KIRA YOUNG

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INTERDISCPLINARY SCIENTIST/ENGINEER

Scientist/chemical engineer with multidisciplinary research and development background. Expertise in molecular biology and analytical chemistry techniques, data collection and analysis. Able to work both independently and as part of a team. Willing to master new techniques as needed.

EDUCATION

PhD, Functional Genomics, University of Maine, Orono, ME Graduation Date: May 2014 *Dissertation:* A proteomics approach to the study of endoglin function and hereditary hemorrhagic telangiectasia

BS, Chemical Engineering, summa cum laude, University of Maine, Orono, ME, 2003 **GPA: 3.74/4.00**

George Mitchell Peace Scholar, University College Cork, Cork, Ireland, Sept-Dec 2001

RESEARCH EXPERIENCE

Jackson Laboratory, Bar Harbor, ME

Postdoctoral Trainee

Research Highlights:

• Conducting research to identify the underlying mechanisms driving MDS-to-sAML progression and order of mutational acquisition in the development of AML

• Techniques include reduced representation bisulfite sequencing to assay genome-wide DNA methylation, mapping of double strand breaks using the BLESS method and open chromatin region identification using transposase-accesible chromatin sequencing.

Maine Medical Center Research Institute, Scarborough, ME Graduate School of Biomedical Sciences

Ph.D. Candidate, Functional Genomics

A full time research graduate student seeking a Ph.D degree in functional genomics focused on understanding molecular signaling during angiogenesis using *in vitro* systems and *in vivo* mouse models.

Research Highlights:

• Conceptualized and designed proteomics/mass spectrometry-based projects aimed at identifying novel gene targets of TGF- β signaling and their effect on angiogenesis and determined specific functions for endoglin involved in the pathology of arterialvenous malformations in the vascular dysplasia, hereditary hemorrhagic telangiectasia (HHT).

• Required methods included analysis of *in vitro* cell culture of primary human cells, viral transduction methods (retro-, lenti- and adenovirus), cell imaging using confocal microscopy, immunohistochemistry of mouse embryos and perfused adult mouse lungs, maintenance and breeding of multiple mouse strains, PCR, western blot and mass spectrometry to determine novel gene targets.

• Mentored undergraduate and graduate level individuals seeking research experience. Responsible for the oversight and success of their projects.

2014-present

2006 - 2014

• Designed experiments involving the interaction of casein kinase-2 (CK2) with bone morphogenetic proteins (BMPs) in immortalized cells using FRET analysis and confocal microscopy.

Assisted in laboratory start-up for Dr. Anja Nohe including ordering and maintenance of supplies and compliance with laboratory safety regulations.

Molecular biology techniques: confocal microscopy, western blots, cloning techniques, FRET analysis, reporter gene assays.

University of Maine, Orono, ME

Department of Chemical and Biological Engineering and Department of Chemistry

Teaching Assistant

Research Assistant

- Assisted in teaching and preparation of molecular biology methods laboratory course (2006).
- Responsible for teaching and providing assistance to students in a general chemistry lab of eighteen students (2001).

University of Western, London, Ontario

Research Assistant

Assisted in *in vitro* research involving the interaction of epidermal growth factor (EGF) with BMPs. •

Molecular biology techniques: quantum dots with live cell imaging, confocal microscopy and image correlation spectroscopy, western blots, immunoprecipitation.

SCHOLARSHIPS/AWARDS

2012	1 st Place, Graduate Student Poster Award, NAVBO, Genetics and Genomics of Vascular Disease
2012	Student travel award, NAVBO, Genetics and Genomics of Vascular Disease
2006-2009	NSF-IGERT Functional Genomics Trainee
2001	George Mitchell Peace Scholarship
1998-2003	Pulp and Paper Scholarship
1998-2003	Top Scholar Award

ABSTRACTS

Young K., Tweedie E., Conley B., Vary C.P. Quantitative proteomics identifies correlation of *in vitro* and *in vivo* analysis of protein identification in response to endoglin expression.

Young K., Krebs L., Tweedie E., Conley B., Gridley T., Vary C.P. Conditional inactivation of endoglin in Pax3 expressing cells leads to intersomitic vessel defects.

PUBLICATIONS

Young, K., Conley B., Tweedie E., Brooks P.C., Vary C.P. (2012) Endoglin signaling regulates angiogenesis via the Hippo pathway. (Plos One, manuscript under revision for publication)

Young, K., Conley, B., Romero D., Tweedie E., O'Neill C., Pinz I., Brogan L., Lindner V., Liaw L., Vary C.P. (2012) BMP9 regulates endoglin-dependent chemokine responses in endothelial cells. Blood. 120(20):4263-73.

2001 and 2006

May - Sept 2003

Romero D., O'Neill C., Terzic A., Contois L., <u>Young K.</u>, Conley BA., Bergan RC., Brooks P.C., Vary C.P. (2011) Endoglin regulates cancer-stromal cell interactions in prostate tumors. *Cancer Res.* 71(10):3482-93.

Bragdon B., Thinakaran S., Moseychuk O., King D., <u>Young K.</u>, Litchfield DW., Petersen N.O., Nohe A. (2010) Casein kinase 2 beta-subunit is a regulator of bone morphogenetic protein 2 signaling. *Biophys J.* 99(3):897-904.

REFERENCES

Calvin Vary, PhD Center for Molecular Medicine MMCRI 81 Research Drive Scarborough ME, 04074 <u>varyc@mmc.org</u> (207) 396-8148 Robert Friesel, PhD Center for Molecular Medicine MMCRI 81 Research Drive Scarborough ME, 04074 <u>friesr@mmc.org</u> (207) 396-8147

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