Microbial Colonization Promotes Model Invertebrate Gut Tissue Growth and Development

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Research  

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Abstract

BACKGROUND. Digestive tissues are essential for diet processing and nutrient accessibility, especially in omnivores, and these functions occur despite and in collaboration with dynamic microbial communities that reside within and upon these tissues. Prolonged host development and reduced digestive tissue sizes have been observed in germ-free animals and normal host phenotypes were recovered following the re-introduction of typical gut microbiomes.

RESULTS. High-resolution histological analyses of Periplaneta americana cockroach digestive tissues revealed that total elimination of gut bacteria had severe impacts on the growth and development of gut tissues, especially the posterior midgut and anterior hindgut subcompartments that are expected to be colonized and inhabited by the greatest number of bacteria. Juveniles that were briefly exposed to normal gut microflora exhibited a partial gut morphological recovery, suggesting that a single inoculation was insufficient. These data highlight gut microbiota as integral to normal growth and development of tissues they are in direct contact with and, more broadly, the organism in which they reside.

CONCLUSIONS. We draw on these data, integrate host life history traits (i.e. multigenerational cohousing, molting, and filial coprophagy and exuvia feeding), and previous studies to propose a host developmental model in which gut tissues reflect a conflict-collaboration dynamic where 1) nutrient-absorptive anterior midgut tissues are in competition with transient and resident bacteria for easily assimilable dietary nutrients and whose growth is least-affected by the presence of gut bacteria and 2) posterior midgut, anterior hindgut, and to a lesser degree, posterior hindgut tissues are significantly impacted by gut bacterial presence because they are occupied by the greatest number of bacteria and the host is relying upon, and thus collaborating with, them to assist with complex polysaccharide catabolism processing and nutrient provisioning (i.e. short-chain fatty acids).

Full-text

Due to technical limitations, full-text HTML conversion of this manuscript could not be completed. However, the manuscript can be downloaded and accessed as a PDF.

Figures
Figure 1

P. americana digestive tract and transverse section measurement schematic. Dissection of complete digestive tract (A) from 5th instar P. americana with the midgut (dashed box) and subcompartments and hindgut (solid box) subcompartments labeled. Exemplars of stained and imaged (100X magnification) transverse sections from lab reared P. americana anterior midgut (B), posterior midgut (C), anterior hindgut (D) and posterior hindgut (E). Diagram depicts the visceral muscle (dark gray), epithelial tissue
(light gray) and lumen (white) in a stylized simplification of a transverse section (100x total magnification) of P. americana gut tissues (F). Masks (black) were generated from processed and imaged thin-sections and used to collect visceral muscle thickness (VMT; F), epithelial cross-sectional area (CSA; G), luminal perimeter (LPR; H), and gut perimeter (GPR; I) measurements.

**Figure 2**

Exposure to normal gut bacterial community impacts gut subcompartment growth and development across several gut physiological measures. ANOVA was conducted on the average value of epithelial cross-sectional area (CSA), gut perimeter (GPR), luminal perimeter (LPR), gut perimeter-to-luminal perimeter ratio (a measure of luminal crypt depth) measurements and visceral muscle thickness (VMT) measurements from each insect. Boxplots represent first and third 500 quartiles, median, minimum and maximum values. Germ-free insects (GF, open fill), Single exposure to conspecific feces (Con, light grey fill); ad libatum access to conspecific feces and cohousing with nonsanitized insects under nonsterile conditions (WT, dark grey fill).

**Supplementary Files**

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• SupplementalMaterialsDataTables.xlsx