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Education

May 2010 Ph.D. Pharmacology/Pharmaceutical Sciences Northeastern University, Boston, MA
April 2006 M.S. Pharmaceutical Sciences Northeastern University, Boston, MA

Positions and Employment

2003 Intern, Quality Control, MARS Therapeutics Ltd., Hyderabad, India
2005-2006 Research Fellow, Institute on Urban Health Research, Northeastern University, Boston, MA
2006 Co-Op, Preclinical Pharmacology, Genzyme, Framingham, MA
2010-2014 Postdoctoral Associate, Dr. Leonard Shultz's lab, The Jackson Laboratory, Bar Harbor, ME
2015-2018 Associate Research Scientist, The Jackson Laboratory, Bar Harbor, ME
2019-present Research Scientist, The Jackson Laboratory, Bar Harbor, ME

Funding

2022 NCI R01
2019 NIH R21
2017 Director's Innovation Fund (DIF), JAX, Bar Harbor, ME
2011 Juvenile Diabetes Research Foundation (JDRF) International Postdoctoral Fellowship
2010 JAX Postdoctoral Scholarship, Bar Harbor, ME

Honors

2005 Fellowship from Northeastern University for outstanding performance in academia (summer and fall)
2005 Scholarship for the 20th Annual New England Biolabs, Inc. "Molecular Biology" summer course, Smith College, Northampton, MA
2006 Best Poster, 35th Annual New England Pharmacologists meeting, Boston, MA
2009 Best Poster Runner-up Award, Research and Scholarship Expo, Northeastern University, Boston, MA
2010 The John and Evelyn Neumeier Research Achievement Award, Northeastern University, Boston, MA

Publications

1. Programmable RNA-Guided Large DNA Transgenesis by CRISPR/Cas9 and Site-Specific Integrase Bxb1. *Front Bioengineering Biotechnology* 2022.
2. Inactive rhomboid proteins RHBDF1 and RHBDF2 (iRhoms): A decade of research in murine models *Burzenski L, Low BE, Kohar V, Shultz LD, Wiles MV, Hosur V. Mammalian Genome* 2021
3. Role of MicroRNA in Inflammatory Bowel Disease: Clinical Evidence and the Development of Preclinical Animal Models. *Suri K, Bubier J, Wiles MV, Shultz LD, Amiji MM, Hosur V. Cells* 2021
4. Genes adapt to outsmart gene-targeting strategies in mutant mouse strains by skipping exons to reinitiate transcription and translation. *Hosur V, Low BE, Li D, Stafford GA, Kohar V, Shultz LD, Wiles MV. Genome Biology* 2020.
5. Improved mouse models and advanced genetic and genomic technologies for the study of neutrophils. *Hosur V, Skelly DA, Francis C, Low BE, Kohar V, Burzenski LM, Amiji MM, Shultz LD, Wiles MV. Drug Discovery Today*, 2020.
6. RHBDF2-regulated growth factor signaling in a rare human disease, tylosis with esophageal cancer: what can we learn from murine models? *Hosur V, Farley ML, Low BE, Burzenski LM, Shultz LD, Wiles MV. Frontiers in Genetics*. 2018
7. ADAM17 is essential for ectodomain shedding of the EGF-receptor ligand amphiregulin. *Hosur V, Farley ML, Burzenski LM, Shultz LD, Wiles MV. FEBS Open Bio*. 2017

8. Early induction of NRF2 antioxidant pathway by RHBDF2 mediates rapid cutaneous wound healing. **Hosur V**, Burzenski LM, Stearns TM, Farley ML, Sundberg JP, Wiles MV, Shultz LD. *Exp Mol Pathology*. 2017
9. Genetic deletion of amphiregulin restores the normal skin phenotype in a mouse model of the human skin disease tylosis. **Hosur V**, Low BE, Shultz LD, Wiles MV. *Biol Open*. 2017
10. Tissue-specific role of RHBDF2 in cutaneous wound healing and hyperproliferative skin disease. **Hosur V**, Lyons BL, Burzenski LM, Shultz LD. *BMC Res Notes*. 2017
11. Genetic modification of mice using CRISPR-Cas9: Best practices and practical concepts explained. **Hosur V**, Low BE, Wiles MV. *Rigor and Reproducibility in Genetics and Genomics 2021*, In press
12. Efficient targeted transgenesis of large donor DNAs in multiple mouse genetic backgrounds using bacteriophage Bxb1 integrase. Low BE, **Hosur V**, Lesbirel S, Wiles MV. *Scientific reports*, In revision. 2022
13. Development of Humanized Mice in the Age of Genome Editing. **Hosur V**, Low BE, Avery C, Shultz LD, Wiles MV. *J Cell Biochemistry*. 2017
14. Human cancer growth and therapy in immunodeficient mouse models. Shultz LD, Goodwin N, Ishikawa F, **Hosur V**, Lyons BL, Greiner DL. *Cold Spring Harbor Protocols*. 2014
15. Subcapsular transplantation of tissue in the kidney. Shultz LD, Goodwin N, Ishikawa F, **Hosur V**, Lyons BL, Greiner DL. *Cold Spring Harbor Protocols*. 2014
16. Rhbdf2 mutations increase its protein stability and drive EGFR hyperactivation through enhanced secretion of amphiregulin. **Hosur V**, Johnson KR, Burzenski LM, Stearns TM, Maser RS, Shultz LD. *Proc Natl Academy Sci U S A*. 2014
17. Retrotransposon insertion in the T-cell acute lymphocytic leukemia 1 (Tal1) gene is associated with severe renal disease and patchy alopecia in Hairpatches (Hpt) mice. **Hosur V**, Cox ML, Burzenski LM, Riding RL, Alley L, Lyons BL, Kavirayani A, Martin KA, Cox GA, Johnson KR, Shultz LD. *PLoS One*. 2013
18. Dystrophin and dysferlin double mutant mice: a novel model for rhabdomyosarcoma. **Hosur V**, Kavirayani A, Riefler J, Carney LM, Lyons B, Gott B, Cox GA, Shultz LD. *Cancer Genet*. 2012
19. Engraftment of human HSCs in nonirradiated newborn NOD-scid IL2 γ null mice is enhanced by transgenic expression of membrane-bound human SCF. Brehm MA, Racki WJ, Leif J, Burzenski L, **Hosur V**, Wetmore A, Gott B, Herlihy M, Ignatz R, Dunn R, Shultz LD, Greiner DL. *Blood*. 2012
20. Humanized mice for the study of type 1 and type 2 diabetes. Greiner DL, Brehm MA, **Hosur V**, Harlan DM, Powers AC, Shultz LD. *Ann NY Acad Sci*. 2012
21. $\alpha 4\beta 2$ nicotinic receptors partially mediate anti-inflammatory effects through Janus kinase 2-signal transducer and activator of transcription 3 but not calcium or cAMP signaling. **Hosur V**, Loring RH. *Mol Pharmacol*. 2011
22. Gene regulation of alpha4beta2 nicotinic receptors: microarray analysis of nicotine-induced receptor up-regulation and anti-inflammatory effects. **Hosur V**, Leppanen S, Abutaha A, Loring RH. *J Neurochem*. 2009

Research Support

NCI R01

07/01/22-06/30/27

Site-specific Integration of Large (10-100 kb) DNA constructs into the mouse genome and human induced pluripotent stem cells using the Cas9-Bxb1 integrase toolbox

Advances in experimental techniques to modify the genetic makeup of mice have enabled development of mice that “model” particular diseases, including cancers, providing live organisms that researchers can use to study disease progression and test therapies. However, a major roadblock in the development of such models is the inefficiency of inserting large tracts of DNA into mouse chromosomes, a limitation that hinders research on the role of many coding and non-coding DNA regions in cancer. In the proposed project we will develop, validate, and make publicly available an innovative technique for precise insertion of large tracts of DNA into mouse chromosomes, enabling researchers to develop improved and new models of cancer.

Role: Principal Investigator

NIH R21

04/01/19-03/31/22

Development and Validation of a Novel Cas13a and Nanoparticle Guide-RNA Delivery System that Allows Precise Ablation of Host Macrophage Populations in a Humanized Mouse Model

Role: Co-I

The goal is to develop a state-of-the-art technology to precisely modulate mouse macrophages *in vivo* in mice, thus enabling heightened engraftment of the human hematopoietic/immune system, including human RBCs.

Completed

JAX-DIF-FY17

06/01/17-05/31/19

The Jackson Laboratory Director's Innovation Fund

NSG-Bald, an improved PDX Model for Rapid Growth, Study and Treatment of Human Tumors

The goal of this project is to test the hypothesis that growth of certain tumor types is constrained by the lack of host oncogenic factors, including amphiregulin (AREG) and heparin-binding EGF (HB-EGF), both linked with angiogenesis.

Role: Principal Investigator

3-2011-391

09/01/11-08/31/14

Juvenile Diabetes Research Foundation International

Human Immune System-Engrafted mice for the study of Islet Rejection and type 1 diabetes.

The goal was to study human islet allograft rejection and the development of type 1 diabetes in novel strains of immunodeficient mice that have functional human immune systems generated by engraftment of human hematopoietic stem cells.

Role: Principal Investigator