



Joint Seminar

Developmental and Stem Cell Biology Program
Cell Biology Program

Division of Gastroenterology, Hepatology & Nutrition

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A tale of two tissues: Lgr5⁺ stem cells in the stomach and uterus

We have identified Lgr5 as a facultative component of the Wnt receptor complex selectively expressed on various homeostatic stem cell populations. Employing new, non-variegated Lgr5-2A-CreERT2/EGFP/DTR mouse models we now identify a subset of Lgr5-expressing chief cells responsible for epithelial repair and cancer initiation in the corpus stomach following parietal cell atrophy. We additionally characterize the transcriptomes Lgr5⁺ stem cells in mouse intestine, colon and stomach, revealing new gastric stem cell-specific markers that can be used to isolate human gastric stem cells for regenerative medicine applications and for use in selectively targeting cancer-causing mutations to the Lgr5⁺ stem cell compartment in mice as a means of evaluating their contribution to gastric cancer initiation and progression. Finally, we identify neonatal Lgr5⁺ cells in the mouse uterus as Wnt-dependent stem cells responsible for epithelial gland development. Adjacent Lgr5⁻ epithelial cells within the neonatal glands function as essential niche components to support the function of Lgr5⁺ stem cells *ex-vivo*.

Wednesday | April 3 | 2019 | 12 pm

Daniels Hollywood Theatre
(1246 Black Wing)

Hosts: Drs. Tae-Hee Kim & Robert Bandsma