

**CURRICULUM VITAE****PERSONAL BACKGROUND**

Name: **LAWRENCE T. REITER**  
 Birthplace: Morristown, New Jersey  
 Date of Birth: April 29<sup>th</sup>, 1969  
 Citizenship: USA  
 Marital Status: Married: wife (Tiffany); son (Vincent)

Address: Department of Neurology  
 University of Tennessee Health Science Center  
 855 Monroe Avenue, Link 415  
 Memphis, TN 38163

Phone/FAX Office: (901) 448-2635  
 Lab: (901) 448-7443  
 FAX: (901) 448-7440  
 Cell: (619) 520-2529

E-mail: [lreiter@uthsc.edu](mailto:lreiter@uthsc.edu)  
 ORCID ID: 0000-0002-4100-2630  
 ERA Commons ID: Ireiter

**EDUCATION****Undergraduate:**

University of Southern California (USC), 1987-1991  
 Major: Molecular Biology; Minor: Chemistry  
 Degree: B.S. (1991)

**Graduate:**

Baylor College of Medicine, 1993 - 1997  
 Major: Cell and Molecular Biology  
 Degree: Ph.D. (1997); Mentor: James R. Lupski, M.D., Ph.D.

**Postdoctoral:**

Department of Biology, University of California, San Diego (UCSD), 1999 - 2005  
 Field: *Drosophila* Genetics; Mentor: Ethan Bier, Ph.D.

**HONORS AND AWARD**

Claude W. Smith Fellowship Award	1996
Boehringer Ingelheim Fellowship Award	1996
Young Scientist Fellowship Award	1996
Postdoctoral Fellowship CMT Association	1998
Sigma Xi Outstanding Dissertation Award	1998
Finalist A.S.H.G. Postdoctoral Presentation Award	1998
Glaucoma Foundation Fellowship	2000-2001
Full Scholarship to CSHL Bioinformatics Course	2001
Winner A.S.H.G. Postdoctoral Presentation Award	2003

Recipient, NIH Loan Repayment Program	2005-2009
Distinguished Mentor Award, Nominee	2011
Vice Chancellor for Research, CORNET Awardee	2017

**PROFESSIONAL EXPERIENCE**

1987-1991	Undergraduate Research, Dr. Carol Miller, USC, Los Angeles, CA
1991-1992	Eighth Grade Earth Science Teacher, Mc Main School, Orleans Parrish, New Orleans, LA
1992-1993	Research Associate, Dr. Max Oeschger, LSU. Dental School, New Orleans, LA
1993-1997	Graduate Student, Dr. James R. Lupski, Baylor College of Medicine, Houston, TX
1997-1998	Postdoctoral Fellow, Dr. James R. Lupski, Baylor College of Medicine, Houston, TX
1999-2005	Postdoctoral Fellow, Dr. Ethan Bier, UCSD, San Diego, CA
2005-2011	Assistant Professor, Dept. of Neurology, UTHSC, Memphis, TN
2011-2017	Associate Professor (w/ tenure), Dept. of Neurology, UTHSC, Memphis, TN
2017-present	Full Professor, Dept. of Neurology, UTHSC, Memphis, TN
2005-present	Adjunct, Anatomy and Neurobiology, UTHSC, Memphis, TN
2006-present	Adjunct, Dept. of Pediatrics, UTHSC, Memphis, TN
2006-2016	Director, <i>Drosophila</i> Transgenic Core, UTHSC, Memphis, TN
2014-2017	Director, 15q Duplication Autism Clinic, Le Bonheur Children's Hosp.

**TEACHING EXPERIENCE**

1991-1992	8 <sup>th</sup> grade Earth Science, Teach for America Program, New Orleans, LA
2000	Course Instructor, Molecular Biology I, UCSD, La Jolla, CA
2003	Guest Lecturer, Biology of Model Organisms, UCSD, La Jolla, CA
2007	Guest Lecturer, Developmental Neuroscience, UTHSC, Memphis, TN
2008	Guest Lecturer, Systems Biology, UTHSC, Memphis, TN
2008	Guest Lecturer, Bioinformatics, UTHSC, Memphis, TN
2008	Guest Lecturer, Behavioral Neuroscience, UTHSC, Memphis, TN
2009	Guest Lecturer, Developmental Neuroscience, UTHSC, Memphis, TN
2009	Guest Lecturer, Bioinformatics, UTHSC, Memphis, TN
2010	Guest Lecturer, Behavioral Neuroscience, UTHSC, Memphis, TN
2012	Co-director, Science as a Profession, UTHSC, Memphis, TN
2012	Guest Lecturer, Developmental Neuroscience, UTHSC, Memphis, TN
2012	Guest Lecturer, Behavioral Neuroscience, UTHSC, Memphis, TN
2012	Guest Lecturer, Bioinformatics/Computational Bio., UTHSC, Memphis, TN
2013	Co-director, Science as a Profession, UTHSC, Memphis, TN
2013	Guest Lecturer, Bioinformatics/Computational Bio., UTHSC, Memphis, TN
2014	Guest Lecturer, Science as a Profession, UTHSC, Memphis, TN
2014	Guest Lecturer, Behavioral Neuroscience, UTHSC, Memphis, TN
2015	Guest Lecturer, Science as a Profession, UTHSC, Memphis, TN
2015	Guest Lecturer, Bioinformatics/Computational Bio., UTHSC, Memphis, TN
2016	Guest Lecturer, Science as a Profession, UTHSC, Memphis, TN
2016	Guest Lecturer, Bioinformatics/Computational Bio., UTHSC, Memphis, TN
2016	Guest Lecturer, Behavioral Neuroscience, UTHSC, Memphis, TN

**UTHSC COMMITTEES AND RELATED INSTITUTIONAL ACTIVITIES**

Molecular Resource Center Advisory Board (2005-present)  
 Clinical and Translational Science Institute Advisory Board (2007-2011)  
 CTSI, New Technologies Maven (2007-2011)  
 Postdoctoral Advisory Committee (2008-2013, Appointed President 2013)  
 Dean's Faculty Advisory Committee (2011-2016, Appointed President 2013)  
 UTHSC Strategic Planning Committee (2013-2014)  
 Molecular Bioinformatics Core Committee, Chair (2016-present)  
 Metabolomics and Proteomics Core Committee (2016-present)

**Advisory and Dissertation Committees:**

Mike DeCupreya (Mentor: LeDoux)	2005-2010
Ryan Delahanty (Mentor: Sutcliffe, Vanderbilt University)	2006-2009
Rawaha Memon, Dental Resident, Masters Student	2012-2013
Ryan Wilson, Dental Resident, Masters Student	2013-2014
Kevin Hope, Graduate Student (Mentor: Reiter)	2015-2019
A. Kaitlyn Victor, Graduate Student (Mentor: Reiter)	2017-present

**TRAINEES AND LAB MEMBERS****Present:****Postdoctoral****Pre-doctoral**

Kaitlyn Victor	Neuroscience Track IPBS Program	2017-present
----------------	---------------------------------	--------------

**Undergraduates**

Yesnia Sanchez-Arrellano	Christian Brothers	2019-present
Walt Krueger	Grad School Applicant	2020-present

**Past:****PhD. Students (1 Total)**

Kevin Hope	Neuroscience Track IPBS Program <i>Postdoc with Clement Chow, Univ Utah</i>	2015-2019
------------	--	-----------

**Technicians**

Sarita Goorha	Part-time Research Technician	2013-2017
Morgan Fuller	Clinical Research Coordinator <i>DO Student</i>	2015-2017

**Postdoctoral (7 Total)**

Nahed Elsis	Postdoctoral Fellow <i>Currently at Florida A&amp;M University</i>	2005-2006
Priti Azad	Postdoctoral Fellow <i>Associate Project Scientist, UCSD</i>	2006-2007
Samiramis Sarkardei	Postdoctoral Fellow <i>Teacher of IB Biology at PASB, Brazil</i>	2008-2009
M. Febin Farook	Postdoctoral Fellow	2009-2012

Sharath Rongali	<i>Postdoctoral Research Associate, Cleveland Clinic</i> Postdoctoral Fellow	2012-2013
Nora Urraca	<i>Assistant Professor, Missouri State University</i> Postdoctoral Fellow	2009-2014
Jungsoo Han	Postdoctoral Fellow	2017-2018
Bidisha Roy	Postdoctoral Fellow	2018-2020

**Masters Student**

Tracy Carter	Master's Program for Bench Research	2017-2018
--------------	-------------------------------------	-----------

**Medical Student Summer Research (6 Total)**

Kelly Ridder	Dystonia genes in flies <i>Former Neurology Residency UTHSC (deceased)</i>	Summer 2008
Beth Nagel	VHL/HIF1alpha in border cell migration <i>Family Practice</i>	Summer 2008
Katlin Ridder	Dystonia genes in flies <i>Former Medical Student UTHSC</i>	Summer 2008
Kayla McCrury	Dube3a/ATPalph5 behavior in flies	Summer 2009
Donald Pierce	Generation of ATPalpha5 constructs	Summer 2011
Rajiv Heda	Identification of glial-GAL4 lines in seizure <i>Third Year Medical Student UTHSC</i>	Summer 2017

**Undergraduates (32 Total)**

Mike Wangler	Masters Student <i>Current: Assistant Professor, BCM, Houston, TX</i> <a href="http://wanglerlab-bcm.org">http://wanglerlab-bcm.org</a>	1999-2001
Mark McElroy	Masters Student	2001-2002
Wendy Ching	SMART Program	Summer 2001
Megan Bowers	Undergraduate – Honors Thesis <i>Current: Resident, Georgetown University Hospital</i>	2001-2002
Alex Ho	Undergraduate	2001-2002
Kevan Akarami	Undergraduate – Honors Thesis	2003-2004
Sanaz Farid	Undergraduate	2002-2004
Courtney Sherman	Undergraduate – Honors Thesis <i>Medical School Graduate</i>	2003-2004
Neena Poole	McNair Summer Program	Summer 2005
Alan Newton	Undergraduate – Thesis <i>Pharmacy School Graduate</i>	Summer 2006
Rebecca Scott	Undergraduate – Thesis <i>Medical School Graduate</i>	2006-2008
Kyle Summers	Undergraduate/Technician <i>Ph.D., UTHSC, 2015;</i> <i>Instructor, Margolin Hebrew Academy</i>	2006-2008
William Bodeen	Undergraduate/Technician <i>PhD Program UTHSC</i>	2008-2009
Rachel Chassan	Rhodes College Student <i>Attended Graduate School, UTHSC – M.S.</i>	2010-2011

Baris Kucukkaraduman (Bilken University, Turkey)	Summer-2012
Savely Zakharenko High School Student	Summer-2012
Charlese Cannon Rhodes College Student	Summer-2013
Addison Jezek Rhodes College Student	Summer-2013-14
	<i>Practicing Physician, Oklahoma</i>
Jeff Harding Mississippi State University	Summer-2013
Anqi Zheng Christian Brothers University	2013-2014
Kelsey Coolican Christian Brothers University	2013-2014
Alan Sun University of Memphis	Summer-2014
Alex Taylor High School Student	Summer-2014
Theresa Borcky Summer Undergraduate Internship	Summer-2014, 2015
Juanma Ramirez Visiting Graduate Student (Spain)	Summer-2014
Morgan Fuller Rhodes College Student	Summer-2015
	<i>Practicing D.O., Georgia</i>
Ken Gentner UT, Chattanooga	Summer-2015
Danny Flatten Christian Brothers University	2015-2016
Ben May Rhodes College	2016-2017
Avani Alapati Rhodes College	2017-2019
Rachel Cox UT, Knoxville	Summer-2018
Isabelle Mikell Rhodes College	Summer-2018
Drew Liess Rhodes College	2019-2020

### **RESEARCH INTERESTS AND PROJECTS**

Autism genetics  
Autism clinical phenotyping  
Angelman syndrome  
Prader-Willi syndrome  
Duplication 15q syndrome  
Epilepsy genetics  
Synaptic plasticity  
UBE3A related disorders  
Epigenetics  
Dental pulp stem cells  
Drosophila models of human disease

### **RESEARCH SUPPORT AND AWARDS**

#### **ACTIVE**

Research Award ( <b>Weese-Mayer - PI</b> )	11/01/2019-10/31/2021	0.0 cal.
ROHHAD Association	\$40,000/yr	
"RAPID-ONSET OBESITY WITH HYPOTHALAMIC DYSFUNCTION, HYPOVENTILATION, AND AUTONOMIC DYSREGULATION (ROHHAD): STEM CELL MODELS TO INVESTIGATE CAUSE AND CONSEQUENCES"		
The goal of these studies is to create a DPSC model of ROHHAD neurons for molecular studies.		
Pilot Research Award ( <b>Reiter - PI</b> )	10/01/2018-09/30/2020	1.2 cal.
Foundation for Prader-Willi Research	\$50,000 (shared grant – 2 sub-awards)	
"Assessment of epigenetic driven circadian rhythm defects in neurons from individuals with PWS"		

The purpose of this study is to detect changes in circadian rhythm and methylation in DPSC derived neurons from PWS deletion cases.

- Pilot Research Award (**Potts - PI**) 06/01/2018-05/30/2020 1.2 cal.  
 Foundation for Prader-Willi Research \$50,000 (sub-contract)  
 "Evaluating endosomal recycling pathways in primary neurons from PWS individuals"  
 The purpose of this study is to use DPSC neurons from subjects with PWS to investigate the role of MAGL2 in the pathogenesis of PWS.
- Shainberg Neuroscience Award (**Reiter-PI**) 04/01/18-03/31/19 0.6 cal.  
 LeBonheur Children's Foundation \$40,000  
 "Assessment of Protein Changes in Epileptic Brain"  
 The purpose of this study is to investigate changes in epilepsy brain related to synaptic proteins and a set of glutathione S transferase proteins. Recent work in our lab indicates that these proteins may not only be changing in Dup15q syndrome, but also other forms of epilepsy.
- R21 HD091541 (**Reiter-PI**) **5%tile** 04/01/2017-03/31/2019 (NCE) 1.2 cal.  
 NIH/NICHD, \$125,000-yr1/\$150,000-yr2  
 "An in vivo chemical screen for seizure suppression in Duplication 15q syndrome."  
 The goal of this project is to screen for FDA approved compounds that can re-activate gene expression for synaptic genes down regulated in our Dup15q fly seizure model.

#### **PENDING:**

- R01 NS099127-0A1 (**Reiter-PI**) 9/01/2020-8/31/2025 2.4 cal.  
 NIH/NINDS \$250,000/yr  
 "The role of glial cells in UBE3A over-expression induced seizures."  
 The goal of this project is to identify UBE3A substrates specifically in glial cells, which regulate seizures in our fly Dup15q seizure model.

#### **PREVIOUS:**

- R21 GRANT12726394 (**Reiter-PI**) 04/01/2019-03/31/2022 1.2 cal.  
 NIH/NINDS \$125,000/\$150,000  
 "A primer on the study of gliopathic epilepsies."  
 The goal of this project is to correlate both UBE3A expression and changes in GST proteins directly in brain material from Dup15q or idiopathic epilepsy subjects. These studies will explore the molecular nature of the gliopathic seizures in Dup15q syndrome.
- Graduate Fellowship Kevin Hope (**Reiter-Mentor**) 09/01/2015-08/31/2019 N/A  
 Duplication 15q Alliance \$100,000  
 "Investigation of Synergistic Interactions Among Genes in the 15q Duplication Syndrome"  
 This is a project using *Drosophila melanogaster* to identify combinatorial effects on the brain through over-expression of two or more 15q Duplication region genes. The goal is to better understand which genes contribute to the autism phenotype and which contribute to the seizure phenotype in 15q Duplication syndrome. Stipend Support Only.

CORNET Award ( <b>Palmer – Reiter Co-I</b> ) Vice Chancellor for Research Office	07/01/2017-06/30/18 \$47,824	N/A
<p>“Identifying chemical modulators of the Ube3a ubiquitin ligase as a therapeutic strategy to treat epileptic seizures as well as HPV related cancers.”</p> <p>The goal is to screen for chemical modulators of UBE3A activity in yeast that may provide efficacious new therapies to treat a variety of neurological disorders, as well as HPV associated cancers. <u>No Faculty Salary Support</u></p>		
Pilot Research Award ( <b>Reiter- PI</b> ) Foundation for Prader-Willi Research	09/01/2016-03/31/2018 \$178,000	1.2 cal.
<p>“Gene Expression Analysis in PWS Subject Derived Dental Pulp Stem Cell Neurons”</p> <p>The purpose of this study is to use DPSC neurons from subjects with PWS both with and without ASD in order to identify ASD-specific differential gene expression in this disorder. We will also investigate the basic changes in both mRNA and miRNA transcripts in PWS neurons versus neurotypical controls.</p>		
Shainberg Neuroscience Award ( <b>Reiter-PI</b> ) LeBonheur Children’s Foundation	03/01/2012-06/30/2017 \$130,000	0.6 cal.
<p>“Identification and Behavior Evaluation of Duplication 15q11-q13 Autism Patients in the Mid-South Region.”</p> <p>The purpose of this study is to identify unique aspects of the interstitial duplication 15q autism phenotype since these patients all share a common molecular etiology related to over-expression of the UBE3A gene. The eventual goal is to use these phenotypic features as predictors for this particular type of autism. <u>Additional funding granted to continue study.</u></p>		
Duplication 15q Alliance ( <b>Reiter-PI</b> ) Contract	04/01/2014-06/30/2018 \$60,000	1.2 cal.
<p>These funds are to direct the development of a medical informatics database for all 15q clinic sites. This database will be housed at Le Bonheur Children’s Hospital in Memphis, TN. It will contain both identifiable PHI and de-identified data from all clinic sites.</p>		
ISIS Pharmaceuticals ( <b>Reiter-PI</b> ) Contract	05/01/2015-11/01/2015 \$5000	N/A
<p>Funds paid for a student to work in Drosophila on a specific gene in my laboratory related to hypertension in humans.</p>		
UTRF Maturation Funding Program ( <b>Reiter – PI</b> ) UT Research Foundation	01/02/2015-02/02/2016 \$16,000	0.6 cal.
<p>“Blood Test for Familial Autoimmune Positive Autism Spectrum Disorder”</p> <p>The goal of this project is to develop an auto-antibody test for children with a certain sub-type of autism prior to the onset of autistic symptoms. The funds are intended for the development of this technology to eventual commercial application. <u>There is no salary support.</u></p>		
Institutional Funding ( <b>Dean</b> ) Grant Stimulus Funding	12/01/2013-11/31/2014 \$100,000	
<p>These funds are meant to supplement the lab while three NIH applications are submitted in 2014.</p>		
Duplication 15q Alliance ( <b>Reiter-PI</b> ) Pilot Research	03/01/2012- 12/01/2014 \$40,000	

“Construction of a Tet Responsive Ube3a Over-expression Mouse”

The intention of this funding is to produce a publicly available Ube3a over-expression mouse model for the study of *Ube3a* duplication autism. This project is in collaboration with Dr. Scott Dindot at Texas A&M University.

- R21 NS075709 (**Reiter-PI**) **8%tile Score** 04/01/2012 – 12/31/2014 1.2 cal.  
 NIH/NINDS, \$125,00-yr1/\$150,000-yr2  
 “Tooth pulp as a source for neuronal precursor cells to study neurogenetic disorders”  
 The goal of this project is to demonstrate that dental tooth pulp can be a good source for studying the molecular and electrophysiological properties of neurons from individuals with neurogenetic diseases.
- R01 NS059902 (**Reiter-PI**) **3.7%tile Score** 7/01/2008-8/30/2014  
 NIH/NINDS \$215,000/yr  
 “Proteomics in *Drosophila* to Identify autism candidate substrates of UBE3A”  
 The goal of this project is to identify UBE3A substrates through proteomic profiling in flies; validate that these targets interact with Dube3a or UBE3A in vitro and in vivo; and to determine if these targets are mis-regulated in the brains of Ube3a over-expression and knock-out mice.
- Center for Integrated and Translational Genomics (**Reiter –PI**) 2011 N/A  
 Pilot Project Ion Torrent Award \$5000  
 “Identification of Dube3a Regulated Transcripts by CHIPseq”  
 The purpose of this study is to screen for transcriptionally regulated targets of Dube3a in fly heads. We will also be able to identify putative binding sites for transcription factors that interact with Dube3a. No salaries paid on this grant.
- Pilot Program for Clinical Neuroscience (**McVicar - PI**) 10/01/2008-7/01/2011 0.6 cal.  
 UTHSC Neuroscience Institute \$50,000  
 “Immune Characterization in Children with Autism with and without Familial Autoimmune History”  
 The purpose of this study is to identify biomarkers in a subset of autism patients with a family history of autoimmune disease, design a protein based assay system for screening serum for these biomarkers and confirm that these serum antibodies are still present at one year’s time. No faculty salaries paid on this grant.
- Shainberg Neuroscience Award (**Reiter-PI**) 10/01/2006-3/01/2012 0.6 cal.  
 LeBonheur Children’s Foundation \$50,000/yr  
 “Identification and Behavior Evaluation of Duplication 15q11-q13 Autism Patients in the Mid-South Region.”  
 The purpose of this study is to identify unique aspects of the interstitial duplication 15q autism phenotype since these patients all share a common molecular etiology related to over-expression of the UBE3A gene. The eventual goal is to use these phenotypic features as predictors for this particular type of autism. No salaries are paid on this grant.
- Pilot Research Award (**Reiter-PI**) 12/01/2007-11/30/2008 4.0 cal.  
 Autism Speaks \$60,000/yr  
 “Identification of UBE3A substrates using proteomic profiling in *Drosophila*.”  
 The major goal of this project is to identify the protein targets of human and fly UBE3A via mis-expression proteomic analysis in *Drosophila* heads. This grant pays salary for a postdoctoral fellow. Ended early due to overlap with R01.

Pilot Research Award (**Reiter- PI**) 12/01/2008-11/30/2009 1.2 cal.  
 Angelman Syndrome Foundation \$77,866  
 “A combined molecular and electrophysiological approach to understanding cerebellar defects in Angelman syndrome.” This study in collaboration with Dr. Detlef Heck in the Department of Anatomy and Neurobiology is designed to identify critical cell types in the cerebellum that are affected by the loss of Ube3a and therefore may contribute to the Angelman syndrome phenotype.

Charcot-Marie-Tooth Disease Association, Postdoctoral Fellowship Award. (**Lupski-PI**) (1998-1999).

Glaucoma Foundation, Postdoctoral Fellowship Award: “Role of cytochrome 450 *CYP1B1* in the pathogenesis of primary congenital glaucoma.” (**Bier-PI**) (2000-2001).

Cure Autism Now, Young Investigator Award: “Identification of genes involved in autism using a *Drosophila* UBE3A mis-expression system.” (**Reiter-PI, Bier-mentor**) (2003-2005)

Cure Autism Now, Pilot Research Award: “A proteomics approach to the Identification and characterization of protein targets regulated by UBE3A” (**Reiter-PI**) \$120,000 (4/1/06-3/31/08).

#### **PROFESSIONAL SOCIETIES**

American Society of Human Genetics  
 Genetics Society of America  
 International Society for Autism Research  
 American Society for Molecular Biology

#### **SCIENTIFIC ADVISORY BOARDS**

15q Alliance – Scientific Advisory Board	2008-2018
idic15 Canada – Scientific Advisory Board	2012-2013

#### **EDITORIAL BOARD APPOINTMENTS**

Open Access (London) – *Autism* (2013-2015)  
*Austin Journal of Autism & Related Disabilities* (2015-present)  
*Scientific Reports* (2016-present)  
 Associate Editor, *Frontiers in Genetics* – Genetic Disorders (2019-present)

#### **MEETINGS ATTENDED (since 2005)**

American Society of Human Genetics, Salt Lake City, UT (2005)  
 Genetics Society of America, Model Organisms Meeting, San Diego, CA (2006)  
 47<sup>th</sup> Annual *Drosophila* Research Conference, Houston, TX (2006)  
 American Society of Human Genetics, New Orleans, LA (2006)  
 IDEAS Conference (chromosome 15q duplications), Boston, MA (2007)  
 Angelman Syndrome Foundation Biennial Conference, St. Louis, MO (2007)  
 American Society of Human Genetics, San Diego, CA (2007)  
 49<sup>th</sup> Annual *Drosophila* Research Conference, San Diego, CA (2008)  
 American Society of Human Genetics, Philadelphia, PA (2008)  
 IDEAS Conference (chromosome 15q duplications), Indianapolis, IN (2009)  
 Angelman Syndrome Foundation Biennial Conference, Orlando, FL (2009)  
 American Society of Human Genetics, Honolulu, HI (2009)

International Meeting for Autism Research, Philadelphia, PA (2010)  
 Angelman Syndrome Foundation Scientific Conference, Chapel Hill, NC (2010)  
 American Society of Human Genetics, Washington, D.C. (2010)  
 International Meeting for Autism Research, San Diego, CA (2011)  
 IDEAS Conference (chromosome 15q duplications), Philadelphia, PA (2011)  
 Society for Neuroscience, Washington, DC (2011)  
 53<sup>rd</sup> Annual Drosophila Research Conference, Chicago, IL (2012)  
 15q Alliance Scientific Meeting, Boston, MA (2012)  
 International Meeting for Autism Research, San Sebastian, Spain (2013)  
 15q Alliance Scientific Meeting, Sacramento, CA (2013)  
 55<sup>th</sup> Annual Drosophila Research Conference, San Diego, CA (2014)  
 PRISMS Meeting, St. Louis, MO (2014)  
 CdLS Meeting, Los Angeles, CA (2014)  
 Joint 15q Alliance and ASF Meeting, Boston, MA (2014)  
 Potocki-Lupski Syndrome Meeting, Houston, TX (2015)  
 15q Alliance Meeting, Orlando, FL (2015)  
 Joint 15q Alliance and ASF Meeting, Boston, MA (2016)  
 The Allied Genetics Conference, Presenter, Orlando, FL (2016)  
 Foundation for Prader-Willi Syndrome Research Meeting, Providence, RI (2016)  
 15q Alliance Meeting, Los Angeles, CA (2017)  
 Foundation for Prader-Willi Syndrome Research Meeting, Indianapolis, IN (2017)  
 American Society for Human Genetics, Orlando, FL (2017)  
 European Society for Human Genetics, Milan, Italy (2018)  
 Joint 15q Alliance and ASF Meeting, Chapel Hill, NC (2018)  
 Invited Speaker, Foundation for Prader-Willi Research, Las Vegas, NV (2018)  
 Invited Session (co-chair), Glial Health and Disease, Porto, Portugal (2019)  
 Angelman Syndrome Foundation Meeting, Invited Speaker, Louisville, KY (2019)  
 Foundation for Prader-Willi Syndrome Research Meeting, New Orleans, LA (2019)

### **INVITED SEMINARS**

Invited Speaker F.A.S.E.B. Meeting	1997
Invited Speaker A.S.H.G. Meeting	1997
Invited Speaker A.S.H.G. Meeting	2002
Invited Speaker A.S.H.G. Meeting	2003
Co-chair model organisms session A.S.H.G.	2005
Invited Speaker, Epigenetics & Neural Developmental Disorders	2006
Invited Speaker, Christian Brothers University	2007
Invited Speaker, Vanderbilt University	2007
Invited Session/Speaker "The Duplication 15q syndrome", A.S.H.G.	2008
Invited Speaker, Vanderbilt University	2008
Invited Speaker, University of Toronto, Toronto, Canada	2009
Invited Speaker, NEMOURS, University of Delaware	2009
Invited Speaker, ASF Meeting, UNC, Chapel Hill, NC	2010
Invited Speaker, Vanderbilt Kennedy Center	2010
Invited Speaker A.S.H.G. Meeting	2010
Co-chair session A.S.H.G. Meeting	2010
Invited Speaker, Baylor College of Medicine, Houston, TX	2011
Invited Mini-Symposium, Society for Neuroscience Meeting	2011
Invited Speaker, ASF Meeting, Rockville, MD	2012

Co-organizer, 15q Alliance Meeting, Boston, MA	2012
Invited Speaker, CIC bioGUNE, Bilbao, Spain	2013
Invited Speaker, ASF Meeting, Orlando, FL	2013
Invited Speaker, 15q Alliance Meeting, MIND Institute, CA	2013
Invited Speaker, co-organizer, joint 15q Duplication and ASF Meeting, Boston	2014
Invited Speaker, St. Louis University Medical School	2014
Invited Speaker, Grand Rounds, Minnesota Epilepsy Group, St. Paul, MN	2015
Invited Speaker, PTLIS Meeting, Houston, TX	2015
Invited Speaker, University of Toronto, Toronto, Canada	2015
Invited Speaker, Foundation for Prader-Willi Research, Austin, TX	2015
Invited Speaker, Pediatric Neurology Conference, Memphis, TN	2016
Invited Speaker, Harvard Mass General Hospital, Neurology, Boston, MA	2016
Invited Speaker, TAGC, Human Disease Models II, Orlando, FL	2016
Invited Speaker, Foundation for Prader-Willi Research, Providence, RI	2016
Invited Speaker, University of Kentucky, Biochemistry, Lexington, KY	2017
Invited Speaker, Foundation for Prader-Willi Research, Indianapolis, IN	2017
Invited Speaker, Universita' degli Studi di Torino, Torino, Italy	2018
Invited Speaker, Foundation for Prader-Willi Research, Las Vegas, NV	2018
Invited Session/Speaker, Glia in Health and Disease Meeting, Porto, Portugal	2019
Invited Speaker, Angelman Syndrome Foundation Meeting, Louisville, KY	2019
Invited Speaker, Neuroscience Institute, UTHSC, Memphis, TN	2019
Invited Speaker, Dept. of Pharmacology, UTHSC, Memphis, TN	2020

### **NIH REVIEWS**

National Institutes of Health, American Recovery & Reinvestment Act of 2009 (ARRA), *ad hoc* 2009  
 National Institutes of Health, Fragile X Syndrome Research Center (FXSRC), *ad hoc* 2013  
 National Institutes of Health, Molecular Neurogenetics [MNG] Study Section, *ad hoc* 2015, 2016, 2017  
 National Institutes of Health, Director's Pioneer Award (DP1), Mail Reviewer, *ad hoc* 2018, 2019, 2020  
 National Institutes of Health, Neurodevelopment, Synaptic Plasticity, and Neurodegeneration panel (F03A) *ad hoc* 2018

### **GRANT REVIEWER** (2005-present; \*within the last year)

Autism Speaks (Postdoctoral Fellowship Awards)  
 CTSI-NTMP Grants  
 Fight for Sight, Grant in Aid Reviewer  
 Foundation for Prader-Willi Research, Pilot Research Grants Reviewer\*  
 German-Israeli Foundation for Scientific Research and Development (Germany/Israel)  
 Israeli National Science Foundation (Israel)  
 Italian Ministry of Health (Italy)  
 KUL Research Professorship  
 Medical Research Council (UK)  
 Netherlands Organization for Health Research and Development, Reviewer  
 Oak Ridge Associated Universities  
 Swiss National Science Foundation (Switzerland)  
 Swiss 3RCC Grants (Switzerland)

**JOURNAL AND ABSTRACT REVIEWER** (1997-present)

Acta Neurologica Belgica  
Acta Neuropathologica Communications  
African Journal of Biotechnology  
American Society for Human Genetics, Abstracts (2010, 2012)  
American Journal of Medical Genetics  
American Journal of Human Genetics  
ASN Neuro  
Autism Research  
BBA - Gene Structure and Expression  
BMC Cell Biology  
BMC Genetics  
BMC Bioinformatics  
Biochemical Journal  
Biological Psychiatry  
Bioinformatics  
BioMed Research International  
Diseases  
Epigenetics  
European Child & Adolescent Psychiatry  
European Journal of Medical Genetics  
Gene  
Genome Research  
Frontiers in Genetics  
Human Molecular Genetics  
International Society for Autism Research Meeting, Abstracts (2012, 2013)  
Journal of Neurodevelopmental Disorders  
Journal of Neurochemistry  
Journal of Neuroscience  
Journal of Pediatric Genetics  
Journal of Visual Experimentation  
Neurobiology of Disease  
Neurogenetics  
Neuroscience  
PLoS One  
PLoS Genetics  
Stem Cell Research  
Stem Cell Research Translational Medicine  
Stem Cell Research and Therapy  
The American Journal of Psychiatry  
Translational Research

**PRESS**

2013-present Spectrum News (Simons Foundation for Autism Research)  
Comments and Articles (13 total) (<https://spectrumnews.org/?s=Lawrence+Reiter>)  
2014 People Behind the Science Interview (<http://www.peoplebehindthescience.com/dr-larry-reiter>)  
2018 Memphis Parent, *Autism Myths* (<https://memphisparent.com/health/autism-myths/>)

**IMPACT ON THE LITURATURE**

h-index (number of publications cited x times) = **27**

i10-index (number of publication with at least 10 citations) = **40**

Publications cited >100 times = **5**

Number of citations (Google Scholar) = **3301**

ResearchGate Score = **44.14**

**LINKS TO SCHOLARLLY RANKINGS**

ResearchGate (**Scientific Community**): ([https://www.researchgate.net/profile/Lawrence\\_Reiter](https://www.researchgate.net/profile/Lawrence_Reiter))

Google Scholar (**Publications**): (<https://scholar.google.com/citations?user=-kaQJ14AAAAJ&hl=en>)

Publons (**Reviews**): (<https://publons.com/a/530014/>)

Associate Editor (*Frontiers in Genetics*): (<https://loop.frontiersin.org/people/374636/editorial>)

**PEER REVIEWED PUBLICATIONS**

- 1) Hope KA, Johnson D, Miller PW, Lopez-Ferrer D, Kakhniashvili D, **Reiter L.T.** Transcriptomic and proteomic profiling of glial versus neuronal Dube3a overexpression reveals common molecular changes in gliopathic epilepsies. *Neurobiol Dis.* 2020 Apr 25:104879. PubMed PMID: 32344153.
- 2) DiStefano, C., Wilson, R.B., Hyde, C., Cook, E.H., Thibert, R.L., **Reiter, L.T.**, Vogel-Farley, V., Hipp, J., Jeste, S. Behavioral characterization of dup15q syndrome: Toward meaningful endpoints for clinical trials. *Am J Med Genet A.* 2019 Oct 26. [Epub ahead of print]
- 3) Hope, K.A., Flatten, D., Cavitch, P., May, B., Sutcliffe, J.S., O'Donnell, J., **Reiter, L.T.**, The *Drosophila* Gene *Sulfateless* Modulates Autism-like Behaviors. *Front Genet.* 2019 10:574.
- 4) Frohlich, J., **Reiter L.T.**, Saravanapandian, V., DiStefano, C., Huberty, S., Hyde, C., Chamberlain, S., Bearden, C.E., Golshani, P., Irimia, A., Olsen, R.W., Hipp, J.F., Jeste, S.S. Mechanisms underlying the EEG biomarker in Dup15q syndrome. *Mol Autism.* 2019 Jul 3;10:29. Erratum in: *Mol Autism.* 2019 Nov 6;10:37.
- 5) Hope, K.A., McGinn, A., & **Reiter, L.T.** A genome wide enhancer/suppressor screen for *Dube3a* interacting genes in *Drosophila melanogaster*. *Sci Rep*, 2019 Feb 20; 9(1):2382.
- 6) Toro C, Hori RT, Malicdan MCV, Tiff CJ, Goldstein A, Gahl WA, Adams DR, Fauni HB, Wolfe LA, Xiao J, Khan MM, Tian J, Hope KA, **Reiter LT**, Tremblay MG, Moss T, Franks AL, Balak C; C4RCD Research Group, LeDoux MS. A recurrent de novo missense mutation in UBTF causes developmental neuroregression. *Hum Mol Genet.* 2018 Apr1;27(7):1310.
- 7) Urraca N., Hope K.A., Victor A.K., Belgard T.G., Memon R., Goorha S., Valdez C., Tran Q.T., Sanchez S., Ramirez J., Donaldson M., Bridges D. and **Reiter L.T.** Significant transcriptional changes in 15q duplication but not Angelman syndrome deletion stem cell-derived neurons. *Molecular Autism* 2018, 9:6.
- 8) Ajayi O.J., Smith E.J., Viangteeravat T., Huang E.Y., Nagisetty N.S.V.R., Urraca N., Lusk L., Finucane B., Arkilo D., Young J., Jeste S., Thibert R., Dup15q Alliance, **Reiter L.T.** Multisite Semiautomated Clinical Data Repository for Duplication 15q Syndrome: Study Protocol and

- Early Uses. *JMIR Res Protoc*. 2017; **6(10)**:e194. PubMed PMID: 29046268.
- 9) Hope K.A., LeDoux M.S., **Reiter L.T.** Glial overexpression of Dube3a causes seizures and synaptic impairments in *Drosophila* concomitant with down regulation of the Na<sup>+</sup>/K<sup>+</sup> pump ATPα. *Neurobiology of Disease*. 2017; **108**:238-248. PMID: 28888970.
  - 10) Copping N., Christian S., Ritter D., Islam M.S., Buschner N., Zolkowska D., Pride M., Berg E., LaSalle J., Ellegood J., Lerch J., **Reiter L.T.**, Silverman J., Dindot S. Neuronal overexpression of *Ube3a* isoform 2 causes behavioral impairments and neuroanatomical pathology relevant to 15q11.2-q13.3 duplication syndrome. *Human Molecular Genetics*. 2017; **26(20)**:3995-4010. PMID: 29016856.
  - 11) Dunaway K., Goorha S., Matelski L., Urraca N., Lein P.J., Korf I., **Reiter L.T.**, LaSalle JM. Dental Pulp Stem Cells Model Early Life and Imprinted DNA Methylation Patterns. *Stem Cells*. 2017; **35(4)**:981-988. PubMed PMID: 28032673.
  - 12) Frohlich J., Senturk D., Saravanapandian V., Golshani P., **Reiter L.T.**, Sankar R, Thibert RL, DiStefano C., Huberty S., Cook E.H., Jeste S.S. A Quantitative Electrophysiological Biomarker of Duplication 15q11.2-q13.1 Syndrome. *PLoS One*. 2016; **11(12)**:e0167179. PubMed: 27977700.
  - 13) N. Urraca, B. Potter, R. Hundley, E. Pivnick, K. McVicar, R. Thibert, C. Ledbetter, R. Chamberlain, L. Miravalle, C. Sirois, S. Chamberlain, **L.T. Reiter**. A rare inherited 15q11.2-q13.1 interstitial duplication with maternal somatic mosaicism, renal carcinoma and autism. *Front. Genet*. 2016; **7**:205. PubMed PMID: 27933089.
  - 14) K.A. Hope, M.S. LeDoux, **L.T. Reiter**. The *Drosophila melanogaster* homolog of *UBE3A* is not imprinted in neurons. *Epigenetics*. 2016; **11**:1-6. PMID: 27599063.
  - 15) C. DiStefano, A. Gulsrud, S. Huberty, C. Kasari, E. Cook, **L.T. Reiter**, R. Thibert, S.S. Jeste. Identification of a distinct developmental and behavioral profile in children with Dup15q syndrome. *J Neurodev Disord*. 2016; **6(8)**:19. PMID: 27158270.
  - 16) I. El-Iyachi, S. Goorha, **L.T. Reiter**, G. Miranda-Carboni. Effects of hTERT immortalization on osteogenic and adipogenic differentiation of dental pulp stem cells. *Data in Brief*. 2016; **6**:696-9. PMID: 26958627.
  - 17) N. Urraca, R. Memon, I. El-Iyachi, S. Goorha, C. Valdez, Q. Tran, R. Scroggs, G. Miranda-Carboni, M. Donaldson, D. Bridges, **L.T. Reiter**. Characterization of neurons from immortalized dental pulp stem cells for the study of neurogenetic disorders. *Stem Cell Res*. 2015; **15(3)**:722-730. PMID: 26599327.
  - 18) R. Wilson, N. Urraca, C. Skobowait, K. Hope, L. Miravalle, R. Chamberlin, M. Donaldson, T. Seagroves, **L.T. Reiter**. Assessment of the tumorigenic potential of spontaneous and hTERT-immortalized cultured dental pulp stem cells. *Stem Cell Trans. Med*. 2015; Jun 1. pii: sctm.2014-0196.
  - 19) I. Hatfield, I. Harvey, E. Yates, J. Redd, **L.T. Reiter**, D. Bridges. The role of TORC1 in muscle development in *Drosophila*. *Sci Rep*. 2015; **13(5)**:9676.
  - 20) C. Valdez, R. Scroggs, R. Chasen, **L.T. Reiter**. Variation in Dube3a expression affects

- neurotransmission at the *Drosophila* neuromuscular junction. *Biology Open*. 2015; May 6. pii: bio.20148045.
- 21) N. Germain, P. Chen, A. Plocik, H. Glatt-Deeley, J. Brown, J. Fink, K. Bolduc, T. Robinson, E. Levine, **L.T. Reiter**, B. Graveley, M. Lalande, S. Chamberlain. Gene expression analysis of human induced pluripotent stem cell-derived neurons carrying copy number variants of chromosome 15q11-q13.1. *Molecular Autism*. 2014; **5**:44.
  - 22) K. Conant, B. Finucane, N. Cleary, A. Martin, C. Muss, M Delany, E. Murphy, O. Rabe, K. Luchsinger, S. Spence, N. C. Schanen, O. Devinsky, E. Cook, J. LaSalle, **L.T. Reiter**, R. Thibert. A survey of seizures and current treatments in 15q duplication syndrome. *Epilepsia*. 2014; **55(3)**:396-402.
  - 23) L. Jensen, M.F. Farook, **L.T. Reiter**. Proteomic profiling in *Drosophila* reveals potential Dube3a regulation of the actin cytoskeleton and neuronal homeostasis. *PLoS One*. 2013; **8(4)**:e61952.
  - 24) N. Urraca, J. Cleary, V. Brewer, E.K. Pivnick, K. McVicar, R.L. Thiber, N.C. Schanen, C. Esmer, D. Lamport, **L.T. Reiter**. The Interstitial Duplication 15q11.2-q13 Syndrome Includes Autism, Mild Facial Anomalies and a Characteristic EEG Signature. *Autism Res*. 2013; **6(4)**:268-79.
  - 25) M. Farook, M. DeCuypere, K. Hyland, T. Takumi, M.S. LeDoux, **L.T. Reiter**. Altered serotonin, dopamine and norepinephrine levels in 15q duplication and Angelman syndrome mouse models. *PLoS One*. 2012;**7(8)**:e43030.
  - 26) H.A. Scoles, N. Urraca, S. W. Chadwick, **L.T. Reiter**, J.M. Lasalle. Increased copy number for methylated maternal 15q duplications leads to changes in gene and protein expression in human cortical samples. *Mol Autism*. 2011 Dec **12;2(1)**:19.
  - 27) S. Roy, Y. Zhao, M. Allensworth, M.F. Farook, M.S. Ledoux, **L.T. Reiter**, D.H. Heck. Comprehensive motor testing in *Fmr1*-KO mice exposes temporal defects in oromotor coordination. *Behav Neurosci*. 2011 Dec;**125(6)**:962-9. Epub 2011 Oct 17. PubMed PMID: 22004265.
  - 28) M. Wangler, **L.T. Reiter**, G. Zimm, J. Trimble-Morgan, J. Wu, E. Bier. The antioxidant proteins TSA and PAG interact synergistically with Presenilin to modulate Notch signaling in *Drosophila*. *Protein & Cell*. **2**: 554-63.
  - 29) M. Allensworth, A. Saha, **L.T. Reiter** and D. Heck. 2011. Normal social seeking behavior, hypoactivity and reduced exploratory range in a mouse model of Angelman syndrome. *BMC Genetics*. **12**: 7.
  - 30) F. Ferdousy, W. Bodeen, K. Summers, O. Doherty, Wright O, N. Elsis, G. Hilliard, J. O'Donnell, **L.T. Reiter**. 2010. *Drosophila* Ube3a regulates monoamine synthesis by increasing GTP cyclohydrolase I activity via a non-ubiquitin ligase mechanism. *Neurobiology of Disease*, **41(3)**: 669-77.

- 31) N. Urraca, L. Davis, E. Cook, N.C. Schanen and **L.T. Reiter**. 2010. A single tube quantitative high resolution melting curve method for parent of origin determination of 15q duplications. *Genetic Testing and Molecular Biomarkers*. **14(4)**:571-6.
- 32) S. Doronkin, I. Djagaeva, M. Nagle, **L.T. Reiter**, T. Seagroves. 2009. Dose-dependent modulation of HIF-1alpha/sima controls the rate of cell migration and invasion in *Drosophila* ovary border cells. *Oncogene*, **29(8)**:1123-34.
- 33) D. Heck, Y. Zhao, S. Roy, M. LeDoux and **L.T. Reiter**. 2008. Analysis of Cerebellar Function in Ube3a Deficient Mice Reveals Novel Genotype Specific Behaviors. *Human Molecular Genetics*, **17**:2181-89.
- 34) **L.T. Reiter**, L. Do, M. Fischer, N. Hong and E. Bier. 2007. Accentuate the negative: Proteome comparisons using the Negative Proteome Database. *Fly*. **1**:164-71.
- 35) **L.T. Reiter**, T. Seagroves, M. Bowers and E. Bier. 2006. Expression of the Rho-GEF Pbl/ECT2 is regulated by the UBE3A E3 ubiquitin ligase. *Human Molecular Genetics*. **15**:2825-35.
- 36) **L.T. Reiter** and E. Bier. 2002. Using *Drosophila melanogaster* to uncover human disease gene function and potential drug target proteins. *Expert Opinion on Therapeutic Targets*. **6**:387-399.
- 37) S. Chien, **L.T. Reiter**, E. Bier and M. Gribskov. 2002. Homophila: human disease gene congenates in *Drosophila*. *Nucleic Acids Research* **30**:149-51.
- 38) K. Inoue, K. Dewar, N. Katsanis, **L.T. Reiter**, E. Lander, K. Devon, D. Wyman, J. Lupski and B. Birren. 2001. The 1.4-Mb CMT1A duplication/HNPP deletion genomic region reveals unique genome architectural features and provides insights into the recent evolution of new genes. *Genome Research* **11**:1018-1033.
- 39) **L.T. Reiter**, L. Potocki, M. Gribskov and E. Bier. 2001. A systematic analysis of human disease associated gene sequences in *Drosophila melanogaster*. *Genome Research* **11**:1114-1125.
- 40) T. Liehr, **L.T. Reiter**, J. Lupski, T. Murakami, U. Cluassen and B. Rautenstrauss. 2001. Regional localization of 10 mariner transposon-like ESTs by means of FISH-evidence for a correlation with fragile sites. *Mammalian Genome* **12**:326-328.
- 41) **L.T. Reiter**, T. Liehr, B. Rautenstrauss, H. Robertson and J. Lupski. 1999. Localization of *mariner* DNA transposons in the human genome by PRINS. *Genome Research* **9**:839-843.
- 42) **L.T. Reiter**, P. Hastings, E. Nelis, P. De Jonghe, C. Van Broekhoven and J. Lupski. 1998. Human meiotic recombination products revealed by sequencing a hotspot for homologous strand exchange in multiple HNPP deletion patients. *American Journal of Human Genetics* **62**:1023-1033.
- 43) V. Timmerman, B. Rautenstrauss, **L.T. Reiter**, T. Koeuth, A. Löfgren, T. Leibr, E. Nelis, K. Bathke, P. De Jonghe, H. Grehl, J.- J. Martin, J. Lupski and C. Van Broeckhoven. 1997.

- Detection of the CMT1A/HNPP recombination "hotspot" in unrelated patients of European descent. *Journal of Medical Genetics* **34**:43-49.
- 44) T. Murakami, **L.T. Reiter** and J. Lupski. 1997. Genomic structure and expression of the human heme A:farnesyltransferase (*COX10*) gene. *Genomics* **42**:161-164.
- 45) **L.T. Reiter**, T. Murakami, T. Koeuth, R. Gibbs and J. Lupski. 1997. The human *COX10* gene is disrupted during homologous recombination between the 24-Kb proximal and distal CMT1A-REPs. *Human Molecular Genetics* **6**:1595-1603.
- 46) **L.T. Reiter**, T. Murakami, T. Koeuth, L. Pentao, M. Muzny, R. Gibbs and J. Lupski. 1996. A recombination hotspot responsible for two inherited peripheral neuropathies is located near a *mariner* transposon-like element. *Nature Genetics* **12**:288-297.
- 47) **L.T. Reiter**, D. Steffen, S. Shapira, J. Lupski, M. Frazier and D. Wheeler. 1995. A searchable video database of dysmorphology. *Journal of the American Medical Informatics Association (symposium supplement SCAMC Proceedings)*, p. 1000.
- 48) J. Hubar, M. Oeschger and **L.T. Reiter**. 1994. Effectiveness of radiographic film barrier envelopes. *General Dentistry* **42**:406-408.

## REVIEWS AND BOOK CHAPTERS

- 1) Hope, K.A. & **Reiter L.T.** Understanding Human Genetic Disease With the Fly. In: Walz K and Young JI, (eds). *Cellular and Animal Models in Human Genomics Research (Translational and Applied Genomics Series)*. San Diego: Elsevier Inc./Academic Press, 2019: pp-pp. 69-87.
- 2) Victor A.K. & **Reiter L.T.** Dental Pulp Stem Cells for the Study of Neurogenetic Disorders. *Human Molecular Genetics*. 2017. **26(R2)**:R166-R171. PMID: 28582499.
- 3) Chow C. & **Reiter L.T.** Etiology of Human Disease On the Fly. *Trends in Genetics*. 2017. **(6)**:391-398. PubMed PMID: 28420493.
- 4) Goorha S. & **Reiter L.T.** Culturing and Neuronal Differentiation of Human Dental Pulp Stem Cells. *Curr Protoc Hum Genet*. 2017. **92**:21.6.1-21.6.10. PubMed PMID: 28075485.
- 5) Finucane BM, Lusk L, Arkilo D, Chamberlain S, Devinsky O, Dindot S, Jeste SS, LaSalle JM, **Reiter L.T.**, Schanen NC, Spence SJ, Thibert RL, Calvert G, Luchsinger K, Cook EH Jr. 15q Duplication Syndrome and Related Disorders. 2016 Jun 16. In: Pagon RA, Adam MP, Ardinger HH, Wallace SE, Amemiya A, Bean LJH, Bird TD, Fong CT, Mefford HC, Smith RJH, Stephens K, editors. *GeneReviews*® [Internet]. Seattle (WA): University of Washington, Seattle; 1993-2016. Available from <http://www.ncbi.nlm.nih.gov/books/NBK367946/> PubMed PMID: 27308687.
- 6) J. LaSalle, **L.T. Reiter**, S. Chamberlain. Epigenetic regulation of UBE3A and roles in human neurodevelopmental disorders. *Epigenomics*. 2015 Oct;**7(7)**:1213-28. PMID: 26585570.
- 7) R. Ajjuri, M. Hall, **L.T. Reiter** and J. O'Donnell. 2014. *Drosophila* In: Mark S. LeDoux (ed): *Movement Disorders, 2<sup>nd</sup> Edition: Genetics and Models*. pp. 77-93.

- 8) N. Urraca and **L.T. Reiter**. 2013. Developmental disabilities, autism and schizophrenia at a single locus: complex gene regulation and genomic instability of 15q11-q13 cause a range of neurodevelopmental disorders. In: J. Rubenstein and P. Rakic (eds): Neural Circuit Development and Function in the Healthy and Diseased Brain. pp. 617-630.
- 9) S. Doronkin and **L.T. Reiter**. 2008. *Drosophila* orthologues to human disease genes: an update on progress. *Prog Nucleic Acid Res Mol Biol.* **82**:1-32. Book Chapter. PMID: 18929137.
- 10) C. Pflieger and **L Reiter**. 2008. Recent efforts to model human diseases in vivo in *Drosophila. Fly (Austin)*. **May 23;2(3)**. Review. PMID:18820469.
- 11) **L.T. Reiter** and E. Bier. 2002. Using *Drosophila melanogaster* to uncover human disease gene function and potential drug target proteins. *Expert Opin Ther Targets.* Jun;**6(3)**:387-99. Review. PMID: 12223075.
- 12) C. Boerkoel, K. Inoue, **L.T. Reiter**, L. Warner, J. Lupski. 1999. Molecular mechanisms for CMT1A duplication and HNPP deletion. *Ann N Y Acad Sci.* **883**:22-35. Review. PMID: 10586226.
- 13) **L.T. Reiter**, T. Murakami, L. Warner and J. Lupski. 1996. DNA rearrangements effecting a dosage sensitive gene are responsible for two commonly inherited neuropathies. In Edward R. B. McCabe (ed): "The Molecular Medicine of Mental Retardation and Developmental Disabilities." Mental Retardation and Developmental Disabilities Research Reviews, John Wiley and Sons, Inc., New York **2**:139-146.
- 14) L. Warner, **L.T. Reiter**, T. Murakami, J. Lupski. 1996. Molecular mechanisms for Charcot-Marie-Tooth disease and related demyelinating peripheral neuropathies. *Cold Spring Harb Symp Quant Biol.* **61**:659-71. Review. PMID: 9246492.
- 15) T. Murakami, C. Garcia, **L.T. Reiter**, and J. Lupski. 1996. Reviews in molecular medicine: Charcot-Marie-Tooth disease and related inherited neuropathies. *Medicine* **75**:233-250. Review. PMID: 8862346.

#### OTHER PUBLICATIONS

- 1) **L.T. Reiter**. 2004. Barstow to Vegas the Hard Way 2003. *Vintage Motocross and Dirt Bike Quarterly VMX* **23**:65-67.
- 2) **L.T. Reiter**. 2016. Your Classics Section: 1964 Triumph Desert Sled. *Classic Bike*. Feb: 24.