JOSEPH F. FOSTER MEMORIAL CHEMICAL BIOLOGY AND BIOCHEMISTRY SEMINAR

Monday, April 15, 2024 3:30 PM, BRWN 4102

"Context Matters: Syndecan-2 Regulates Hematopoietic Stem Cell Self-Renewal"



CHRISTINA TERMINI Assistant Professor Translational Science & Therapeutics Division Fred Hutchinson Cancer Center Affiliate Assistant Professor Dept. of Laboratory Medicine & Pathology University of Washington School of Medicine

Abstract:

Glycans are the most abundant and least studied macromolecule. Proteoglycans are a specific class of glycan-bearing molecules that orchestrate cellular signaling by binding growth factors to control their receptor interactions and activation. We previously discovered a unique population of adult blood stem cells (hematopoietic stem cells), that express the heparan sulfate proteoglycan, Syndecan-2, at high levels. Increased Syndecan-2 expression renders hematopoietic stem cells with increased self-renewal capacity, which is ablated upon genetic knockdown. However, the bone marrow microenvironment, or niche, is also a rich source of proteoglycans, especially Syndecan-2, suggesting interplay between hematopoietic cell-intrinsic and microenvironment-derived proteoglycans exists. In this talk, I will summarize our recent findings regarding how the cellular source of Syndecan-2 influences hematopoietic stem cell self-renewal.



Department of Chemistry