## KELIE M'LISS REECE, Ph.D.

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#### **CAREER OBJECTIVE**

To obtain a scientific editing position that will allow me to use my knowledge, research, and editing experience to review grants and manuscripts.

#### **QUALIFICATIONS**

- Extensive experience writing, editing, and reviewing scientific documents including grants and manuscripts
- Extensive experience working as a freelance editor and editing documents from non-native English speakers
- · Very organized and detail-oriented
- Excellent written communication skills
- Expertise with Microsoft Office Suite
- Expertise in Molecular Pharmacology, Cancer Biology, Proteomics, Cell and Molecular Biology

#### **EDUCATION**

Postdoctoral Fellow, Molecular Pharmacology Section, Medical Oncology Department,
 National Cancer Institute, Bethesda, MD

Ph.D., Pharmacology, Vanderbilt University, Nashville, TN 2009

B.A., Biology, Bryn Mawr College, Bryn Mawr, PA

## **WORK EXPERIENCE**

### Regulatory Health Project Manager: US Food and Drug Administration, Silver Spring, MD

2017-present

Manages the drug review process from initial submission to the time of approval, termination, or withdrawal; implements regulations regarding new drugs and drug products and biologics approved by the organization in the application process; manages and coordinates the work of review team members (scientific, regulatory, and management) for specific drug application reviews; develops project plans including setting time frames, milestones, and endpoints.

#### Scientific Review Manager: SRA International, Inc., Frederick, MD

2013-2016

Managed all aspects of the scientific peer review of grant applications for the Product Development and Prevention Programs for the Cancer Prevention & Research Institute of Texas (CPRIT); developed an annual project plan for the scientific peer review of applications submitted for consideration; served as the primary client liaison for all technical aspects regarding the review process; configured peer review panels; managed the peer review meetings and oversaw the scientific integrity of the review process; ensured that program objectives were met on time.

### **RESEARCH EXPERIENCE**

# Molecular Pharmacology Section, Medical Oncology Branch, National Cancer Institute

Postdoctoral Fellow 2010-2013

Mentor: William D. Figg, Pharm.D., M.B.A.

- Project 1: Investigated the role of ERCC1 polymorphisms in sensitivity to treatment with platinum agents.
- **Project 2:** Investigated the effect of hypoxia-inducible factor  $1\alpha$  (HIF- $1\alpha$ )/p300 complex disruption in prostate cancer.
- **Project 3:** Investigated the cooperative regulation of HIF- $1\alpha$  and androgen receptor-mediated signaling by dual treatment with MDV3100 and HIF- $1\alpha$  inhibitors.
- **Project 4:** Investigated the effect of hypoxia on the expression of organic anion transporting polypeptide 1B3 in cancer.
- Responsibilities: Designed, implemented, managed, and executed multiple research projects; advised, supervised, and
  directed multiple undergraduate, graduate, and postbaccalaureate students on various projects; wrote, reviewed, and
  edited grant proposals and manuscript submissions.

### Department of Pharmacology, Vanderbilt University

Thesis Mentor: Brian Wadzinski, Ph.D. 2001-2009

**Doctoral Research** 

Project: "Regulation of the CaMKIV/PP2A Signaling Module." Investigated the molecular mechanisms governing the
inactivation of CaMKIV by PP2A. This work laid the foundation for future studies directed towards understanding how the
manipulation or intervention of this pathway could impact key biological processes regulated by this signaling complex,
such as neuronal growth and function, and memory formation and maintenance. My research and expert writing skills led
to the award of an NIH training grant.

Responsibilities: Designed and executed experiments; organized graduate thesis research, including collaborations and
mentoring committees; supervised undergraduate and high school students; contributed and assisted with the
development and writing of grants and manuscripts.

#### **TEACHING EXPERIENCE**

Guest Lecturer, Molecular Targets and Cancer, Johns Hopkins University

Topic: Overview of Protein Kinases and Protein Phosphatases

Topic: Angiogenesis Targets

Teacher, AIDS Education Outreach Program

2002-2009

High school tutor, all subjects

2002-2008

#### **EDITORIAL APPOINTMENTS**

Scientific Editor: 2011-2017

Responsibilities include grant and manuscript copyediting, critiquing, polishing, and proofreading

**Enago English Proofreading** 

Paper Talks

Compuscript Ltd.

Editorial Board: 2011-2013

Frontiers in Cancer Molecular Targets and Therapeutics

**Reviewer:** 2010-2013

Cancer Biology and Therapy
Oncogene
Diochimie

Clinical Cancer Research
Journal of Molecular Medicine
American Journal of Pathology

Molecular Cancer Therapeutics BMC Cancer

PNAS Future Medicinal Chemistry

# **SELECTED PUBLICATIONS:**

\*Fernandez E.V., \*Reece K.M., Ley A.M., Troutman S.M., Sissung T.M., Price D.K., Chau C.H., Figg W.D. (2015) Dual targeting of the androgen receptor and hypoxia-inducible factor 1a pathways synergistically inhibits castration-resistant prostate cancer cells. Mol. Pharmacol. 87(6):1006-12

Reece, K.M., Richardson, E.D., Cook, K.M., Campbell, T.J., Pisle, S.T., Holly, A.J., Venzon, D.J., Liewehr, D.J., Chau, C.H., Price, D.K., Figg, W.D. (2014) Epidithiodiketopiperazines (ETPs) exhibit antiangiogenic and antitumor activity by disrupting the HIF-1a/p300 complex in a preclinical model of prostate cancer. Mol. Cancer. 13(91)

Reece, K.M., Figg, W.D. (2013) UGT1A1 polymorphisms and mutations in irinotecan-induced toxicity. In S. Yang & J. Dancey (Ed.) Handbook of Therapeutic Biomarkers in Cancer (pp 353-368). Chicago, IL: Pan Stanford Publishing

Gao, R., Reece, K.M., Sissung, T.M., Fu, S.H., Venzon, D., Price, D.K., Reed, E., Spencer, S.D., Price, D.K., Figg, W.D. (2012) Are race-specific ERCC1 haplotypes in melanoma cases versus controls related to the predictive and prognostic value of ERCC1 N1118N? BMJ Open. 3(1)

<sup>\*</sup>both authors contributed equally to this work