Chang K. Sung, PhD

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MOLECULAR CANCER CELL BIOLOGIST

Significant experience in protein destruction- and small RNA-mediated pathological signaling pathways Taught various biology courses and improved educational materials

Guided and supervised junior researchers and students to develop research projects

Discovered protein destruction mechanisms of a tumor suppressor that could lead to development of novel therapeutic approaches in human ovarian carcinomas

Performed Q RT PCR-based RNA arrays and identified microRNAs that could be used as biomarkers for early detection of human cancers and tumor progression

TEACHING EXPERIENCE

Texas A&M University-Kingsville Kingsville, TX Assistant Professor Genetics Lecture and Recitation Fall 2013 University of Illinois at Chicago Chicago, IL **Teaching Assistant** -Taught laboratory and discussion sections for Lab and Lecture courses for seven academic semesters

-Developed educational materials and curricula aimed at encouraging student participation such as research article group discussions and research data/background presentations

Cell Biology Lab	Fall 2003
Genetics Lab	Fall 2002, Fall 2000, Spring 2000
Genetics Lecture	Fall 2001, Spring 2001
General Biology Lab	Fall 1999

RESEARCH EXPERIENCE

Texas A&M University-Kingsville

ASSISTANT PROFESSOR in Biology

Research interests include tumorigenesis and programmed cell death mechanisms utilizing the mouse polyoma virus experimental system. Current studies also focus on functional roles of the p150 protein, a tumor suppressor in normal human ovarian surface epithelial cells

Harvard Medical School

INSTRUCTOR in Microbiology and Immunobiology

Investigated tumorigenesis and programmed cell death mechanisms with various cell based assays including protein-protein interaction studies, Q RT PCR arrays and pyrosequencing analyses.

-Screened de-ubiquitinating enzymes (DUBs) that may preserve a tumor suppressor p150 in normal human ovarian surface epithelial cells

2011-2013 Boston, MA

2013-Present

Kingsville, TX

1

- -Examined epigenetic silencing (promoter methylation) of the tumor suppressor p150 in human ovarian carcinomas
- -Screened murine microRNAs in response to the oncogenic polyoma virus infection
- to identify novel biomolecular markers involved in oncogenesis and tumor progression
- -Investigated how polymorphisms in toll-like receptor 4 (Tlr4) lead

Sung, C.K.*, Yim, H.*, Gu, H., Li, D., Andrews, E., Duraisamy, S., Li, C., Drapkin, R. and Benjamin, T.L. (2012) The Polyoma Virus Large T Binding Protein p150 is a Transcriptional Repressor of c-MYC. **PLoS ONE** 7(9): e46486 *<u>contributed equally</u>

Andrews, E.*, Velupillai, P.*, **Sung, C.K.**, Beier, D. and Benjamin, T.L. (2012) Production of a Natural Antibody to the Mouse Polyoma Virus is a Multigenic Trait. **G3: Genes, Genomes, Genetics** 2(3):353-5 *<u>contributed equally</u>

Sung, C.K.*, Dahl, J.*, Yim, H., Rodig, S. and Benjamin, T.L. (2011) Transcriptional and Post-translational Regulation of the Quiescence Factor and Putative Tumor Suppressor p150^{Sal2}. **FASEB J** 25(4):1275-83 *<u>contributed equally</u>

Yim, H., Sung, C.K.