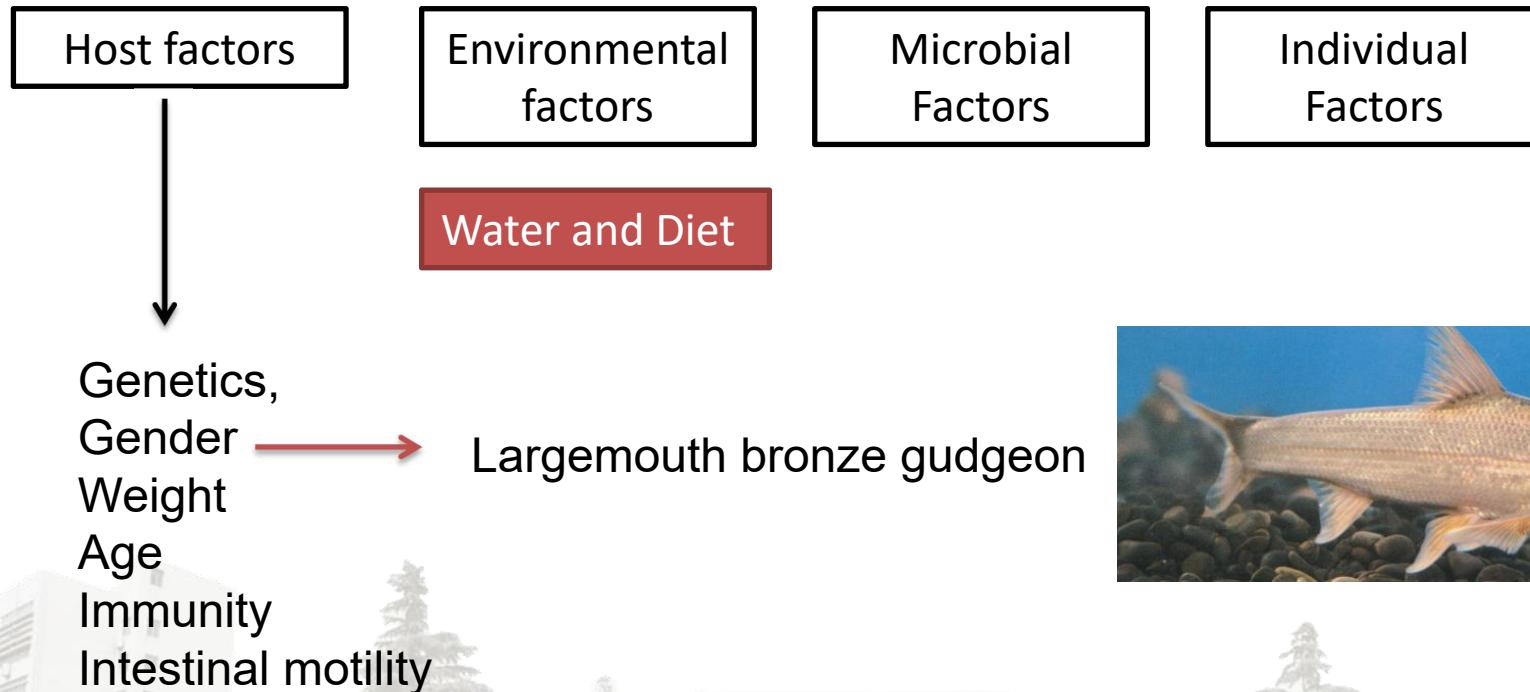


*Cytophaga,
Flavobacterium
pseudomonas*



Part I 鱼类肠道微生物的结构

Factors affecting GI microbiota of fish





Part I 鱼类肠道微生物的结构

The influence of water



Temperature

Diversity and seasonal changes in lactic acid bacteria
in the intestinal tract of cultured freshwater fish

Tatsuro Hagi, Daichi Tanaka, Yasutada Iwamura, Takayuki Hoshino*

Institute of Applied Biochemistry, University of Tsukuba, 1-1-1 Ten-nodai Tsukuba, Ibaraki, 305-8572, Japan

Received 8 December 2003; received in revised form 15 January 2004; accepted 15 January 2004

Seasonal Variation

ORIGINAL ARTICLE

**Pyrosequencing-based characterization of gastrointestinal
bacteria of Atlantic salmon (*Salmo salar* L.) within a
commercial mariculture system**

K.Z. Zarkasi^{1,2}, G.C.J. Abell³, R.S. Taylor³, C. Neuman⁴, E. Hatje⁴, M.L. Tamplin¹, M. Katouli⁴ and
J.P. Bowman¹

**The effect of diet and environmental temperature on
the faecal microbiota of farmed Tasmanian Atlantic
Salmon (*Salmo salar* L.)**

Christina Neuman¹, Eva Hatje¹, Kamarul Z Zarkasi², Richard Smullen³, John P Bowman² &
Mohammad Katouli¹



Part I 鱼类肠道微生物的结构

Salinity

Response of gut microbiota to salinity change in two euryhaline aquatic animals with reverse salinity preference

Meiling Zhang^a, Yuhong Sun^a, Yukun Liu^a, Fang Qiao^a, Liqiao Chen^a, Wen-Tso Liu^b, Zhenyu Du^a, , Erchao Li^a, ,

decreased when the host facing hyposaline or hypersaline stress. The alteration of intestinal microbiota was likely attributed to the environmental selection for microbes that could grow better under high or low salinity, the host response to the salinity stress and subsequent stress exerting on the gut microbiota, or both.

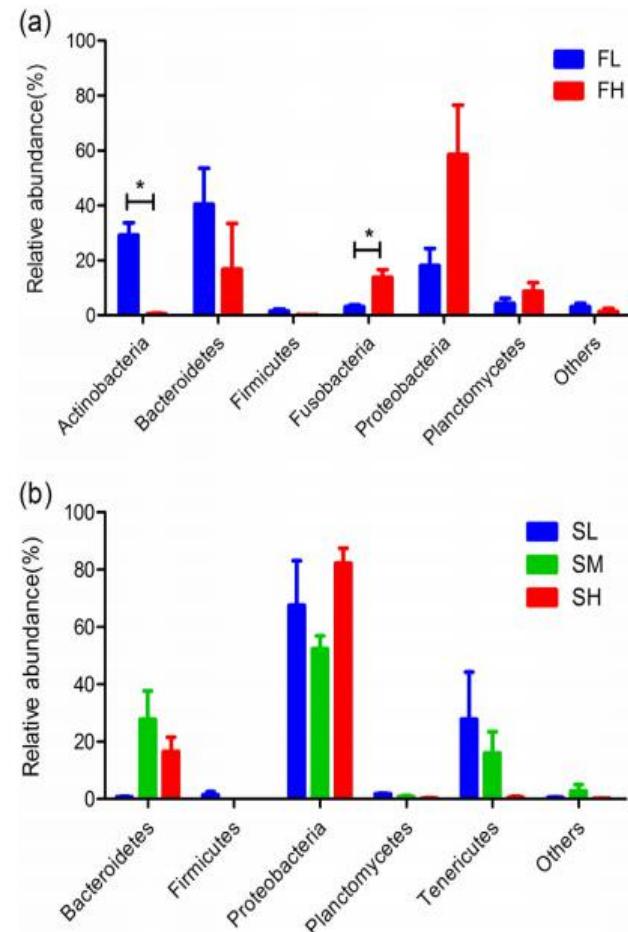


Fig. 2. Relative abundance of the intestinal flora at phylum level in Nile tilapia and Pacific white shrimp reared at different salinities. The most dominant phyla (> 1% of the total sequences) were shown. Asterisk (*) represents the difference between groups is significant based on student t-test, $p < 0.05$.



Part I 鱼类肠道微生物的结构

The influence of diet and feeding habit



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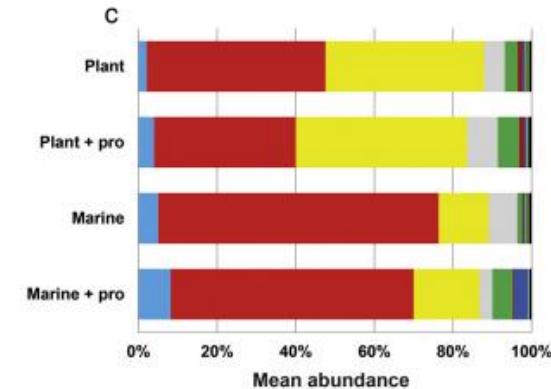
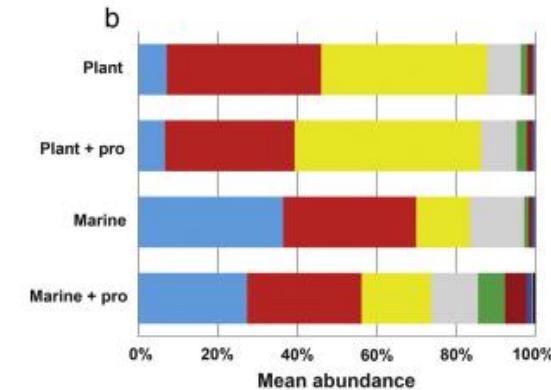
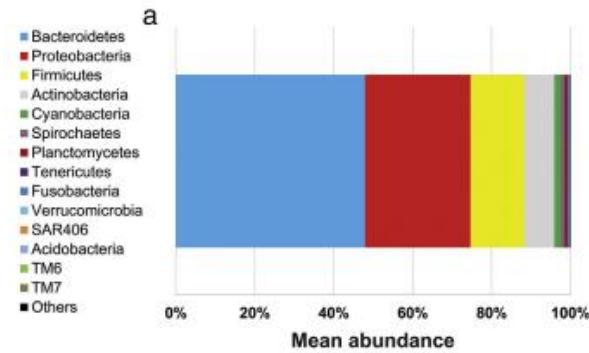
The development of the gut microbiota in rainbow trout (*Oncorhynchus mykiss*) is affected by first feeding and diet type



H.-C. Ingerslev ^{a,*}, L. von Gersdorff Jørgensen ^b, M. Lenz Strube ^a, N. Larsen ^c, I. Dalsgaard ^a, M. Boye ^a, L. Madsen ^a

f.f: 1 day before first feeding
26 days after f.f
49 days after f.f

Firmicutes dominate the plant source oil
Proteobacteria dominate the fish oil





Part I 鱼类肠道微生物的结构

The regional difference of GI microbiota

Density

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鮰鱼消化道细菌群落结构及多样性初步研究

陶诗，王健鑫，刘雪珠，何芳芳，俞凯成

(浙江海洋学院，海洋生物资源与分子工程实验室，浙江舟山 316000)

摘要：采用常规分离纯化与构建 16S rDNA 克隆文库相结合的方法对**鮰鱼** (*Miichthys miiuy*) 消化道细菌多样性进行了初步研究。结果表明，**鮰鱼**消化道各部位可培养细菌的菌落数量从大到小依次为前肠 > 幽门胃 > 中肠 > 后肠 > 前胃 > 后胃 > 口咽腔，其中肠道部位菌落数量最多。分离细菌菌株主要为杆状细菌和革兰氏阴性菌，对其中的 50 株典型菌株进行分子鉴定，发现它们主要属于变形菌门的 γ -变形菌纲(占 52.7%) 和 β -变形菌纲(占 36.8%)，以及厚壁菌门的芽孢杆菌纲(占 10.5%)；不可培养细菌 16S rDNA 文库克隆子主要属于 γ -变形菌纲、 α -变形菌纲和脱铁杆菌纲，部分序列与已报道的物种相似性较低，说明消化道中可能存在新的有待开发的肠道菌株。



Part I 鱼类肠道微生物的结构

Composition varied among different GI tract regions

**Bacterial microflora in the gastrointestinal tract of Nile tilapia,
Oreochromis niloticus, cultured in a semi-intensive system**

Lígia Maria Molinari¹, Denise de Oliveira Scoaris², Raíssa Bocchi Pedroso³, Nilza de Lucas Rodrigues Bittencourt⁴, Celso Vataru Nakamura⁴, Tânia Ueda-Nakamura⁴, Benício Alves de Abreu Filho⁴ and Benedito Prado Dias Filho^{4*}

Table 1. Predominant bacteria isolated from gastrointestinal tract of the tilapia *Oreochromis niloticus*.

Microorganism	Stomach	Anterior gut	Posterior gut
<i>Aeromonas hydrophila</i>	0.6	0	0
<i>Aeromonas veronii</i>	0	0.2	0.3
<i>Burkholderia cepacia</i>	0	0	3.7
<i>Chromobacterium violaceum</i>	90.0	55.5	0
<i>Citrobacter freundii</i>	0	0	13.0
<i>Escherichia coli</i>	7.4	0	0
<i>Flavimonas oryzihabitans</i>	1.1	0	0
<i>Plesiomonas shigelloides</i>	0.6	4.8	76.0
Unidentified sp	0	39.3	6.6

Values are mean percentages from four independent experiments



Part II 鱼类肠道微生物功能研究

Gnotobiotic zebrafish gut microbiota transplants models

Gnotobiotic response:

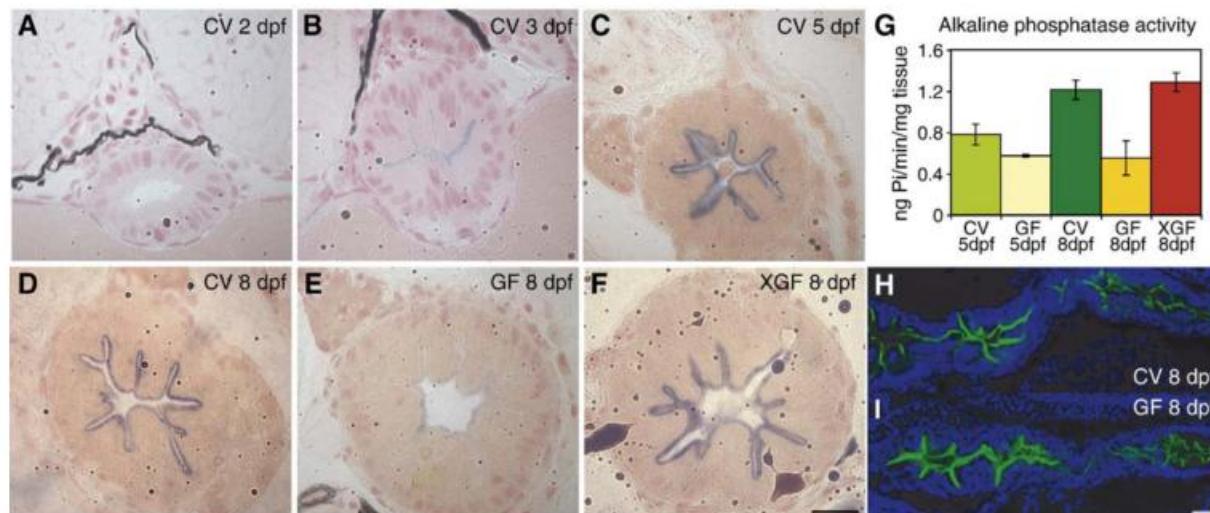
John F. Rawls, Buck

Department of Molecular

Contributed by Jeffrey I. G

Distinct sig

Jennifer M. E

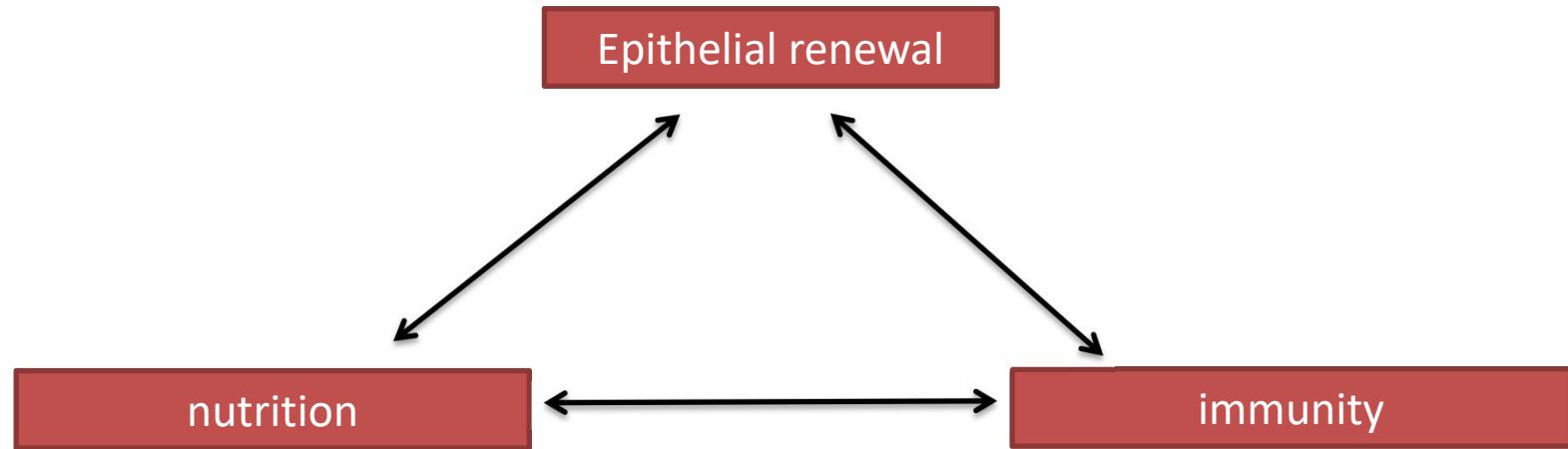


Hatch from 3 days post fertilization (dpf), microbes colonized from 3-4dpf. At 5 dpf, begins food ingestion.



Part II 鱼类肠道微生物功能研究

Role of GI microbiota in fish : gnotobiotic approaches





Part II 鱼类肠道微生物功能研究

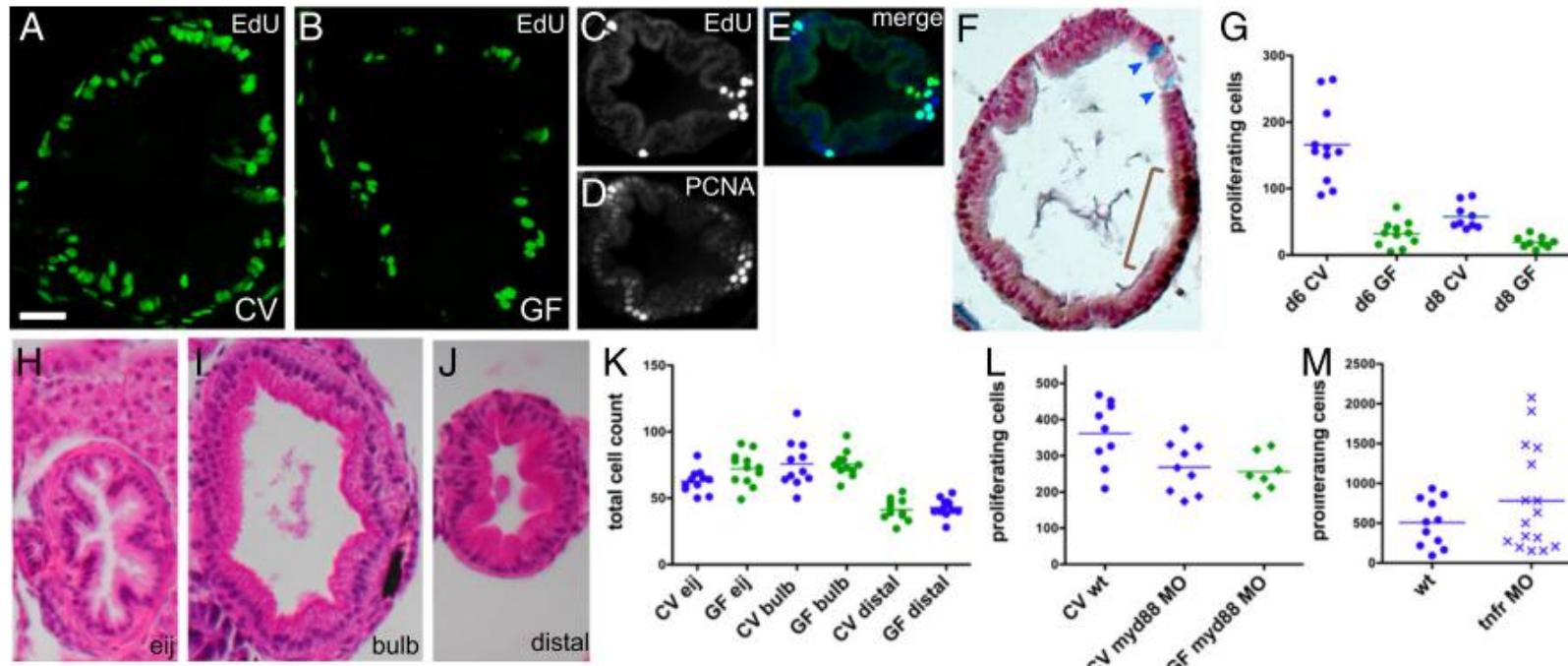


Fig. 1. Microbes induce epithelial cell proliferation in the zebrafish larval intestine via Myd88 and not inflammation. Transverse sections of the intestinal bulb of 6-dpf larvae reared CV (A) or GF (B) labeled with EdU to reveal S-phase cells are shown. A transverse section of the distal intestine shows colocalization of EdU-labeled (C) and PCNA-labeled (D) cells (E, merge). (F) Distinct cell populations are stained with Alcian blue, marking differentiated goblet cells (arrowheads), and anti-PCNA, marking mitotic cells (brown bracket). (G) Total numbers of S-phase cells over 30 serial sections in the intestinal bulb of individual 6-dpf larvae are represented for the treatment groups and genotypes indicated. Here and in subsequent figures, the genotype and microbial exposure of each larva are indicated by the shape and color of the data point, respectively. Significantly fewer BrdU-labeled cells were found in 6- and 8-dpf larvae reared GF vs. CV ($P < 0.001$). H&E sections of 8-dpf GF intestines at three locations within the intestinal tract are shown: the esophageal intestinal junction (H, eiJ), defined as position 0; the bulb, 30 sections caudal to the junction (I); and the distal intestine, 60 sections caudal to the junction (J). (K) There was no significant difference between total intestinal epithelial cell counts at the three positions described above in 8-dpf CV vs. GF larvae. (L) Significantly fewer EdU-labeled cells were found in CV or GF myd88 MO vs. CV WT (wt) ($P < 0.05$). (M) There was no significant difference in the numbers of EdU-labeled cells between CV vs. GF myd88 MO or between wt vs. tnfr MO. (Scale bars: A and B, 15 μ M; C-F and H-J, 25 μ M.)



Part II 鱼类肠道微生物功能研究

REVIEW ARTICLE

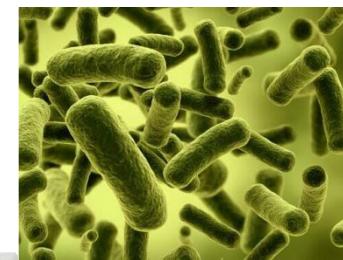
Enzyme-producing bacteria isolated from fish gut: a review

A.K. RAY¹, K. GHOSH² & E. RINGØ^{3,4}

¹ Fisheries Laboratory, Department of Zoology, Visva-Bharati University, Santiniketan, West Bengal, India; ² Aquaculture Laboratory, Department of Zoology, University of Burdwan, Burdwan, West Bengal, India; ³ Norwegian College of Fishery Science, Faculty of Biosciences, Fisheries and Economics, University of Tromsø, Tromsø, Norway; ⁴ Aquaculture Protein Centre (a CoE), Department of Aquatic Medicine and Nutrition, Norwegian School of Veterinary Medicine, Oslo, Norway

Amylase, Cellulase, Lipase, Proteases, Chitinase, Phytase

Significant roles in digestion



Protein macromolecules

Fat storage



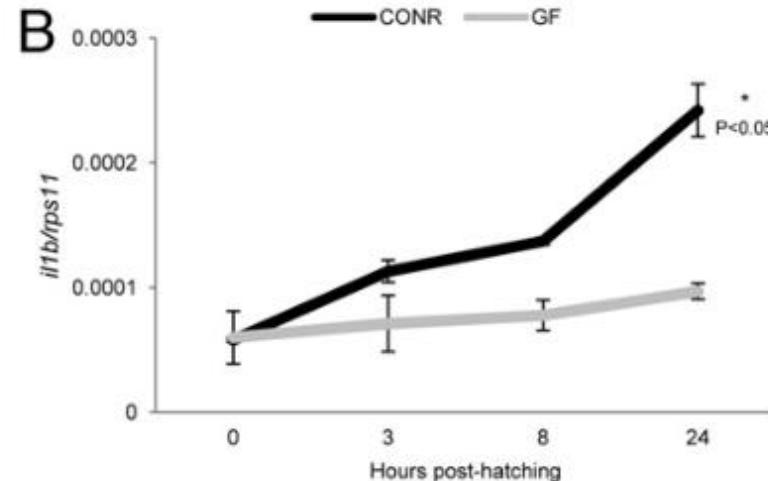
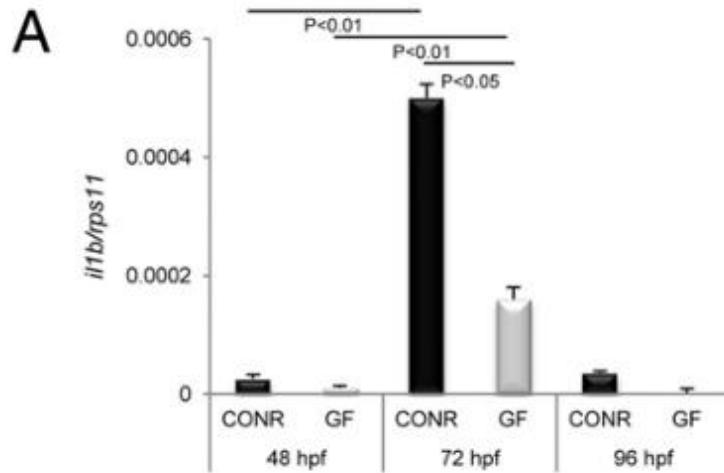
Part II 鱼类肠道微生物功能研究

PNAS

Regulation of immunity and disease resistance by commensal microbes and chromatin modifications during zebrafish development

Jorge Galindo-Villegas^{a,b,1}, Diana García-Moreno^{a,b,1}, Sofia de Oliveira^{a,c}, José Meseguer^{a,b}, and Victoriano Mulero^{a,b,2}

^aDepartamento de Biología Celular e Histología, Universidad de Murcia, 30100 Murcia, Spain; ^bInstituto Murciano de Investigación Biosanitaria, 30120 Murcia, Spain; and ^cUnidade de Biologia Microvascular e Inflamação, Instituto de Bioquímica-Instituto de Medicina Molecular, Faculdade de Medicina, Universidade de Lisboa, 1649-028 Lisbon, Portugal

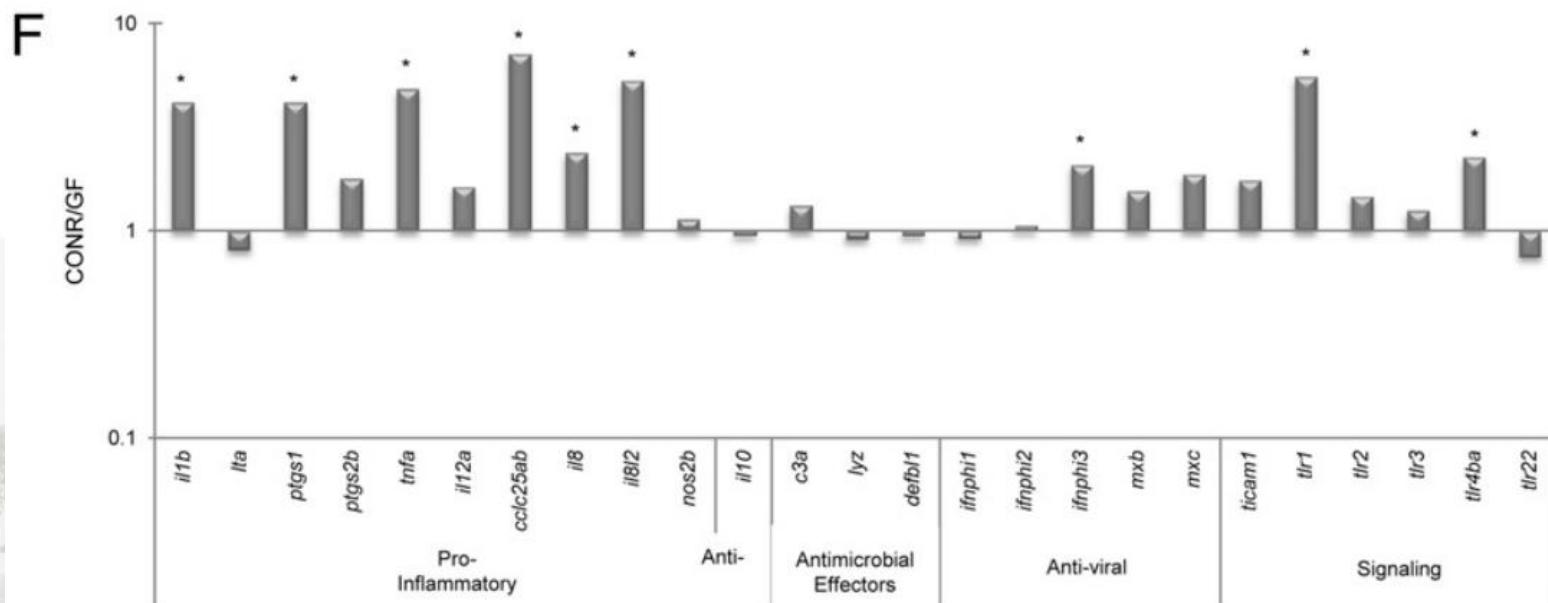
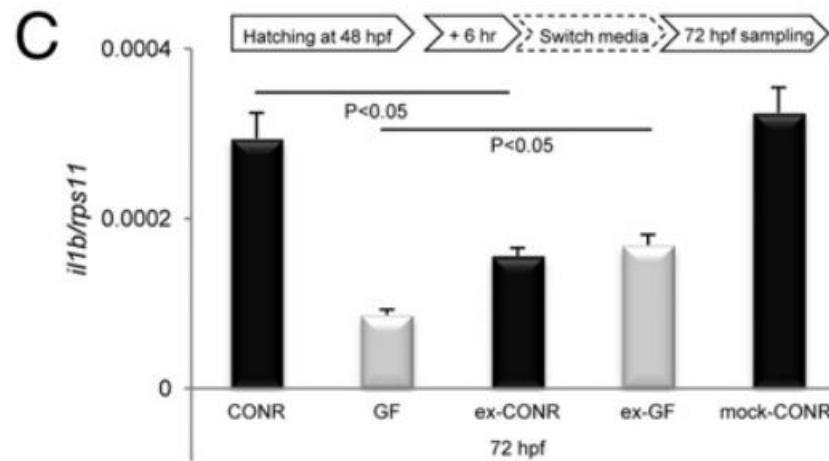


Postfertilization

Post hatching



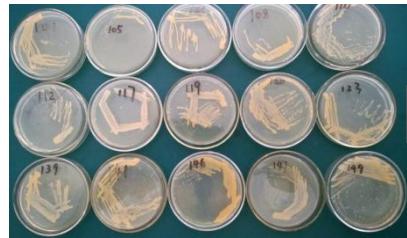
Part II 鱼类肠道微生物功能研究



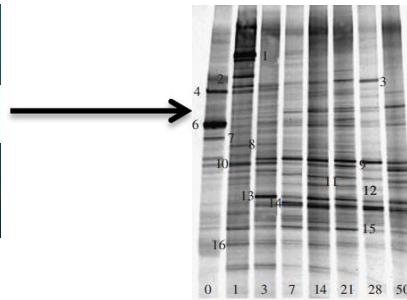


Part III 评价微生物群落的方法

Methods used to assess the bacterial communities

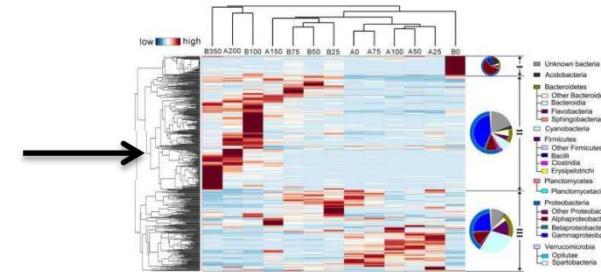
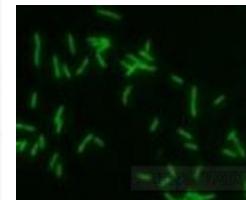


<0.1% of total species of fish



2006-2016

1. Clone library
2. Finger printing
3. qPCR or FISH
4. FISH and immunohistochemistry



NGS



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