

CURRICULUM VITAE

NAME: Xuejun Wang, M.D., Ph.D.

NICKNAME: “XJ” Wang

CITIZENSHIP: U.S.A.

AFFILIATION: Division of Basic Biomedical Sciences
Sanford School of Medicine
University of South Dakota
414 E Clark St., Lee Medical Science Building
Vermillion, SD 57069
USA
Phone: 605 658-6345
Fax: 605 677-6381
E-mail: xuejun.wang@usd.edu
<http://www.usd.edu/faculty-and-staff/Xuejun-Wang>

RANK AND/OR TITLE: Tenured Full Professor of Basic Biomedical Sciences
Director, the MD/PhD Program

ORCID: <https://orcid.org/0000-0001-9267-1343>

Scopus Profile: <https://www.scopus.com/authid/detail.uri?authorId=35235510300>

Google Scholar: <https://scholar.google.ca/citations?user=ipQguqMAAAAJ&hl=en>

EDUCATION

September 1980 to July 1985

Hubei Medical University (now Wuhan University College of Medicine), Wuhan, Hubei, China
Bachelor of Medicine in Clinical Medicine (M.D. equivalent)

September 1985 to July 1988

Hubei Medical University (now Wuhan University College of Medicine), Wuhan, Hubei, China
M.S. in Pathophysiology (Advisor: Chuanren Dong, M.D.)
Thesis Title: Hastened Plasma Coagulation and Thrombosis Contribute to The Induction of Myocardial Infarction/Necrosis by Isoproterenol in Rats.

January 1996 to August 1998

University of South Dakota College of Medicine, Vermillion, South Dakota, USA
Ph.D. in Anatomy and Structural Biology (Advisor: A. Martin Gerdes, Ph.D.)
Dissertation Title: Cardiomyocyte Remodeling in Chronic Pressure Overload Cardiac Hypertrophy and Heart Failure in Guinea Pigs.

POSTDOCTORAL TRAINING

September 1998 to September 2001

Postdoctoral Fellow

Advisor: Jeffrey Robbins, Ph.D.

Division of Molecular Cardiovascular Biology
Cincinnati Children's Hospital Medical Center
Cincinnati, Ohio

AHA Postdoctoral Fellowship project title: Dissecting Desmin-related Cardiomyopathy with Mouse Transgenesis.

ACADEMIC/ADMINISTRATIVE APPOINTMENTS

September 2006 – present

Professor and Director
The MD/PhD Program
Sanford School of Medicine
University of South Dakota
Vermillion, SD

September 2010 - July 2016

Director
The Interim PQCD Research Center
USD Sanford School of Medicine
Vermillion, SD

June 2006

Tenure granted by University of South Dakota

June 2005 – August 2006

Associate Professor (Tenure Track/Tenured)
Cardiovascular Research Institute,
Sanford School of Medicine of the University of South Dakota
Sioux Falls, SD

October 2001 – May 2005

Assistant Professor (Tenure Track)
Cardiovascular Research Institute
University of South Dakota School of Medicine
Sioux Falls, SD

September 1998 – September 2001

Research Fellow
Division of Molecular Cardiovascular Biology
Cincinnati Children's Hospital Medical Center
Cincinnati, Ohio

January 1996 – August 1998

Graduate Assistant
Department of Anatomy and Structural Biology
University of South Dakota School of Medicine
Vermillion, SD

November 1994 – December 1995

Research Associate
Department of Anatomy and Structural Biology
University of South Dakota School of Medicine
Vermillion, SD

July 1993 – October 1994

Associate Professor
Department of Pathophysiology
Hubei Medical University (now Wuhan University College of Medicine)
Wuhan, Hubei, China

July 1988 – June 1993

Instructor
Department of Pathophysiology
Hubei Medical University (now Wuhan University College of Medicine)
Wuhan, Hubei, China

September 1985 – June 1988

Graduate Teaching Assistant
Department of Pathophysiology
Hubei Medical University (now Wuhan University College of Medicine)
Wuhan, Hubei, China

SPECIAL HONORS OR RECOGNITIONS

- | | |
|------|---|
| 1988 | Best Original Research Article Award
Society of Pathophysiology of Hubei Province, P.R. China |
| 1988 | Best Original Research Articles in Natural Sciences (3 rd prize)
The Science & Technology Association of Hubei Province, P. R. China |
| 1990 | Best Original Research Articles in Natural Sciences (2 nd prize)
The Science and Technology Association of Hubei Province, P. R. China |
| 1991 | Best Original Scientific Research Articles (2 nd prize)
The Science and Technology Association of Wuhan, P.R. China |
| 1993 | The Medical Sciences and Technology Advancement Awards (2 nd prize)
The Department of Health, Hubei Province, China |
| 1995 | Best Original Scientific Research Articles (3 rd prize)
The Science and Technology Association of Wuhan, the Commission of Science and Technology of Wuhan, and the Department of Personnel of Wuhan, China |
| 1995 | The Sciences and Technology Advancement Awards (3 rd prize)
The Government of Hubei Province, China |

- 2000 Postdoctoral Fellowship
American Heart Association (AHA) Ohio Valley Affiliate
- 2000 Young Investigator Award
Heart Failure Society of America (HFSA)
- 2001 Young Investigator Award
International Society for Heart Research-North American Section (ISHR-NAS)
- 2002 Best Abstract Award Finalist
ISHR-North American Section Annual Meeting
- 2002 Scientist Development Award
American Heart Association (AHA) National Center
- 2003 Distinguished Performance Award
Vice President for Health Affairs and Office of the Dean, USD School of Medicine
- 2005 The Protein Folding Scientific Advisory Committee Poster Award
“Inaugural Annual Symposium of Protein Folding Disorders”
Cambridge Healthtech Institute
- 2007 AHA Established Investigator Award
American Heart Association National Center
- 2008 President Award for Excellence in Research of the Established Faculty
University of South Dakota
- 2010 Fellow
AHA Council on Basic Cardiovascular Sciences
- 2011 Fellow
American Physiological Society: Cardiovascular Section
- 2012 Distinguished Service Award
The Academy of Cardiovascular Research Excellence (ACRE)
- 2019 Fellow (FISHR)
International Society of Heart Research (ISHR)
- 2020 Professor of the Game
University of South Dakota (USD Men’s basketball game vs. Purdue Fort Wayne,
1/23/2020)

MEMBERSHIPS AND OFFICES IN PROFESSIONAL SOCIETIES

Membership

- 2000- American Heart Association (AHA)

- 2000- International Society for Heart Research (ISHR): North American Section
- 2003- Academy of Cardiovascular Research Excellency (ACRE)/Chinese American Heart Association (CNAHA), lifetime member
- 2005- American Physiological Society (APS)

Offices

- 2008-2010 Board Director, ACRE
- 2010 Chair, Nomination/Election Committee, ACRE
- 2010-2014 Research Committee of AHA Midwest Affiliate
- 2010-2014 The Committee for Melvin L. Marcus Young Investigator Award in Cardiovascular Sciences
- 2014 The Committee for the Junior Young Investigator Award, ISHR-North American Section
- 2013-2015 Nominating Committee of APS Cardiovascular Section
- 2013-present Scientific Advisory Board of the International Academy of Cardiology
- 2016-present Awards Committee, APS Cardiovascular Section
- 2019- Chair, Awards Committee, APS Cardiovascular Section
- 2019- Steering Committee, APS Cardiovascular Section
- 2020- President-elect, Midland Society of Physiological Sciences (a chapter of APS)
- 2021- President-elect, ACRE

CONSULTING POSITIONS**A. PEER REVIER**

Acta Pharmaceutica Sinica B
 American Journal of Cardiology
 American Journal of Pathology
 American Journal of Physiology: Cellular Physiology
 American Journal of Physiology: Heart and Circulatory Physiology
 Antioxidants & Redox Signaling
 Archives of Biochemistry and Biophysics
 Autophagy
 BBA-Gene Regulatory Mechanisms

BBA-Molecular Basis of Disease
BBA-Molecular Cell Research
BBRC
BioMed Central-Cardiovascular Disorders
Biomolecules
Brain Research Bulletin
Cardiovascular Research
Cardiovascular Toxicology
Cell Physiology & Biochemistry
Cell Motility and Cytoskeleton
Circulation
Circulation Research
Circulation: Heart Failure
Clinica Chemica Acta
Comprehensive Physiology
Coronary Artery Disease
Current Molecular Medicine
Developmental Biology
EBiomedicine
eLife
EMBO Molecular Medicine
European Journal of Heart Failure
European Pharmacology Research
Experimental Cell Research
Free Radical Biology and Medicine
Frontiers in Cell and Developmental Biology
Frontiers in Physiology
Frontiers in Public Health
Growth Hormone and IGF Research
Human Molecular Genetics
Hypertension
International Journal of Medicine
International Journal of Molecular Medicine
International Journal of Molecular Science
International Journal of Obesity
International Journal of Nanomedicine
Journal of American College of Cardiology
Journal of American Heart Association
Journal of Cardiac Failure
Journal of Cardiovascular Pharmacology
Journal of Cell Physiology
Journal of Cell Science

Journal of Cellular and Molecular Medicine
 Journal of Clinical Investigation
 JCI-Insight
 Journal of Investigative Medicine
 Journal of Molecular and Cellular Cardiology
 Journal of Molecular and Cellular Biology
 Journal of Physiology
 Journal of Translational Medicine
 Journal of Vascular Research
 Journal of Visualized Experiments
 Medical Hypotheses
 Molecular Cellular Biochemistry
 Molecular and Cellular Biology
 Molecular Medicine
 Molecular Therapy Nucleic Acids
 Nature Communication
 Nature Review of Cardiology
 Pharmacology Research
 Physiology Reports
 PLoS Genetics
 PLoS One
 Protein and Cell
 Redox Biology
 Scientific Reports
 Trends in Cardiovascular Medicine
 Trends in Molecular Medicine

B. EDITORIAL BOARDS

2008-present	<i>International Journal of Physiology, Pathophysiology and Pharmacology</i>
2009-present	<i>American Journal of Translational Research</i>
2009-2010	Guest Editor for a Spotlight Issue on “The Ubiquitin-Proteasome Pathway in Cardiovascular Disease” for <i>Cardiovascular Research</i>
2011-present	Associate Editor-in-Chief, <i>American Journal of Cardiovascular Disease</i>
2012-present	<i>American Journal of Physiology- Heart & Circulatory Physiology</i>
2012-present	Review Editor for <i>Frontiers in Clinical and Translational Physiology</i>
2013-2019	Review Editor for <i>Frontiers in Striated Muscle Physiology</i>

2013-2017	<i>Circulation Research</i>
2019 – present	Associate Editor for <i>Frontiers in Striated Muscle Physiology</i>
2020 – present	<i>Journal of Molecular and Cellular Cardiology</i>
2020 – present	Associate Editor for <i>Frontiers in Ageing - Aging, Metabolism and Redox Biology</i>
2020 -	Co-Guest Editor for <i>Cells</i> Special Issue on “Molecular Mechanisms Underlying Cardiac Dysfunction”
2020 -	<i>Engineering</i>

C. NATIONAL/INTERNATIONAL COMMITTEES

2003-2020	International Expert Panel Singapore National Medical Research Council
2005-2007	External Expert Grant Reviewer Philip Morris External Research Program
2006	National Institutes of Health National Heart, Lung & Blood Institute; Ad hoc Reviewer for PPG
2007-2009	American Heart Association (AHA) National Center Cardiac Biology/Regulation Study Section II; Chartered Member
2009	AHA National Center Cardiac Biology/Regulation Study Section 2; Chairman
2007-present	China National Natural Science Foundation Key Research Projects; Oversea Expert Reviewer
2008-2009	National Institutes of Health Cardiac Contractility & Heart Failure Study Section; Ad hoc Member
2009-2013	National Institutes of Health Cardiac Contractility & Heart Failure Study Section; Chartered Member
2009-present	Abstract Review AHA Annual Scientific Sessions
2012-2015	AHA National Center Cardiac Biol. & Regulation – Basic Science Study Section; chartered member

- 2014-present Abstract Reviewer
AHA Basic Cardiovascular Sciences Annual Scientific Session
- 2014 International Society for Heart Research North American Section
Junior Young Investigator Award; Judge
- 2015 -present Grant Reviewer
Israel Science Foundation, Israel
- 2015 Grant Reviewer
Fondazione Cariparo, Italy
- 2015 NIH Special Emphasis Panel
Cardiovascular and Respiratory Sciences Member Conflict Applications (ZRG1
CVRS-B 02)
- 2015 NIH Special Emphasis Panel
Cardiovascular and Respiratory Sciences Member Conflict Applications (ZRG1
CVRS-E (02) M)
- 2015 The Ministry of Science and Technology, China
The State Science and Technology Awards of China; Oversea Reviewer
- 2015 The Ministry of Education, China.
The Chang-Jiang Scholar Program; Oversea Reviewer
- 2015 International Society for Heart Research-North American Section (ISHR-NAS)
2015 Annual Meeting (Seattle, WA) Poster Awards; Judge
- 2016 NIH National Heart Lung and Blood Institute (NIH/NHLBI)
Program Project Grant Peer Review Panel (2016/05 HLBP 1); Ad hoc Reviewer
- 2017 NIH National Heart Lung and Blood Institute (NIH/NHLBI)
Program Project Review Committee (HLBP 1 Workgroup 005, 2017/05
HLBP1); Ad hoc Reviewer
- 2017 - 2018 AHA National Center
Cardiac Biol. & Regulation – Basic Science Study Section; chartered member
- 2018 NIH Center for Scientific Review ZRG1 F05-U (20)
Fellowships: Cell Biology, Developmental Biology, and Bioengineering

- 2018 - UK Medical Research Council, Ad Hoc Reviewer
- 2018 The Netherlands Organization for Scientific Research (NWO/ZonMw), Ad Hoc Reviewer
- 2020 - AHA National Center Career Development Awards Cardiac Basic Sciences 2 committee, chartered member
- 2020 National Institutes of Health
Myocardial Ischemia and Metabolism (MIM) Study Section; *Ad hoc* Member (2020 June meetings)
- 2020 National Institutes of Health
Special Emphasis Panel for the dissolving MIM: ZRG1 MIM-R 01 (meeting date October 29-30, 2020)
- 2020- International Expert Panel
Singapore National Medical Research Council
- 2020 Judge Panel, Young Investigator Awards
Great Wall International Congress of Cardiology 2020 (GW-ICC 2020 virtual)/Asian Heart Society Congress 2020, October 18, 2020
- 2021 National Institutes of Health
Special Emphasis Panel, Fellowship: Cell Biology, Developmental Biology, and Bioengineering (F05-U) (meeting date Feb 18-19, 2021)

COMMITTEE ASSIGNMENTS

University of South Dakota (USD) Sanford School of Medicine

- 2001-2003 Coordinator, Cardiovascular Research Institute Seminar and Journal Club Series
- 2002-2004 Coordinator for Graduate Studies, Cardiovascular Research Institute
- 2002-2004 Director, Molecular Biology Core of Cardiovascular Research Institute
- 2002-2005 Internal Advisory Committee for the Cardiovascular COBRE
- 2003-2009 Research Committee of School of Medicine
- 2005-2007 Chair, Research Committee of School of Medicine
- 2004-2007 Graduate Committee of USD School of Medicine
- 2005-2007 University Senate
- 2005-2006 Conference of the Senate
- 2005 The Task Force of the USD Senate on USD Conflict of Interest Policy Draft
- 2005-2008 Medical Student Research Committee
- 2006 The Task Force for the Creation of Medical Student Scholarship Pathway
- 2006-2016 Chair, the MD/PhD Admissions Committee
- 2007 Pre-Tenure Review Committee for Dr. Yifan Li

- 2007 Pre-Tenure Review Committee for Dr. Alexander Erkine
- 2007 Chair, the Taskforce to Establishing a Graduate Course on *Grant and Scientific Writing*, for Basic Biomedical Sciences Graduate Program
- 2007 Convener, the Molecular Pathogenesis group SWOT (strength, weakness, opportunities, and threat) Analysis for Division of Basic Biomedical Sciences Research Retreat 2007.
- 2007-2014 Research Council of the Division of Basic Biomedical Sciences
- 2007-2008 Chair, the Search Committee for a tenure-track faculty position in Protein Quality Control (PQC) of the Division of Basic Biomedical Sciences
- 2008-2009 The Search Committee for a tenure-track faculty position in Protein Quality Control (PQC) of the Division of Basic Biomedical Sciences
- 2008 USD Committee investigating alleged research misconduct of a faculty member
- 2009-2018 Sanford School of Medicine P&T Committee
- 2009-2014 Chair, Monthly PQC Roundtable Meeting
- 2009-2010 LCME Task Force Sub-Committee
- 2010-2011 Chair, the Organizing Committee for the Inaugural Symposium on Ubiquitin, Protein Quality Control and Molecular Pathogenesis, Vermillion, SD, June 22-24, 2011.
- 2010-2011 The Search Committee for a tenure-track faculty position in Protein Quality Control and Degradation (PQCD)
- 2010-2016 Director, the Interim Center for PQCD Research
- 2009-2016 Chair, the Finance/Budget Committee for PQCD Research & Development
- 2011-2012 Task Force for Medical Curriculum Reform (Skin & Musculoskeletal Block)
- 2011 The Search Committee for VP of Health Affairs/Dean of Sanford School of Medicine
- 2011 Pre-tenure Review Committee of Dr. Victor Huber
- 2012 Chair, The Organizing Committee for the Second Symposium on Ubiquitin, Protein Quality Control and Molecular Pathogenesis, Deadwood, SD, June 13-15, 2012
- 2012 Pre-tenure Review Committee of Dr. Hongmin Wang
- 2013 The Primary Committee for Dr. Carlos Telleria's Promotion to Full Professor
- 2013 Pre-tenure Review Committee of Dr. Khosrow Rezvani
- 2013 The Search Committee for two tenure-track faculty positions in the Department of Biomedical Engineering

2014	Chair, The Organizing Committee for the Third Symposium on Ubiquitin, Protein Quality Control and Molecular Pathogenesis, Deadwood, SD, June 4-6, 2014
2014-2019	Grant Applications Pre-submission Internal Review Committee of Division of Basic Biomedical Sciences
2014	Pre-tenure Review Committee of Dr. J Scott Pattison
2016-present	Graduate Committee of USD Sanford School of Medicine
2016-present	The MD/PhD Admissions Standing Committee
2018	Member of The Search Committee for a tenure-track Assistant or Associate Professor in infectious disease for the Division of Basic Biomedical Sciences
2018	Member of The Primary Committee for Dr. S. N. Sathyanesan's Promotion to Full Professor
2019	Member of The Search Committee for a tenure-track Assistant Professor for the Division of Basic Biomedical Sciences
2019	Chair, The Primary Committee for Dr. H. Wang's Promotion to Full Professor.
2021-	P&T Committee, Sanford School of Medicine, USD.

COMMUNITY SERVICE

2002-2006	Judge, Sioux Valley Hospital & Health System Annual Quality Fair
2003	Volunteer Speaker, AHA Sioux Falls Regional Gala
2006 – 2017	Faculty Advisor, USD Association of Chinese Students and Scholars
2015-	Food Server, USD Lee Medical Building Annual Welcome Table Christmas Dinner

TEACHING AND ADVISING

A. COURSES TAUGHT

1. ANAT 511	<i>Gross Anatomy</i>	USD School of Medicine and Health Sciences
2. ANAT 521	<i>Microanatomy</i>	USD School of Medicine and Health Sciences
3. BIOCHEM	<i>Medical Biochemistry</i>	USD Sanford School of Medicine
4. PHPH 792	<i>Genetic Approaches</i>	USD Graduate School (Course Director)
5. CPHD 740	<i>Protein Quality Control</i>	USD Graduate School (Course Director)
6. BIOC 798	<i>Heat Shock Proteins</i>	USD Graduate School
7. PHPH 728	<i>Signal Transduction</i>	USD Graduate School
8. CPHD 788	<i>Res Basic Biomed Sci</i>	USD Graduate School
9. CPHD 898	<i>Thesis Research</i>	USD Graduate School
10. CPHD 620	<i>Foundations of Cardiovasc. Sci.</i>	USD Graduate School (Course Director)
11. CPHD 792	<i>Heat Shock Proteins and Dis.</i>	USD Graduate School
12. IMC 601	<i>Skin and Musculoskeletal Block</i>	USD Sanford School of Medicine

B. GRADUATE STUDENTS MENTORING

1. Served as the Primary Mentor

Graduate student names	Degree obtained (pursued)	Dates	Current positions
Wei Huang (visiting student)	Ph.D.	10/2001 – 10/2002	Professor of Medicine, Nanjing Medical University, Nanjing, Jiangsu, China
Assangi R. K. Kumarapeli	Ph.D.	9/2002 - 8/2006	Associate Professor, Department of Pathology, University of Arkansas for Medical Sciences, Little Rock, AR
Mingxin Tang	M.S.	7/2003 – 8/2006	Research Scientist University of Hawaii, Honolulu, HI, PA
Jie Li	Ph.D.	9/2003 - 12/2008	Assistant Professor, Medical College of Georgia Augusta University, Augusta, GA, USA
Qingwen Zheng	Ph.D.	9/2005 – 12/2010	Physician, Kaiser Permanente Roseville Medical Center Roseville, California
Mark J. Ranek, BS	Ph.D.	9/2006 – 5/2012	Assistant Professor, Johns Hopkins University
Lei Zhang, MS	Ph.D. (co-mentor)	8/2015 – 8/2016	Postdoc at Medical University of South Carolina, Charleston, SC
Hanming Zhang, BS	(PhD)	8/2013 – 5/2019	Postdoc at Yale University, New Haven, Connecticut
Penglong Wu, BM, MS	Visiting PhD Student from Shanghai Jiao Tong University	6/22/2015 – 6/21/2017	Postdoctoral fellow, Guangzhou Medical University and USD Sanford School of Medicine.
Chao Suo, DDS	(PhD)	8/2015 – 5/2017	Taking a 2-year leave of absence for a Dentistry Licensure training program in University of Las Vegas, Las Vegas, NV.
Liuqing Yang, BS, MS	PhD	8/22/2017 -	
Samiksha Giri, BA	PhD	8/22/2017-	
Megan T. Lewno, BA	PhD	8/22/2019-	

Mingqi Cai, MBBS	Visiting Graduate Student	1/22/2020 - 1/21/2021	USD PhD student
Mingqi Cai, MBBS	PhD	1/06/2021 -	
MdSalim Ahammed, MS	PhD	1/06/2021-	

2. Served as a Member of the Graduate Study Advisory Committees

2001-2003	James Kuzman, PhD Advisory Committee, BBS
2005-2006	Leah Callahan, MS thesis committee, BBS
2008-2010	Lili Guo, PhD Advisory/Thesis Committee, BBS
2010-2011	Rui Du, PhD Advisory Committee, BBS
2010-2014	Shuai Li, PhD Advisory/Thesis Committee, BBS
2013-2016	Lei Zhang, PhD Advisory Committee, BBS
2013-2017	Xianhua Meng, MS/PhD Thesis Committees, Dept. of Chemistry, USD
2015-2018	Hongbo Gao, PhD Advisory/Thesis Committee, BBS

C. NON-CLASSROOM TEACHING/ADVISING

1. List of Junior Faculty Mentored

2008-2012	H. Su, PhD, Research Assistant Professor; currently R01-funded tenured Associate Professor at Medical College of Georgia of Augusta University, Augusta, Georgia
2009-2015	H. Wang, PhD, Assistant Professor; currently R01-funded tenured Associate Professor at University of South Dakota Sanford School of Medicine, Vermillion, SD 57069
2011-2016	K. Rezvani, PhD, Assistant Professor; currently Tenured Associate Professor at USD Sanford School of Medicine, Vermillion, SD
2011-2016	J.S. Pattison, PhD, Assistant Professor
2013 -2015	C. Wang, MD, PhD, Research Assistant Professor; currently Professor and Chairman of the Dept. of Pathophysiology, Wuhan University Medical School, Wuhan, Hubei, China
2017- 2019	Nirmal Parajuli, PhD, Research Assistant Professor, currently Senior Research Associate at the Immunology Research Program, Henry Ford Health System, Detroit, MI.
2019 - 2020	Xing Liu, PhD, Assistant Professor, Department of Biochemistry, Purdue University, West Lafayette, IN, (secondary mentor for AHA CDA grant, funded on July 1 st , 2020)

2021- William Chen, MD, PhD, Assistant Professor, Division of Basic Biomedical Sciences, USD Sanford School of Medicine.

2. List of Postdoctoral Trainees

Name of trainees	Training period	Current position
Quanhai Chen, MD	1/2003 - 11/2005	Senior Scientist, GlaxoSmithKline, Heart Failure (Development Performance Unit), King of Prussia, PA, USA
Hanqiao Zheng, MD, PhD	8/2003 - 5/2008	Res. Scientist, Harvard University School of Public Health, Boston, MA, USA
Jinbao Liu, MD, PhD	11/2003 - 10/2005	Professor and Vice President for Research, Guangzhou Medical University, Guangzhou, Guangdong, China
Huabo Su, PhD	11/2004 – 6/2008 7/2008 – 7/2012	Associate Professor (tenured), Vascular Biology Center and Department of Pharmacology, Medical College of Georgia Augusta University, Augusta, GA, USA
Daoxiong Lei, MD, PhD	11/2005-10/2008	Professor, Tianjin 4 th Hospital, Tianjin, China
Youn-Chul Ryu, PhD	9/2006 – 4/2009	Associate Professor, Jeju National University, Jeju-si, Jeju-do, Korea
Wei Huang, MD, PhD	10/2007-10/2008	Professor, Nanjing Medical University, Nanjing, Jiangsu, China
Jie Li, MD, PhD	1/2009 – 7/21/2012	Assistant Professor, Medical College of Georgia Augusta University, Augusta, GA, USA
Zongwen Tian, MD, PhD	8/2009 – 8/21/2012	Associate Professor & Chair, Dept. of Anatomy, Wuhan University Medical School, Wuhan, Hubei, China
Changhua Wang, MD, PhD	10/2010-10/2011	Professor & Chair, Dept. of Pathophysiology, Wuhan University Medical School, Wuhan, Hubei, China
Mark J. Ranek, PhD	6/1/2012-11/21/2012	Assistant Professor at Johns Hopkins University, Baltimore, MD
Hongxin Xu, MD, PhD	11/1/2012-11/21/2013	Associate Professor and Cardiologist, Renming Hospital, Wuhan University School of Medicine, Wuhan, Hubei, China
Chengjun Hu, MD, PhD	3/22/2012 – 3/31/2014	Associate Professor, Dept. of Anatomy, Wuhan University College of Basic Biomedical Sciences, Wuhan, Hubei, China
Yihao Tian, MD, PhD	3/22/2012 – 3/31/2014	Assistant Professor, Dept. of Anatomy, Wuhan University College of Basic Biomedical Sciences, Wuhan, Hubei, China
Erin J. Terpstra, PhD	2/1/2012 – 6/30/2015	Medical Student, USD SSOM

Bo Pan, PhD	12/22/2014 – 6/21/2019	Senior Postdoctoral Research Associate, Department of Physiology, Wayne State University College of Medicine, Detroit, MI
Peng Xiao, PhD	1/22/2015 – 9/21/2018	Postdoctoral Fellow, The Wistar Institute, Philadelphia, PA 19104
Ammara Abdullah, PhD	6/22/2015 – 6/21/2017	Research Scientist II, ONC ODD/Oncology NIBR, Novartis Pharmaceuticals, 3000 Kent Ave. Ste. #1950, West Lafayette, IN 47906
Penglong Wu, MD, PhD	8/3/2017- 6/21/2020	Physician Scientist, The Cardiovascular Institute of Xiamen University, Xiamen, Fujian, China
Mohamed Hussain, PhD	9/22/2020 - 6/21/2021	
Mark J. Bouska, PhD	1/3/2021 -	

3. List of Visiting Scientists

Name & Degrees	Training period	Parent Institution
Changhua Wang, MD, PhD	1/2014 ~ 8/21/2015	Professor & Chair, Dept. of Pathophysiology, Wuhan University College of Basic Medical Sciences, Wuhan, Hubei, China
Maggie Gong, MD, PhD	2/2014 ~ 1/21/2015	Professor, Harbin Medical University, Harbin, Heilongjiang, China
Feng Yao, MD, PhD	4/2009- 11/2009	Professor & Chair, Dept. of Breast Surgery, Wuhan University Medical College, Wuhan, Hubei, China

4. Medical Students and Medical Resident Research

2002	Matt Mahowald, USD medical student of Class 2005, summer research
2002	Louis W. Lim, MD, Internal Medicine Chief Resident, research rotation
2004	Paul King, USD medical student of Class 2007, summer research
2005	Weitian Liu, MD, Internal Medicine Resident, research rotation
2007	Marius Vulcan, USD medical student of Class 2010, summer research
2011, 2012	Sigurd E. Hartnett, USD MD/PhD program student, summer research
2015	Nickolas Pekas, USD MD/PhD student, summer research
9/2021 -	Andrew L. Guymon, USD medical student of Class 2025, research volunteer.

5. Undergraduate Student Researchers

Name of trainees	School attended	Distinction	Training period
Mark List	Augustana College, Sioux Falls, SD	BRIN Summer Scholars	2005 & 2006 summer

Mark List	Augustana College	Part-time undergrad researcher	1/2005-12/2006
Lindsey Gerdes	Augustana College,	COBRE Summer Research Scholar	2005 summer
Andy Nelson	Augustana College	Undergrad Summer Researcher	2006 summer
Heath Eggleston	Dakota Wesley	BRIN Summer Scholar	2007 summer
Blake Alberts	USD	The Honors' Thesis Research	2008-2010
Blake Alberts	USD	NIH summer research scholarship	2009 summer
Morgan Hanson	USD	The Honors Thesis Research	2008-2010
Levi Froke	USD	NIH Summer Research Scholarship	2009 summer
Levi Froke	USD	Part-time undergrad. researcher and The Honors Thesis Research	2009-2011
Yun Zou	USD	Part-time undergrad. researcher	12/2009-5/2010
Michael Freitag	USD	Part-time undergrad. researcher	1/2010-4/2010
Michael Freitag	USD	Undergrad summer researcher	2010 summer
Jiwen Li	Rice University	Rice/Baylor Medical Scholar, NIH Summer Research Scholarship	2010 summer
Michael Freitag	USD	Undergrad summer researcher	2011 summer
Levi Froke	USD	Undergrad summer researcher	2011 summer

Lance M. Ranek	USD	Work study (undergrad research assist)	2012/2013 school year
Lance M. Ranek	USD	Work study (undergrad research assist)	2013/2014 school year
Lance M. Ranek	USD	Undergrad summer researcher	2014 summer
Casey A. Reihe	USD	Undergrad researcher	2015 Spring – 2017 Spring
Caleb Ray Wenz	USD	Undergrad researcher	2015 Spring-2016 Spring
Andrew V.Y. Yevugah	USD	Undergrad researcher	2016 Spring- 2017 Spring
Tanner James Redlin	USD	Undergrad researcher	2016 Fall-2017 Spring
Taylor Grace Faw	USD	Undergrad researcher	2016 Fall-2018 Summer
Kasha Merie Shear	USD	Undergrad researcher	2016 Fall-2018 Spring

6. Undergraduate Honors Thesis Advised

2008-2010 Blake Alberts, USD
 2008-2010 Morgan Hanson, USD
 2009-2011 Levi Froke, USD
 2014-2017 Casey A. Reihe, USD
 2017-2018 Taylor Grace Faw, USD
 2019-2019 Mary Ann Doom, USD

7. Research Technicians Mentored

2002-2003 Niels Harden, currently practicing physician
 2003-2005 Joseph W. Glasford, currently Research Operation Manager of Sanford Research/USD
 2004-2008 Kathleen M. Horak, currently homemaker
 2006-2008 Mingxin Tang, currently Director of Physiology Core of University of Hawaii, Honolulu, Hawaii, USA
 2007-2008 Amy J. Stephenson
 2007-2008 John R. Bosch, went to Osteopathic medical school at St Paul, MN
 2008-2010 Emily McDowell, currently Research Associate in USDSSOM
 2011-2012 Travis Bjordahl, went to dental school
 2009-2014 Suleman Said, Research Associate III
 2008-2015 Andrea Jahn, 2008-11/8/2015, Research Associate III; currently Assistant to the Dean of BBS.
 2016-2019 Megan T. Lewno, BS, Research Associate I and II.
 2018- Jack O. Sternberg, BS, Research Associate I (7/2018-6/2020) and II (7/2020-).

2020 - Jose Lira, BS, Research Associate I (8/21/2020 -)
 2021 - Renae Sieck, MS, Research Associate I (7/26/21 -)

D. DEVELOPMENT OF TEACHING STRATEGIES, ASSESSMENTS, METHODS

2004-2005 Developed and directed a graduate course “Genetic Approaches”
 2007 Chair, the Working Group on establishing a graduate course on Grant and Scientific Writing, USD Sanford School of Medicine Division of Basic Biomedical Sciences (2007).
 2011 Member, Medical Curriculum Innovation (Skin and musculoskeletal system section)
 2012 Developed and directed a graduate course “CPHD 740: Protein Quality Control and Degradation (PQCD)”
 2013 Chair, Taskforce to develop the curriculum for the Cardiovascular Sciences Specialty of the Basic Biomedical Sciences graduate program
 2014 Developed and directed a graduate course “CPHD 620: Foundation of Cardiovascular Sciences”

PRESENTATIONS

Invited Seminars/Presentations (National and International)

1. “Cardiac myocyte remodeling in pressure overloaded cardiac hypertrophy and failure”; The Institute of Muscle, Arthritis, and Skin Diseases, NIH, Bethesda, MD. April, 1998.
2. “Cardiac myocyte remodeling in chronic pressure overload-induced cardiac hypertrophy and failure”; Gladstone Cardiovascular Institute, University of California at San Francisco, CA, May, 1998.
3. “The alteration of intercalated disk-associated proteins during the progression from compensated cardiac hypertrophy to congestive heart failure in pressure overloaded guinea pigs”; Department of Pharmacology in University of Minnesota, Minneapolis, MN. March, 1998.
4. “Intercalated disk remodeling in pressure overloaded cardiac hypertrophy and failure”; Division of Molecular Cardiovascular Biology, Children’s Hospital Research Foundation, Cincinnati, OH. April, 1998.
5. “Cytoskeletal remodeling of cardiac myocytes in pressure overload hypertrophy and failure”; Department of Pharmacology, East Tennessee State University School of Medicine, Johnson City, TN, July, 1999.
6. "*In vivo* Modeling Desmin-related Cardiomyopathies with Transgenics"; the 4th Annual Scientific Meeting of Heart Failure Society of America, Boca Raton, FL, September 10-13, 2000.
7. "Transgenic Models of Desmin-related Cardiomyopathies"; South Dakota Health Research foundation-Cardiovascular Research Institute, University of South Dakota,

Sioux Falls, SD, October 16, 2000.

8. “Desmin filaments and heart diseases”; Department of Biomedical Sciences, Florida Atlantic University, Boca Raton, FL, May 21, 2001.
9. “Intermediate Filaments and Cardiac Diseases: Cause and Effects”; Division of Cardiology, University of California at Davis, Davis, CA, May 25, 2001.
10. “Intermediate Filaments and Cardiac Diseases: Cause and Effects”; Department of Basic Biomedical Sciences, Mercer University Medical School, Macon, GA, June 14, 2001.
11. “Intermediate Filaments and Cardiac Diseases: Cause and Effects”; Center of Excellence in Genomics and Bioinformatics, University of Tennessee, Memphis, TN, June 25, 2001.
12. “Intermediate filaments and cardiac diseases: Cause and Effects”; Midwestern University, Glendale, AZ, June 29, 2001.
13. “Desmin in cardiac remodeling”; A Symposium on “Remodeling and Progression of Heart Failure” (an official satellite meeting of the 17th World Congress of International Society for Heart Research), Minneapolis, MN, July 12-15, 2001.
14. “Desmin filaments and cardiac diseases: cause and effects”; University of South Dakota School of Medicine, Vermillion, SD, July 16, 2001.
15. “Intermediate filaments and cardiac diseases: cause and effects”; Department of Physiology, University of Texas Health Science Center, San Antonio, TX, July 23, 2001.
16. “Ubiquitin-Proteasome System in Pathogenesis and Therapeutics”; Wuhan University College of Basic Medical Sciences, Wuhan, Hubei, China, September 20, 2002.
17. “Ubiquitin-Proteasome System and Cardiovascular Diseases”; South Dakota State University College of Veterinary Science, Brookings, SD, October 4, 2002.
18. “Intermediate filaments and cardiac disease: establish causality”; Wuhan University College of Medicine, Wuhan, Hubei, China, Oct. 29, 2002.
19. “Ubiquitin-Proteasome System in Pathogenesis and Therapeutics”; Guangzhou Medical College, Guangzhou, Guangdong, China, Nov. 1, 2002.
20. “Desmin filaments and cardiac diseases”; the VII Meeting of International Society for Heart Research: China Section, Guangzhou, China, November 1-5, 2002.
21. “Modulation of the ubiquitin-proteasome system by an alpha B-crystallin mutant”; The 1st Annual COBRE Symposium. Rapid city, SD, June 28-30, 2003.
22. “In Situ Monitoring Dynamic Changes in the Ubiquitin-Proteasome System in vitro and in vivo”; The 1st Symposium of the Academy of Cardiovascular Research Excellency (ACRE), Washington, DC, April 20, 2004.

23. “Trashmen and police on strike in Alzheimer’s disease of the heart”; The 2nd Annual COBRE Symposium. West Yellow Stone, MT, August 4-7, 2004.
24. “Trashmen on strike in a mouse model of cardiac Alzheimer’s”; Division of Cardiology, University of Utah, Salt Lake City, UT, November 2, 2004.
25. “A novel transgenic mouse model reveals deregulation of the ubiquitin-proteasome system in the heart by Doxorubicin”; The 2nd ACRE annual scientific meeting, Vancouver, Canada, July 15, 2005.
26. “The Ubiquitin-Proteasome System in Cardiac Remodeling and Failure”; Cardiovascular Distinguished Lecture Series, University of California at Los Angeles School of Medicine, LA, CA January 31, 2006.
27. “Dissecting the Ubiquitin-Proteasome System in the Heart with Genetic Approaches”; Long Island Jewish Medical Center, New Hyde Park, NY, March 7, 2006.
28. “The Ubiquitin-Proteasome System in Cardiac Remodeling and Failure” at the Department of Molecular and Cellular Pharmacology, University of Miami, Miami, FL, March 16, 2006.
29. “The Ubiquitin-Proteasome System in Cardiac Physiology and Pathophysiology”; The 5th International Ascona Workshop on Cardiomyocyte Cell Biology, Monte Verita, Ascona, Switzerland, April 2-6, 2006.
30. “Inadequate Protein Quality Control in Heart Failure”; Division of Molecular Medicine, UCLA School of Medicine, Los Angeles, CA, April 18, 2006.
31. “The Ubiquitin-Proteasome System in Cardiac Remodeling and Failure”; Department of Molecular Genetics, University of Cincinnati, Cincinnati, OH May 2, 2006.
32. “Inadequate Protein Quality Control in Heart Failure”. The Center for Translational Medicine, Jefferson Medical College, Philadelphia, PA, May 10, 2006.
33. “The Ubiquitin-Proteasome System in Cardiac Remodeling and Failure”; The Center of Cardiovascular Sciences, Albany Medical College, Albany, NY, May 12, 2006.
34. “The Ubiquitin-Proteasome System in Cardiac Remodeling and Failure”; Department of Pharmacology, Loyola University Medical Center, Maywood, IL, May 15, 2006.
35. “The COP9 Signalosome and Protein Quality Control”, ZOMES IV: The 4th International Symposium on COP9 Signalosome, Proteasome, and eIF3: at the interface between signaling & proteolysis. New Haven, CT, June 18-21, 2006.
36. “The Ubiquitin-Proteasome System in Cardiac Remodeling and Failure”; University of Texas Houston Medical School, Houston, TX, May 11, 2007.
37. “Protein quality control in cardiac remodeling and failure”; Wuhan University College of Basic Biomedical Sciences, Wuhan, China, June 4, 2007.
38. “The ubiquitin proteasome system in cardiac remodeling and failure”; Marie Curie

- Symposium on the Ubiquitin-proteasome System in Cardiovascular Disease, Hamburg, Germany, June 9, 2007.
39. “Ubiquitin-proteasome system dysfunction in cardiomyopathies”; Heart Failure 2007 (the annual meeting of Heart Failure Association of the European Society of Cardiology), Hamburg, Germany, June 10, 2007.
 40. “Cardiac remodeling and protein quality control”; The 29th Meeting of the North American Section of the International Society for Heart Research (ISHR), Bologna, Italy, June 21-22, 2007.
 41. “Proteasomal degradation”; The 4th Annual Symposium of the American Heart Association Council on Basic Cardiovascular Sciences. Keystone, CO, USA, July 30-August 2, 2007.
 42. “The Proteasome and Cardiac Disease”; *Sunday Morning Program*, American Heart Association Scientific Sessions, Orlando, Florida, USA, November 5, 2007.
 43. “Proteasomal degradation in cardiomyopathy”; *Cardiac Seminar*, American Heart Association Scientific Sessions, Orlando, Florida, USA, November 7, 2007.
 44. “The ubiquitin-proteasome system for protein degradation”; *European Winter Meeting on Translational Cardiology*, organized by the Heart Failure Association (HFA) of the European Society of Cardiology (ESC), Garmisch-Partenkirchen, Germany, January 23-26, 2008.
 45. “Proteasomes in cardiac remodeling and failure”; Late Breaking Sciences-ISHR-North America Section 2008 meeting, Cincinnati, OH, June 17-21, 2008.
 46. “The role of the UPS in cardiac disease”; Sunday Morning Program Session on Protein Misfolding, Proteolysis, and Cardiac Disease, AHA Scientific Sessions, New Orleans, LA, November 8, 2008.
 47. “COP9 Signalosome, Proteasome, and Lysosome”; The Graduate Seminar Series, Guangzhou Medical College, Guangzhou, Guangdong, China, December 18, 2008
 48. “COP9 Signalosome, Proteasome, and Lysosome: All in the Same Zomes”; Wuhan University College of Basic Medical Sciences, Wuhan, Hubei, China, December 19, 2008.
 49. “Protein Quality Control in Cardiac Remodeling and Failure”; University of British Columbia, Vancouver, BC, Canada, April 24, 2009.
 50. “A molecular pathway underlying cardiac pathogenesis of inadequate PQC”; the Division of Basic Biomedical Sciences of Guangzhou Medical College, Guangzhou, Guangdong, China, June 19, 2009.
 51. “Proteasome, lysosome, and signalosome: all in the same ZOME”; Wuhan University College of Medicine, Wuhan, Hubei, China, June 24, 2009

52. “COP9 signalosome in the heart”; the Sunday Morning Program on “Protein Quality Control in Heart Disease”, AHA Scientific Sessions, Orlando, FL, November 15, 2009.
53. “Ubiquitin-proteasome system in heart disease”; the Graduate School of Guangzhou Medical College, Guangzhou, Guangdong, China, December 17, 2009.
54. “The ubiquitin-proteasome system in cardiac proteinopathy”; Department of Cellular Physiology and Neurosciences, Loyola University, Maywood, IL, April 20, 2010.
55. “Proteasome functional insufficiency in cardiac proteinopathy”; The session on “The Role of the Ubiquitin Proteasome System in Cardiac Disease, Diabetes, and Aging” of the 2010 EB meeting, Anaheim, CA, April 24-28, 2010.
56. “The ubiquitin-proteasome system in cardiac proteinopathy”; The Department of Physiology of University of Oklahoma College of Medicine, Oklahoma City, OK, May 10, 2010.
57. “COP9 Signalosomes regulate proteolysis in the heart”; a state-of-the-art lecture given to the Symposium "Genes, Proteins, and Translational Medicine" hosted by UCLA in conjunction with the AHA BCVS 2010 meeting, Rancho Mirage, CA, July 18, 2010.
58. “The ubiquitin-proteasome system in cardiac remodeling and failure”; Department of Biology, San Diego State University, San Diego, CA, November 4, 2010.
59. “The Ubiquitin-Proteasome System in Cardiac Remodeling and Failure”; Department of Molecular and Integrative Physiology University of Illinois at Urbana-Champaign, Urbana, IL, April 28, 2011.
60. “Protein Quality Control and Heart Disease”; Molecular Biology and Biotechnology Seminar Series, The Center of Molecular Biology and Biotechnology and College of Medicine, Florida Atlantic University, Boca Raton, FL, November 16, 2011.
61. “The COP9 Signalosome Regulates Autophagy”; Sanford Research/USD, Sioux Falls, SD, November 18, 2011.
62. “Proteasome Functional Insufficiency in Cardiac Pathogenesis”; Lillehei Heart Institute Lecture, Lillehei Heart Institute at the University of Minnesota Medical School, Minneapolis, MN, December 7, 2011.
63. “Can we treat proteinopathy by upregulating 11S proteasomes?” A State-of-the-Art Lecture at the Conference on the Protein Degradation Pathways in Health and Diseases , San Diego, CA., January 23, 2012.
64. “The ubiquitin-proteasome system in cardiac pathogenesis”; Department of Pathology, University of Cincinnati, Cincinnati, OH, May 18, 2012.
65. “Proteasome functional insufficiency in cardiac pathogenesis”; Department of Cell Biology and Molecular Medicine, UMDNJ – New Jersey Medical School, Newark, NJ, May 23, 2012.

66. “The ubiquitin-proteasome system in heart disease”; Department of Biomedical Sciences, New York College of Osteopathic Medicine at New York Institute of Technology, Old Westbury, NY, May 25, 2012.
67. “Protein Quality Control and Disease: Focus on the COP9 Signalosome”; the Educational Ministry Key Laboratory Seminar Series, Shanghai Jiaotong University, Shanghai, China. June 20, 2012.
68. “Intracellular Protein Quality Control and Pathogenesis: the COP9 signalosome”; Pathophysiology Invited Seminars of Wuhan University, Wuhan, Hubei, China. June 25, 2012
69. “Ubiquitination and Proteasomes: Mechanism of Heart Failure”; The 2012 Scientific Session of American Heart Association Council on Basic Cardiovascular Sciences (AHA-BCVS): Frontiers in Cardiovascular Science and Novel Therapy. New Orleans, LA, July 24, 2012.
70. “The COP9 Signalosome Polices the Heart”; Department of Biological Sciences, University of Illinois at Chicago, Chicago, IL, September 18, 2012.
71. “Proteasome Dysfunction in Cardiac Pathogenesis”; the Feinberg Cardiovascular Research Institute, Northwestern University, Chicago, IL, September 19, 2012.
72. “Proteasome Dysfunction in Pathogenesis”; the Biochemistry and Molecular Biology Seminar at Mayo Clinic in Rochester, Minnesota, October 2, 2012
73. “Protein Quality Control and Pathogenesis”; *University of Wisconsin Department of Pathology and Laboratory Medicine Seminar*, Madison, Wisconsin, October 10, 2012.
74. “UPS regulation and dysfunction in heart failure”, an invited lecture to a Session on Protein Quality Control and Homeostasis in Cardiac Physiology and Disease, AHA Scientific Sessions, Los Angeles, CA, November 4, 2012.
75. “Proteasome dysfunction in cardiac pathogenesis”; Keystone Symposium on Cardiac Remodeling, Signaling, Matrix and Heart Function (D4-2013), Snowbird, UT, April 9, 2013.
76. “Interaction of ubiquitin proteasome system and autophagy in the heart”; Cardiac Seminars on Dynamics of Protein Degradation Machinery in Cardiac Function. AHA Scientific Sessions, Dallas, TX, November 18, 2013.
77. “Loss of Function of an Extraproteasomal Ubiquitin Receptor Ubiquilin1 in Cardiomyocytes Exacerbates Cardiac Proteotoxicity”; The 35th International Society for Heart Research North American Section Meeting, Miami Beach, Florida, USA on May 12 -15, 2014.
78. “Protein Quality Control and Cell Death”; Department of Anatomy and Cell Biology, University of South Carolina School of Medicine, Columbia, SC, USA, May 19, 2014.
79. “Inadequate Coupling between Ubiquitination and Proteasomal Degradation in Cardiac

- Pathogenesis”; Wuhan University College of Basic Medical Sciences, Wuhan, Hubei, China, June 13, 2014.
80. “Protein Quality Control and Disease”; the 2nd Affiliated Hospital of Zhongnan University Xiangya Medical School, Changsha, Hunan, China on June 20, 2014.
 81. “Protein Quality Control and Degradation in Cardiac Disease”; the 2nd Affiliated Hospital of Zhejiang University Medical College, Hangzhou, Zhejiang, China, June 25, 2014.
 82. “Protein Quality Control in Cardiac Pathogenesis”; the Institute of Molecular Medicine of Peking University, Beijing, China, July 1, 2014.
 83. “Priming the proteasome by PKG: a novel cardioprotective mechanism of sildenafil”; The 19th World Congress on Heart Disease, Boston, MA, USA on July 28, 2014.
 84. “Inadequate Protein Quality Control in Cardiac Pathogenesis”; the Invited Seminars at Molecular Medicine, University of Oklahoma Health Science Center, Oklahoma City, OK, August 7, 2014.
 85. “Inadequate Protein Quality Control in Cardiac Pathogenesis”; the School of Veterinary Medicine and Biomedical Sciences, University of Nebraska, Lincoln, Nebraska, September 29, 2014
 86. “Protein degradation and heart failure: The NRF2-p62 axis in the cross-talk between proteasomal and lysosomal degradation”; the 36th International Society for Heart Research North American Section (ISHR-NAS) Annual Meeting, Seattle, WA, June 9, 2015.
 87. “Neddylolation/Deneddylolation, Protein Quantity & Quality Control, and Cardiomyocyte Necroptosis”; the Session on Protein Folding and ER Stress, AHA Scientific Sessions, Orlando, FL, November 9, 2015.
 88. “The COP9 Coerces Lysosomes and Proteasomes to Police the Heart”; Department of Biomedical Sciences, New York Institute of Technology College of Osteopathic Medicine, Old Westbury NY, March 14, 2016.
 89. “The Interplay between Autophagy and the Ubiquitin-Proteasome System in Cardiac Proteotoxicity”; American Society for Investigative Pathology (ASIP) 2016 Annual Meeting at Experimental Biology, San Diego, CA, April 2-6, 2016.
 90. “Inadequate protein quality control and heart failure”; The 4th International Conference on Cardio-metabolic Science, Wuhan, Hubei, China, May 11-14, 2016.
 91. “The COP9 signalosome in the heart”; Department of Pathophysiology at Guangzhou Medical University, Guangzhou, China, May 20, 2016
 92. “The COP9 coerces lysosomes and proteasomes to police the heart”; College of Life Science at Shanghai Ocean University, Shanghai, China, May 23, 2016

93. “Proteasome functional insufficiency in cardiac pathogenesis”, the Division of Cardiology at the 6th People’s Hospital of Shanghai, Shanghai, China, May 24, 2016
94. “Cardiac protein quality control and necroptosis”; Department of Pathology and Translational Pathobiology, LSU at Shreveport, Shreveport, LA, May 31, 2016.
95. “The COP9 Signalosome: A Posttranscriptional Cop in the Heart”; the 2016 Scientific Sessions of American Heart Association and the Council on Basic Cardiovascular Sciences, Phoenix, Arizona, July 18-21, 2016.
96. “Cardiac protein quality control in health and disease”; the ADVS/CIB Seminar Series of Utah State University at Logan, UT, September 22, 2016.
97. “Proteasome functional insufficiency in pathogenesis”; the Molecular and Cellular Pathology Seminar Series of the Graduate Program of the Department of Pathology at University of Alabama at Birmingham, Birmingham, AL, October 11, 2016.
98. “The State-Of-The-Art in Cardiac Protein Misfolding”; Cardiac Seminars on Misfolded Proteins of the 2016 Scientific Sessions of American Heart Association, New Orleans, Louisiana, November 15, 2016.
99. “Ubiquitin and ubiquitin-like proteins in the heart: an overview”; to the Symposium entitled “Ubiquitin and ubiquitin-like proteins in cardiovascular physiology and disease” of the Experimental Biology meeting, Chicago, IL, April 22-26, 2017.
100. “Protein Degradation in Heart Failure”; the 36th Annual Conference of the North American Section of the International Society of Heart Research (NAS-ISHR), New Orleans, LA, May 30-June 2, 2017.
101. “The COP9 Signalosome and the Heart”, Wuhan University College of Basic Medical Sciences, Wuhan, Hubei, China, June 17, 2017.
102. “Proteasome and Heart Failure”, Guangzhou Medical University College of Basic Medical Science and College of Pharmacy, Guangzhou, Guangdong, China, June 20, 2017.
103. “Proteasome Functional Insufficiency in Cardiac Pathogenesis”, Invited Seminar Series of the Department of Physiology, Peking University Health Science Center, Beijing, China, July 7, 2017.
104. “The Pathophysiological Significance of Cardiac Proteasome Functional Insufficiency (PFI)”, Department of Pharmacology, Harbin Medical University School of Pharmacy, Harbin, Heilongjiang, China, July 13, 2017.
105. “The Ubiquitin-Proteasome System in Cardiac Pathogenesis”, The 7th Cold Region Cardiology Conference (CRCC) and the 3rd China-Russia Jointed Pharmacology Conference, Harbin, Heilongjiang, China, July 13-16, 2017.

106. “Targeting protein quality control to treat heart disease”, Department of Pharmacology and Toxicology, Medical College of Georgia, Augusta University, Augusta, GA, Jan 22 2018.
107. “Pathophysiology of cardiac protein quality control”, The Seminar Series for Advancing Scientific Research at the No. 1 Affiliated Hospital of Guangxi University of Traditional Chinese Medicine, Nanning, Guangxi, China, August 14, 2018.
108. “Pathophysiology of Cardiac Protein Quality Control”, Nebraska Physiological Society Annual Scientific Meeting, Omaha, NE, October 20, 2018.
109. “Proteinopathies and Heart Disease”, the Session entitled: “Proteostasis Meets Protein Trafficking in the Heart” in the AHA Scientific Sessions 2018 to be held in Chicago IL November 12, 2018.
110. “Priming the Proteasome to Treat Cardiac Proteotoxicity”, Invited Seminar Series of the Department of Cell and Molecular Physiology, Stritch School of Medicine, Loyola University Chicago, Maywood IL, April 18, 2019.
111. “Duo-activation of PKG and PKA by PDE Inhibition to Treat Heart Disease with Increased Proteotoxic Stress”, Distinguished Lecturer Seminar Series of The Institute of Biosciences & Technology (IBT), the Texas A&M University College of Medicine, in Houston TX, May 6, 2019.
112. “Priming the proteasome to treat heart failure”, Distinguished Medical Lecturer Seminar Series, Wuhan University School of Basic Medical Sciences, Wuhan, Hubei, China, May 22, 2019
113. “Priming the proteasome to treat heart failure”, Lectures by Oversea Chinese Medical Elites Session 8, The 13th Oriental Congress of Cardiology (OCC 2019), Shanghai, China, June 2, 2019.
114. “Aberrant Protein Aggregation in Cardiac Muscle”, a panelist for a Visual Keystone Symposium (VKS) “[Intracellular Aggregates: Across the Spectrum of Health and Disease](https://virtual.keystonesymposia.org/ks/live/290/page/1580)”, July 18, 2019. (<https://virtual.keystonesymposia.org/ks/live/290/page/1580>)
115. “Dual Activation of PKA and PKG by PDE1 Inhibition Facilitates Proteasomal Degradation of Misfolded Proteins and Protects against Proteinopathy-Based HFpEF”, 2019 AHA BCVS Scientific Sessions –Session 1A, Boston, MA, July 29, 2019.
116. “Phosphoregulation of the Proteasome”, an invited lecture to the Cardiovascular Seminars Session entitled “Maintaining Protein Integrity Under Stress” of 2019 AHA Scientific Sessions (November 18, 2019; Philadelphia, PA).
117. “Harness proteasome phosphoregulation to protect against proteotoxicity”, Invited Seminar Series, Department of Surgery, The Ohio State University College of Medicine, Columbus, OH, December 10, 2019.

118. “Interplay between the ubiquitin-proteasome system and autophagy”, International Society for Heart Research (ISHR) (Quarantine) Cardiovascular Webinar Series. May 22, 2020. <https://www.youtube.com/watch?v=xiys1qoCMjI&t=45s>
119. “Crosswalk between proteasomal and lysosomal degradation”, 14th Oriental Congress of Cardiology (OCC2020 Virtual Conference), Channel-10 Session 6 WACC/CNAHA Session: Novel Technologies and Translational Medicine in the Diagnosis and Treatment of the Cardiovascular Diseases; June 1, 2020. <https://occ.1mice.net/live/play/144818>.
120. “Priming the proteasome to protect against proteotoxicity”, a Keynote Lecture to the 7th ACRE-APS Scientific Symposium (virtual), Aug. 8th, 2020.
121. “Activation of the proteasome by PKA protects the heart under stress”, an invited lecture to BCVS@GW-ICC: Kinase Signaling and Cardiac Injury, Great Wall International Congress of Cardiology 2020 (GW-ICC 2020 virtual)/Asian Heart Society Congress 2020, October 19, 2020.
122. “Catecholamine surges cause cardiomyocyte necroptosis via a RIPK1-RIPK3 dependent pathway”, an invited lecture to CNAHA@GW-ICC: Basic/Translational Cardiovascular Research and Novel Technologies, Great Wall International Congress of Cardiology 2020 (GW-ICC 2020 virtual)/Asian Heart Society Congress 2020, October 21, 2020.
123. “Priming the proteasome to protect against proteotoxic stress”, Virtual Seminar Series of the Center for Diagnostics and Therapeutics, Georgia State University, May 25, 2021.
124. “Cardiac UCHL1 protects against post-MI remodeling”, an invited lecture (virtual) to Session 1 of Chinese International Forum in the 15th Oriental Congress of Cardiology (OCC 2021): Translational Cardiovascular Research: from the bench-side to the bedside (virtual), May 29, 2021.
125. “Priming the proteasome to protect against proteotoxicity”, an invited lecture (in person) to the 40th ISHR-NAS Scientific Conference: “*Novel Mechanisms of Heart Failure: Advancing New Therapies*”. September 12-16th 2021 in Denver, Colorado.
126. “Priming the proteasome to treat heart failure”, an invited lecture (virtual) to BCVS-ACRE@GW-ICC: Molecular Mechanisms and Intervention of Cardiac Injury/Repair, Great Wall International Congress of Cardiology 2021 (GW-ICC 2021)/Asian Heart Society Congress 2021, October 28, 2021 (confirmed).
127. “The proteasome in cardiac health and disease”, an invited seminar to the Seminar Series of the McAllister Heart Institute at the University of North Carolina at Chapel Hill. November 10th, 2021 (scheduled).
128. “The proteasome hypothesis of heart failure”, an invited lecture (virtual) to the 13th Cross-Strait Cardiovascular Symposium and the 7th China Wine-City International Congress of Cardiology, Luzhou, Sichuan, China; November 11-19, 2021 (Scheduled).

Seminars Given in Employer Institutions

129. “Measurement of regional myocardial blood flow with unlabeled microspheres and Coulter Channelyzer” at University of South Dakota School of Medicine, Vermillion, SD. May 8, 1996.
130. “Dissecting desmin-related cardiomyopathy with mouse transgenesis”; Division of Pediatric Cardiology, Children’s Hospital, Cincinnati, OH, February 7, 2000.
131. “Ubiquitin-Proteasome System and Diseases”, Division of Basic Biomedical Sciences, University of South Dakota School of Medicine, Vermillion, SD, September 17, 2002.
132. “The COP9 Signalosome: A New Initiative from Plants to Mammalian Hearts”, Faculty Seminar, Division of Basic Biomedical Sciences, University South Dakota School of Medicine, Vermillion, SD September 27, 2005.
133. “Protein turnover, cardiomyopathy and the young”; Pediatric Grand Rounds, Department of Pediatrics, University of South Dakota School of Medicine, Sioux falls, SD, March 17, 2005.
134. “Targeted proteolysis in conformational disease”; Faculty Seminar Series, Division of Basic Biomedical Sciences, USD Sanford School of Medicine, October 28, 2008.
135. “Protein quality control and degradation in the heart”; The Inaugural Symposium on Ubiquitin, Protein Quality Control and Molecular Pathogenesis hosted by Sanford School of Medicine of University of South Dakota, Vermillion, SD, June 22-24, 2011.
136. “Histopathology in the post-genomics era”, Faculty Seminar Series of Division of Basic Biomedical Sciences, Sanford School of Medicine of University of South Dakota, Vermillion, SD 57069, March 16, 2012.
137. “Can we boost cardiac proteasomes by stimulating PKG?” The 2nd Annual PQCD Symposium, Deadwood, SD, USA. June 14, 2012.
138. “The Ubiquitin-Proteasome System in Cardiac Pathogenesis: Beyond the Proteasome”, Faculty Seminar Series, USD Sanford School of Medicine, November 30, 2017.
139. “Protein quality control and degradation in the heart”, Faculty Seminar Series, Department of Biomedical Engineering, University of South Dakota, Sioux Falls, SD, USA, January 18, 2018.
140. “Pathophysiological significance of priming the proteasome by PKA”, Faculty Seminar Series, USD Sanford School of Medicine, Vermillion, SD, September 17, 2020

INVITED MODERATOR (National/International)

- 2007 Co-chair, Sunday Morning Program-“Protein Conformation, Degradation and Cardiac Disease”, AHA Scientific Sessions 2007, Orlando, FL.

- 2008 Chair, the session on Genetic Models of Human Disease, AHA Scientific Sessions, Nov 8-10, 2008, New Orleans, LA.
- 2009 Co-Chair and invited speaker, Sunday Morning Program “Protein Quality Control in Heart Disease”, AHA Scientific Sessions, Orlando, FL Nov 14, 2009
- 2010 Co-Chair, the Symposium on "The Role of the Ubiquitin Proteasome System in Cardiac Disease, Diabetes, and Aging" for the Experimental Biology 2010 meeting. April 24-28 in Anaheim, CA.
- 2010 Co-Chair, Melvin L. Marcus Young Investigator Award in Cardiovascular Sciences. AHA Scientific Sessions, Nov. 15, 2010 in Chicago, IL
- 2014 Co-Chair, Session title: “Mitochondrial Biology and Protein Misfolding and/or Proteotoxicity” in the AHA BCVS 2014 "Pathways to Cardiovascular Therapeutics" conference July 14-17, 2014 in Las Vegas, Nevada.
- 2015 Co-Chair, Session Title: “Cellular Quality Control Mechanisms” in the AHA 2015 BCVS Scientific Sessions: Pathway to Cardiovascular Therapeutics. July 13-16, 2015 New Orleans, Louisiana.
- 2017 Co-Chair, a Symposium entitled: “Ubiquitin and Ubiquitin-Like Proteins in Cardiovascular Physiology and Disease” in 2017 Experimental Biology (EB) meeting, Chicago, IL, April 24, 2017.
- 2018 Co-Modulator, The Symposium 2 on Basic biomedical Science and Translational Medicine in the 11th Tongji Cardiovascular Disease Forum/2018China Precision Cardiology 2018/Central China International Congress of Cardiology, Wuhan, Hubei, China, August 11, 2018.
- 2018 Chair, the session entitled “Proteostasis Meets Protein Trafficking in the Heart” in AHA Scientific Sessions 2018, Chicago, IL November 12, 2018.
- 2019 Modulator, Concurrent Session 2B: Beyond Myocytes and Fibroblasts: Forgotten Cells of the Heart. BCVS 2019 Scientific Sessions - Integrative Approaches to Complex Cardiovascular Diseases, Boston, Massachusetts, July 29- Aug 1, 2019.
- 2020 Modulator, Modulator, Channel-10, Session 6 WACC/CNAHA Session: Novel Technologies and Translational Medicine in the Diagnosis and Treatment of the Cardiovascular Diseases (June 1, 2020). The 14th Oriental Congress of Cardiology (OCC2020 Virtual Conference), May 31-June 2, 2020.

BIBLIOGRAPHY

NCBI Bibliography Link:

<https://www.ncbi.nlm.nih.gov/myncbi/xuejun.wang.1/bibliography/public/>

A. ORIGINAL ARTICLES IN SCHOLARLY JOURNALS**(A) In Peer-Reviewed Chinese Journals**

1. Dong C, **Wang X**, Tu S, and Deng G. Alterations of plasma coagulation and extremity blood flow in erythralgia. *Natl Med J China* 1988; 68(5): 292-293.
2. **Wang X** and Dong C. A preliminary study on the relationship between epidemic erythralgia and El Niño. *Natl Med J China* 1988; 68(5): 266-268.
3. **Wang X**, Dong C, and Xiang J. Effects of ribavirin on coagulation-anticoagulation process in patients with epidemic hemorrhagic fever. *Natl Med J China* 1988; 68(12):699-701.
4. Dong C, **Wang X**, Xiang J, and Cosgraff T. The kinetic alterations of coagulation, anticoagulation and fibrinolytic system of patients with epidemic hemorrhagic fever and their significance. *Natl Med J China* 1988; 68(12): 678-681.
5. Dong C and **Wang X**. The alterations of coagulation, fibrinolysis, kinin, and complement system in epidemic hemorrhagic fever with DIC and its clinical Value. *Chinese J Pathophysiol* 1989; 5(5): 285-288.
6. Tu S, **Wang X**, Dong C, and Ling H. The significance of electrocardiogram on the estimation of myocardial infarction size induced by isoproterenol in rats. *Acta Academiae Medicinae Hubei* 1989; 10(4): 306-308.
7. **Wang X**, Dong C, and Xiang J. Effects of ribavirin on hemorrhagic tendency and fatality rate of patients with epidemic hemorrhagic fever. *Acta Academiae Medicinae Hubei* 1989; 10(3): 193-196.
8. **Wang X**, Dong C, and Ling H. The kinetic alterations of plasma prekallikrain and antithrombin 3 in rats following isoproterenol-induced myocardial infarction. *Chinese J Pathophysiol* 1990; 6(2): 100-103.
9. **Wang X**, Dong C, and Xiang J. Studies on the anion gap of epidemic hemorrhagic fever (I): The kinetic alterations of anion gap in EHF. *Acta Academiae Medicinae Hubei* 1990; 11(12): 285-289.
10. Dong C, **Wang X**, and Xiang J. Studies on anion gap of epidemic hemorrhagic fever (II): The mechanism and clinical significance of the increase in anion gap in EHF. *Acta Academiae Medicinae Hubei* 1990; 11(12): 289-293.

11. **Wang X**, Dong C, and Xiang J. Studies on anion gap of epidemic hemorrhagic fever (III): The mechanism and clinical significance of the decrease in anion gap in EHF. *Acta Academiae Medicinae Hubei* 1991; 12(1): 1-4.
12. Dong C, **Wang X**, and Xiang J. Studies on the anion gap of epidemic hemorrhagic fever (IV): The significance of simultaneous changes in anion gap, urine volume, and serum concentration of Na⁺ on the estimations of EHF prognosis. *Acta Academiae Medicinae Hubei* 1991; 12(1): 5-7.
13. **Wang X**, Dong C, and Ling H. Effects of Agkistrodon Halys on plasma coagulation in rats following isoproterenol-induced myocardial infarction. *Pace and Heart (Chinese)* 1991; 5(1): 29-31.
14. **Wang X**, Dong C, and Ling H. The kinetic alterations and pathophysiological significance of plasma coagulation in rats following isoproterenol-induced myocardial infarction. *Acta Academiae Medicinae Hubei* 1991; 12(4): 299-303.
15. **Wang X**, Ouyang J, Liu J, and Dong C. A pathogenesis of isoproterenol-induced occlusion of cardiac microvasculature in rats. *J Microcirculation* 1992; 2(1):8-11.
16. Yu X, Dong C, Ouyang J, Li D, and **Wang X**. An experimental study on the protective modification of soybean phospholipid liposomes enclosed superoxide dismutase (SOD) on the membrane of ischemic and reperfused myocardium in rats. *Chinese J Pathophysiol* 1993; 9(7): 804.
17. Dong C, Chen X, Wang C, Zhong Y, and **Wang X**. Experimental studies of the effects of soybean phospholipid liposomes against the myocardial membrane injury by ischemia/reperfusion. *Acta Academiae Medicinae Hubei* 1993; 14(4): 323-329.
18. **Wang X** and Dong C. Studies on the anion gap of epidemic hemorrhagic fever: V. Effects of intravenous ribavirin therapy on the changes of anion gap of patients with EHF. *Acta Academiae Medicinae Hubei* 1994; 15(3):247-249.
19. Dong C, Yu X, and **Wang X**. Myocardial membrane injury of myocardial ischemia and lipoideamia in rats. *Prog Biochem Biophys* 1994; 21:347-350.
20. **Wang X**, Dong C, Tu S, Zhang Y, Ouyang J, and Liu Y. Effects of soybean phospholipids liposomes on the left ventricular function and infarct size of ischemic-reperfused hearts in rabbits. *Chinese J Pathophysiol* 1994; 10(6):583-585.
21. Huang W, Ma W-Z, and **Wang X**. Intercalated disc remodeling in a transgenic mouse model of desmin-related cardiomyopathy. *Chin J Cardiol* 2003; 31(11):859-864.

(B) In Peer-Reviewed English Journals

22. **Wang X**, Li F, Said S, Capasso JM, and Gerdes AM. Measurement of regional myocardial blood flow in rats by unlabeled microspheres and Coulter Channelyzer. *Am J Physiol* 1996; 271:H1656-1665.
23. Li F, **Wang X**, Capasso JM, and Gerdes AM. Rapid transition of cardiac myocytes from hyperplasia to hypertrophy during postnatal development. *J Mol Cell Cardiol* 1996; 28: 1737-1746.
24. Gerdes AM, Onodera T, **Wang XJ**, McCune SA, and Capasso JM. Myocyte remodeling during the progression to failure in rats with hypertension. *Hypertension* 1996; 28:609-614.
25. Li F, **Wang X**, Bunger PC, and Gerdes AM. Formation of binucleated myocytes in rat heart: I. role of actin-myosin contractile ring. *J Mol Cell Cardiol* 1997; 29:1541-1551.
26. Li F, **Wang X**, and Gerdes AM. Formation of binucleated myocytes in rat heart: II. Cytoskeletal organization. *J Mol Cell Cardiol* 1997; 29:1553-1565.
27. **Wang X**, Li F, and Gerdes AM. Chronic pressure overload cardiac hypertrophy and failure in guinea pigs: I. Regional hemodynamics and myocyte remodeling. *J Mol Cell Cardiol* 1999; 31(2):307-317.(cover illustration)
28. **Wang X**, Li F, Campbell SE, and Gerdes AM. Chronic pressure overload cardiac hypertrophy and failure in guinea pigs: II. Cytoskeletal remodeling. *J Mol Cell Cardiol* 1999; 31(2): 318-331. (cover illustration)
29. **Wang X** and Gerdes AM. Chronic pressure overload cardiac hypertrophy and failure in guinea pigs: III. Intercalated disk remodeling. *J Mol Cell Cardiol* 1999; 31(2): 332-343. (cover illustration)
30. Milner DJ[§], Taffet GE,[§] **Wang X**[§], Pham T, Tamura T, Hartley C, Gerdes AM, and Capetanaki Y. The absence of desmin leads to cardiomyocyte hypertrophy and cardiac dilation with compromised systolic function. *J Mol Cell Cardiol* 1999; 31(11):2063-76.
31. Sanbe A, Nelson D, Gulick J, Setser E, Osinska H, **Wang X**, Hewett TE, Klevitsky R, Hayes E, Warshaw D, and Robbins J. In Vivo Analysis of an Essential Myosin Light Chain Mutations Linked to Familial Hypertrophic Cardiomyopathy. *Circ Res* 2000; 87: 296-302.
32. Yang Q, Hewett TE, Klevitsky R, Sanbe A, **Wang X**, and Robbins J. PKA dependent phosphorylation of myosin binding protein C in transgenic mice. *Cardiovasc Res* 2001; 51(1): 80-88.
33. **Wang X**, Osinska H, Dorn 2nd GW, Nieman M, Lorenz JN, Gerdes AM, Witt S, Kimball T, Gulick J, and Robbins J. Transgenic mouse model of desmin related cardiomyopathy. *Circulation* 2001; 103(19): 2402-2407.

34. **Wang X**, Osinska H, Klevitsky R, Gerdes AM, Nieman M, Lorenz JN, Hewett T, and Robbins J. Expression of R120G- α -B-crystallin causes aberrant desmin and α -B-crystallin aggregation and cardiomyopathy in mice. *Circ Res* 2001; 89(1): 84-91. (with companion editorial)
35. Wu G, Yussman MG, Barrett TJ, Hahn HS, Osinska H, Hilliard GM, **Wang X**, Toyokawa T, Yatani A, Lynch RA, Robbins J, and Dorn 2nd GW. Increased Myocardial Rab GTPase Expression. A Consequence and Cause of Cardiomyopathy. *Circ Res* 2001; 89:1130-1137.
36. Yi XP, **Wang X**, Gerdes AM, and Li F. Subcellular redistribution of focal adhesion kinase and its related nonkinase in hypertrophic myocardium. *Hypertension* 2003; 41: 1317-1323.
37. **Wang X***, Klevitsky R, Huang W, Glasford JW, Li F, and Robbins J. α B-Crystallin Modulates Protein Aggregation of Abnormal Desmin. *Circ Res* 2003; 93: 998-1005. (*corresponding author)
38. Dong X, Liu J, Zheng HQ, Glasford JW, Huang W, Chen QH, Harden NR, Li F, Gerdes AM, and **Wang X***. *In Situ* Dynamically Monitoring the Proteolytic Function of the Ubiquitin-Proteasome System in Cultured Cardiac Myocytes. *Am J Physiol Heart Circ Physiol* 2004; 287:H1417-H1425.
39. Yi XP, Zhou J, Baker J, **Wang X**, Gerdes AM, and Li F. Myocardial expression and redistribution of GRKs in hypertensive hypertrophy and failure. *Anatomic Record* 2005; 282A: 13-23.
40. Gard JJ, Yamada K, Green KG, Eloff BC, Rosenbaum DS, **Wang X**, Robbins J, Schuessler RB, Yamada KA, and Saffitz JE. Remodeling of gap junctions and slow conduction in a mouse model of desmin-related cardiomyopathy. *Cardiovasc Res* 2005; 67(3): 539-547.
41. Kumarapeli ARK, Horak KM, Glasford JW, Li J, Chen Q, Liu J, Zheng Q, and **Wang X***. A novel transgenic mouse model reveals deregulations of the ubiquitin-proteasome system in the heart by doxorubicin. *FASEB J* express article 10.1096/fj.05-3973. Published online October. 7, 2005.
42. Tang YD, Kuzman JA, Said S, Anderson BE, **Wang X**, and Gerdes AM. Low thyroid function leads to cardiac atrophy with chamber dilatation, impaired myocardial blood flow, loss of arterioles, and severe systolic dysfunction. *Circulation* 2005; 112: 3122-3130.
43. Chen Q, Liu J-B, Horak KM, Zheng H, Kumarapeli ARK, Li J, Li F, Gerdes AM, Wawrousek EF, and **Wang X***. Intrasarcolemmal amyloidosis impairs proteolytic function of proteasomes in cardiomyocytes by compromising substrate uptake. *Circ Res* 2005; 97: 1018-1026. (with companion editorial)
44. Liu J-B, Chen Q, Huang W, Horak KM, Zheng H, Mestril R, and **Wang X***. Impairment of the ubiquitin-proteasome system in desminopathy mouse hearts. *FASEB J* 2006; 20: 362-364.

45. Yi XP, Zhou J, Huber L, Qu J, **Wang X**, Gerdes AM, and Li F. Nuclear compartmentalization of FAK and FRNK in cardiac myocytes. *Am J Physiol Heart Circ Physiol* 2006; 290: H2509-H2515.
46. Liu J-B, Tang M, Mestril R, and **Wang X***. Aberrant protein aggregation is essential for a mutant desmin to impair the proteolytic function of the ubiquitin-proteasome system in cardiomyocytes. *J Mol Cell Cardiol* 2006; 40(4): 451-454. (with companion editorial)
47. Lindsten K, Menendez-Benito V, Masucci MG, Dantuma NP, Kumarapeli AR, Horak KM, Zheng H, and **Wang X***. GFP reporter mouse models of UPS proteolytic function. *FASEB J* 2006; 20(7):1027-1028.
48. Zhou J, Qu J, Yi XP, Graber K, Huber L, **Wang X**, Gerdes AM, and Li F. Up-regulation of gamma-catenin compensates for the loss of beta-catenin in adult cardiac myocytes. *Am J Physiol Heart Circ Physiol*. 2007; 292(1):H270-276.
49. Kobayashi S, Mao K, Zheng H, **Wang X**, Patterson C, O'Connell TD, and Liang Q. Diminished GATA4 protein levels contribute to hyperglycemia-induced cardiomyocyte injury. *J Biol Chem* 2007; 282:21945-52.
50. Qu J, Zhou J, Ping Yi X, Dong B, Zheng H, Miller LM, **Wang X**, Schneider MD, and Li F. Cardiac-specific haploinsufficiency of beta-catenin attenuates cardiac hypertrophy but enhances fetal gene expression in response to aortic constriction. *J Mol Cell Cardiol* 2007; 43:319-26. (cover illustration)
51. Powell SR, Samuel SM, Wang P, Divald A, Thirunavukkarasu M, Koneru S, **Wang X**, and Maulik N. Upregulation of myocardial 11S-activated proteasome in experimental hyperglycemia. *J Mol Cell Cardiol* 2008; 44(3):618-21.
52. Kumarapeli ARRK, Su H, Huang W, Horak MK, Tang M, Zheng H, Li M, and **Wang X***. Alpha B-Crystallin suppresses pressure overload cardiac hypertrophy. *Circ Res* 2008; 103(12): 1473-1482. (with editorial)
53. Liu J, Zheng H, Tang M, Ryu Y-C, and **Wang X***. A therapeutic dose of doxorubicin activate ubiquitin-proteasome system mediated proteolysis by acting on both ubiquitination apparatus and the proteasome. *Am J Physiol Heart Circ Physiol* 2008; 295 (6): H2541-2550.
54. Tydlacka S, Li S, **Wang X**, Li S, and Li X-J. Differential activities of the ubiquitin-proteasome system in neurons and glia may account for the preferential accumulation of misfolded proteins in neurons. *J Neuroscience* 2008; 28(49): 13285-13295.
55. Su H, Huang W, and **Wang X***. The COP9 signalosome negatively regulates proteasome proteolytic function and is essential to transcription. *Int J Biochem Cell Biol* 2009; 41(3):615-24.

56. Yang H, Zhou P, Huang H, Ma N, Shen S, Dong W, **Wang X**, Dou Q, Liu J. Shikonin exerts antitumor activity via proteasome inhibition and cell death induction in vitro and in vivo. *Int J Cancer* 2009; 124(10): 2450-2459. (cover illustration).
57. Kumarapeli AR, Horak KM, **Wang X***. Protein quality control in protection against systolic overload cardiomyopathy: the long term role of small heat shock proteins. *Am J Transl Res* 2010; 2(4):390-401.
58. Zheng H[§], Tang M[§], Zheng Q, Kumarapeli ARK, Horak KM, Tian Z, **Wang X***. Doxycycline Attenuates Protein Aggregation in Cardiomyocytes and Improves Survival of a Mouse Model of Cardiac Proteinopathy. *J Am Coll Cardiol* 2010; 56(17):1418-26. (with companion editorial)
59. Tang M[§], Huang W[§], Li J[§], Su H, Horak KM, Liang Q, Molkenin JD, and **Wang X***. Proteasome Functional Insufficiency Activates the Calcineurin-NFAT Pathway in Cardiomyocytes and Mouse hearts. *Cardiovasc Res* 2010; 88(3):424-33. (with companion editorial)
60. Lei D, Li F, Su H, Ye B, Wei N, and **Wang X***. COP9 Signalosome subunit 8 is essential to postnatal hepatocyte survival and effective proliferation. *Cell Death Diff* 2010; 18(2): 259-270.
61. Huang H, Zhang X, Li S, Liu N, Lian W, McDowell E, Zhou P, Zhao C, Guo H, Zhang C, Yang C, Wen G, Dong X, Lu L, Ma N, Dong W, Dou QP, **Wang X***, and Liu J*. Physiological levels of ATP negatively regulate proteasome function. *Cell Res* 2010; 20(12):1372-85.
62. Su H, Li J, Menon S, Liu J-B, Kumarapeli AR, Wei N, and **Wang X***. Perturbation of cullin deneddylation via conditional Csn8 ablation impairs the ubiquitin-proteasome system and causes cardiomyocyte necrosis and dilated cardiomyopathy in mice. *Circ Res* 2011; 108(1):40-50.
63. Li J, Powell SR, and **Wang X***. Enhancing proteasome function via overexpressing PA28 α protects against oxidative stress. *FASEB J* 2011; 25(3):883-893. PMID: 21098724
64. Lu L, Qin A, Huang H, Zhou P, Zhang C, Liu N, Li S, Wen G, Zhang C, Dong W, **Wang X**, Dou QP, and Liu J. Shikonin extracted from medicinal Chinese herbs exerts anti-inflammatory effect via proteasome inhibition. *Eur J Pharmacol* 2011; 658(2-3): 242-7.
65. Huang H, Liu N, Zhao K, Zhu C, Lu X, Li S, Lian W, Zhou P, Dong X, Zhao C, Guo H, Zhang C, Yang C, Wen G, Lu L, Li X, Guan L, Liu C, **Wang X**, Dou QP, and Liu J. Sanggenon C decreases tumor cell viability associated with proteasome inhibition. *Front Biosci (Elite Ed)*. 2011; 3:1315-25.
66. Zheng Q, Su H, Ranek MJ, and **Wang X***. Autophagy and p62 in cardiac proteinopathy. *Circ Res* 2011; 109 (3): 296-308.

67. Li J, Horak KM, Su H, Sanbe A, Robbins J, and **Wang X***. Enhancement of proteasomal function protects against cardiac proteinopathy and ischemia/reperfusion injury in mice. *J Clin Invest* 2011; 121(9):3689-700.
68. Zheng Q, Su H, Tian Z, **Wang X***. Proteasome malfunction activates macroautophagy in the heart. *Am J Cardiovasc Dis* 2011; 1(3):214-226.
69. Su H, Li F, Ranek MJ, Wei N, **Wang X***. The COP9 signalosome regulates autophagosome maturation. *Circulation* 2011; 124(19): 2117-2128.
70. Tian Z[§], Zheng H[§], Li J, Li YF, Su H, and **Wang X***. Genetically Induced Moderate Inhibition of the Proteasome in Cardiomyocytes Exacerbates Myocardial Ischemia-Reperfusion Injury in Mice. *Circ Res* 2012; 111(5): 532-542. (with companion editorial)
71. Huang H, Liu N, Guo H, Liao S, Li X, Yang C, Liu S, Song W, Liu C, Guan L, Li B, Xu L, Zhang C, **Wang X**, Dou QP, and Liu J. L-Carnitine Is an Endogenous HDAC Inhibitor Selectively Inhibiting Cancer Cell Growth In Vivo and In Vitro. *PLoS One* 2012; 7(11):e49062. doi: 10.1371/journal.pone.0049062.
72. Li X, Liu S, Huang H, Liu N, Zhao C, Liao S, Yang C, Liu Y, Zhao C, Li S, Lu X, Liu C, Guan L, Zhao K, Shi X, Song W, Zhou P, Dong X, Guo H, Wen G, Zhang C, Jiang L, Ma N, Li B, Wang S, Tan H, **Wang X**, Dou QP, and Liu J. Gambogic acid is a tissue-specific proteasome inhibitor in vitro and in vivo. *Cell Reports* 2013; 3(1): 211-22.
73. Rajagopalan V, Zhao M, Reddy S, Fajarado G, **Wang X**, Dewey S, Gomes AV, Bernstein D. Altered ubiquitin-proteasome signaling in right ventricular hypertrophy and failure. *Am J Physiol Heart Circ Physiol* 2013; 305(4):H551-62. doi: 10.1152/ajpheart.00771.2012.
74. Ranek MJ, Terpstra EJ, Li J, Kass DA, and **Wang X***. Protein kinase G positively regulates proteasome-mediated degradation of misfolded proteins. *Circulation* 2013; 128: 365-376. (with companion editorial).
75. Liu C, Guo L, Menon S, Jin D, Pick E, **Wang X**, Deng XW, and Ning W. COP9 signalosome subunit Csn8 is required to prevent premature G1 to S phase transition. *J Biol Chem* 2013; 288(28): 20443-52. doi: 10.1074/jbc.M113.468959.
76. Lei D, Li F, Su H, Wei N, and **Wang X***. Hepatic Deficiency of COP9 Signalosome Subunit 8 Induces Ubiquitin-Proteasome System Impairment and Bim-mediated Apoptosis in Murine Livers. *PLoS ONE* 2013; 8(7): e67793. doi:10.1371/journal.pone.0067793.
77. Zhao C[§], Liu S[§], Yang C, Li S, Huang H, Liu N, Li S, **Wang X***, and Liu J*. Gambogic acid moderates cardiac responses to chronic hypoxia likely by acting on the proteasome and NF-κB pathway. *Am J Cardiovasc Dis* 2013; 3(3): 135-145.

78. Su H, Li J, Osinska H, Li F, Robbins J, Wei N, and **Wang X***. The COP9 signalosome is required for autophagy, proteasome-mediated proteolysis, and cardiomyocyte survival in adult mice. *Circ Heart Fail* 2013; 6(5): 1049-1057.
79. Li S, **Wang X**, Li Y, Kost CK Jr., and Martin DS. Bortezomib, a proteasome inhibitor, attenuates angiotensin II-induced hypertension and aortic remodeling in rats. *PLoS ONE* 2013; 8(10): e78564. doi:10.1371/journal.pone.0078564.
80. Liu S[§], Zhao C[§], Yang C, Li X, Huang H, Liu N, Li S, **Wang X***, and Liu J*. Gambogic acid suppresses pressure overload cardiac hypertrophy in rats. *Am J Cardiovasc Dis* 2013; 3(4): 227-238.
81. Shi X, Chen X, Li X, Lan X, Zhao C, Liu S, Huang H, Liu N, Liao S, Song W, Zhou P, Wang S, **Wang X**, Dou QP, and Liu J. Gambogic acid induces apoptosis in chronic myeloid leukemia cells resistant to imatinib via inducing proteasome inhibition and caspase-dependent Bcr-Abl cleavage. *Clin Cancer Res* 2014; 20(1):151-63. doi: 10.1158/1078-0432.CCR-13-1063.
82. Liu Y, Hettinger CL, Zhang D, Rezvani K, **Wang X**, Wang H. The Proteasome Function Reporter GFPu Accumulates in Young Brains of the APP^{swe}/PS1^{dE9} Alzheimer's Disease Mouse Model. *Cell Mol Neurobiol* 2014; 34(3):315-322.
83. Liu Y, Hettinger CL, Zhang D, Rezvani K, **Wang X**, and Wang H. Sulforaphane enhances proteasomal and autophagic activities in mice and is a potential therapeutic reagent for Huntington's disease. *J Neurochem* 2014; 129(3):539-47. doi: 10.1111/jnc.12647.
84. Liu Y, Lü L, Hettinger C, Dong G, Zhang D, Rezvani K, **Wang X**, and Wang H. Ubiquilin-1 protects cells from oxidative stress and ischemic stroke caused tissue injury in mice. *J Neurosci* 2014; 34(8):2813-21. doi: 10.1523/JNEUROSCI.3541-13.2014.
85. Ranek MJ, Kost CK, Martin DS, and **Wang X***. Muscarinic 2 receptors modulate cardiac proteasome function in a protein kinase G-dependent manner. *J Mol Cell Cardiol* 2014; 69:43-51. doi: 10.1016/j.yjmcc.2014.01.017.
86. Liu N, Li X, Liao S, Huang H, Zhao C, Yang C, Liu S, Song W, Lu X, Xu L, Jiang L, Zhao C, Dong X, Zhou P, Wang S, **Wang X**, Dou QP, and Liu J. A novel proteasome inhibitor inhibits tumor growth *in vitro*, *in vivo* and *ex vivo* via targeting both 19S proteasome deubiquitinases and 20S proteolytic peptidases. *Sci Rep* 2014; 4:5240. doi: 10.1038/srep05240.
87. Tian Z, Wang C, Hu C, Tian Y, Liu J, and **Wang X***. Autophagic-lysosomal inhibition impairs cardiac ubiquitin-proteasome system performance in a p62 dependent manner. *PLoS ONE* 2014; 9(6):e100715. doi: 10.1371/journal.pone.0100715.
88. Liu N[§], Li X[§], Huang H[§], Zhao C[§], Liao S[§], Yang C[§], Liu S[§], Song W, Lu X, Lan X, Chen X, Xu L, Jiang L, Zhao C, Dong X, Zhou P, Li S, Wang S, Shi X, Dou QP, **Wang X**, and Liu J.

- Clinically used antirheumatic agent auranofin is a proteasomal deubiquitinase inhibitor and inhibits tumor growth. *Oncotarget* 2014; 5(14):5453-71.
89. Gupta MK, Gulick J, Liu R, **Wang X**, Molkenin JD, and Robbins J. SUMO E2 enzyme UBC9 is required for efficient protein quality control in cardiomyocytes. *Circ Res.* 2014; 115(8):721-9. doi: 10.1161/CIRCRESAHA.115.304760. (with companion editorial)
90. Chen X, Shi X, Zhao C, Li X, Lan X, Liu S, Huang H, Liu N, Liao S, Zang D, Song W, Liu Q, Carter BZ, Dou QP, **Wang X**, and Liu J. Anti-rheumatic agent auranofin induced apoptosis in chronic myeloid leukemia cells resistant to imatinib through both Bcr/Abl-dependent and -independent mechanisms. *Oncotarget.* 2014 Oct 15; 5(19):9118-32.
91. Shi X, Lan X, Li X, Chen X, **Wang X**, and Liu J. 2-tert-butyl-1,4-benzoquinone induces apoptosis in chronic myeloid leukemia cells resistant to imatinib via inducing caspase-dependent Bcr-Abl downregulation. *Med Chem* 2014; 4 (12): 784-790. doi:10.4172/2161-0444.1000231
92. Shi X, Lan X, Chen X, Zhao C, Li X, Liu S, Huang H, Liu N, Zhang D, Liao Y, Zhang P, **Wang X**, Liu J. Gambogic acid induces apoptosis in diffuse large B-cell lymphoma cells via inducing proteasome inhibition. *Sci Rep.* 2015 Apr 8; 5:9694.
93. Ranek MJ[§], Zheng H[§], Huang W[§], Kumarapeli AR, Li J, Liu J, and **Wang X***. Genetically induced moderate inhibition of 20S proteasomes in cardiomyocytes facilitates heart failure in mice during systolic overload. *J Mol Cell Cardiol.* 2015; 85 (8):273-81. (with companion editorial)
94. Li J, Ma W, Li H, Hou N, **Wang X**, Kim IM, Li F, and Su H. NEDD8 Ultimate Buster-1 Long (NUB1L) Protein Suppresses Atypical Neddylation and Promotes Proteasomal Degradation of Misfolded Proteins. *J Biol Chem* 2015; 290(39):23850-62.
95. Su H, Li J, Zhang H, Ma W, Wei N, Liu J*, and **Wang X***. The COP9 signalosome controls the degradation of cytosolic misfolded proteins and protects against cardiac proteotoxicity. *Circ Res* 2015 117(11):956-66. (with companion editorial)
96. Qin Q, Qu C, Niu T, Zang H, Qi L, Lyu L, **Wang X**, Janicki J, Wang XL, Cui T. Nrf2-mediated cardiac maladaptive remodeling and dysfunction in a setting of autophagy insufficiency. *Hypertension* 2016 Jan; 67(1):107-17. doi: 10.1161/HYPERTENSIONAHA.115.06062.
97. Huang H[§], Liao Y[§], Liu N, Hua X, Cai J, Yang C, Zhao C, Chen X, Lan X, Zang D, Wu J, Li X, Shi X, **Wang X**, and Liu J. Two clinical drugs deubiquitinase inhibitor auranofin and aldehyde dehydrogenase inhibitor disulfiram trigger synergistic anti-tumor effects in vitro and in vivo. *Oncotarget* 2016; 7(3): 2098-2808. doi: 10.18632/oncotarget.6425.

98. Jiang L[§], Zang D[§], Yi S[§], Li X[§], Yang C, Dong X, Zhao C, Lan X, Chen X, Liu S, Liu N, Huang H, Shi X, **Wang X***, and Liu J*. A microRNA-mediated decrease in eukaryotic initiation factor 2 α promotes cell survival during endoplasmic reticulum stress. *Sci Rep*. 2016; 6:21565. DOI: [10.1038/srep21565](https://doi.org/10.1038/srep21565)
99. Zhao C[§], Chen X[§], Zang D, Lan X, Liao S, Yang C, Zhang P, Wu J, Li X, Liu N, Liao Y, Huang H, Shi X, Jiang L, Liu X, Dou QP, **Wang X**, and Liu J. A novel nickel complex works as a proteasomal deubiquitinase inhibitor for cancer therapy. *Oncogene* 2016; 35(45):5916-5927. doi: 10.1038/onc.2016.114.
100. Zhang L, Hapona MB, Goyeneche AA, Srinivasan R, Carlos D. Gamarra-Luques CD, Callegari EA, Drappeau DD, Terpstra EJ, Knapp J, Chien J, **Wang X**, Eyster KM, and Telleria CM. Mifepristone induces endoplasmic reticulum stress, triggers the unfolded protein response, increases autophagic flux, and kills ovarian cancer cells in combination with bortezomib or chloroquine. *Molecular Oncology* 2016; 10(7):1099-1117. doi: 10.1016/j.molonc.2016.05.001.
101. Hou N[§], Ye B[§], Li X, Margulies KB, Xu H, **Wang X**, and Li F. TCF7L2 Mediates Canonical Wnt/ β -Catenin Signaling and c-Myc Upregulation in Heart Failure. *Circ Heart Fail* 2016 Jun; 9(6). pii: e003010. doi: 10.1161/CIRCHEARTFAILURE.116.003010. Epub 2016 Jun 14
102. Zhao C, Chen X, Zang D, Lan X, Liao S, Yang C, Zhang P, Wu J, Li X, Liu N, Liao Y, Huang H, Shi X, Jiang L, He Z, Liu X, **Wang X***, and Liu J*. Platinum-containing compound platinum pyrithione is stronger and safer than cisplatin in cancer therapy. *Biochemical Pharmacology* 2016; 116: 22-38.
103. Lan X, Zhao C, Chen X, Zhang P, Zang D, Wu J, Chen J, Long H, Yang L, Huang H, Carter BZ, **Wang X**, Shi X*, Liu J*. Nickel pyrithione induces apoptosis in chronic myeloid leukemia cells resistant to imatinib via both Bcr/Abl-dependent and -independent mechanisms. *J Hematol Oncol* 2016 Nov 25; 9(1):129. DOI: [10.1186/s13045-016-0359-x](https://doi.org/10.1186/s13045-016-0359-x).
104. Li J, Ma W, Yue G, Johnson J, Kim I-M, Weintraub N, **Wang X**, and Su H. Cardiac proteasome functional insufficiency plays a pathogenic role in diabetic cardiomyopathy. *J Mol Cell Cardiol* 2016 Nov 30; 102:53-60. DOI: <http://dx.doi.org/10.1016/j.yjmcc.2016.11.013>
105. Zhao C[§], Chen X[§], Yang C[§], Zang D, Lan X, Liao S, Zhang P, Wu J, Li X, Liu N, Liao Y, Huang H, Shi X, Jiang L, Liu X, **Wang X***, and Liu J*. Repurposing an antidandruff agent to treating cancer: zinc pyrithione inhibits tumor growth via targeting proteasome-associated deubiquitinases. *Oncotarget*. 2017 Feb 21; 8(8):13942-13956. doi: 10.18632/oncotarget.14572.
106. Liao Y, Liu N, Hua X, Cai J, Xia X, **Wang X**, Huang H, Liu J. Proteasome-associated deubiquitinase ubiquitin-specific protease 14 regulates prostate cancer proliferation by deubiquitinating and stabilizing androgen receptor. *Cell Death Dis*. 2017 Feb 2; 8(2):e2585. DOI: [10.1038/cddis.2016.477](https://doi.org/10.1038/cddis.2016.477)

107. Balla C, Assenza GE, Subramanian K, **Wang X**, Volpe M, Dec GW, del Monte F. Presenilin-2, a Novel Excitation Contraction Coupling Protein, Modulates Both Ca²⁺ Release and Ca²⁺ Reuptake in Cardiomyocytes. 2015 (revised)
108. Xiao L, Lan X, Shi X, Zhao K, **Wang X**, Li F, Huang H, Liu J. Cytoplasmic RAP1 mediates cisplatin resistance of non-small cell lung cancer. *Cell Death Dis.* 2017 May 18; 8(5):e2803. (DOI: [10.1038/cddis.2017.210](https://doi.org/10.1038/cddis.2017.210)). PubMed PMID: 28518145.
109. Huang H[§], Guo M[§], Liu N[§], Zhao C[§], Chen H, Wang X, Liao S, Zhou P, Liao Y, Chen X, Lan X, Xu D, Li X, Shi X, **Wang X***, Zhang C-E*, and Liu J*. Bilirubin neurotoxicity is associated with proteasome inhibition. *Cell Death Dis.* 2017; 8(6): e2877 (DOI: [10.1038/cddis.2017.274](https://doi.org/10.1038/cddis.2017.274))
110. Cai J[§], Xia X[§], Liao Y[§], Liu N, Guo Z, **Wang X**, Huang H*, Liu J*. A novel deubiquitinase inhibitor b-AP15 triggers apoptosis in both androgen receptor-dependent and -independent prostate cancers. *Oncotarget* 2017 Jun 28; 8(38):63232-63246. doi: [10.18632/oncotarget.18774](https://doi.org/10.18632/oncotarget.18774). eCollection 2017 Sep 8.
111. Chen X, Wu J, Zhang P, Liao S, Lan X, Chen J, Li X, Huang H, Liao Y, Zhang X, Yang Q, Shi X, Jiang L, Liu N, He Z, **Wang X**, Zhao C, and Liu J. Cadmium pyrithione suppresses tumor growth in vitro and in vivo through inhibition of proteasomal deubiquitinase. *Biometals* 2017 Nov 3; <https://doi.org/10.1007/s10534-017-0062-6>. [Epub ahead of print]
112. Chen X, Zhang X, Chen J, Yang Q, Yang L, Xu D, Zhang P, Lan X, Liao Y, Long H, Cai J, Li X, Huang H, Jiang L, **Wang X**, Liu J. Hinokitiol copper complex inhibits proteasomal deubiquitination and induces paraptosis-like cell death in human cancer cells. *European J Pharmacol* 2017 2017 Sep 5. pii: S0014-2999(17)30578-2. doi: 10.1016/j.ejphar.2017.09.003. [Epub ahead of print]
113. Lan X, Zhao C, Chen X, Zhang P, Zang D, Wu J, Chen J, Long H, Yang L, Huang H, **Wang X**, Shi X, Liu J. Platinum pyrithione induces apoptosis in chronic myeloid leukemia cells resistant to imatinib via DUB inhibition-dependent caspase activation and Bcr-Abl down-regulation. *Cell Death Dis.* 2017; 8(7): e2913. (DOI: [10.1038/cddis.2017.284](https://doi.org/10.1038/cddis.2017.284)).
114. Wu P, Yuan X, Li F, Zhang J, Zhu W, Wei M, Li J*, **Wang X***. Myocardial upregulation of cathepsin D by ischemic heart disease promotes autophagic flux and protects against cardiac remodeling and heart failure. *Circ Heart Fail* 2017; 10(7) (DOI: [10.1161/CIRCHEARTFAILURE.117.004044](https://doi.org/10.1161/CIRCHEARTFAILURE.117.004044))
115. Abdullah A, Eyster KM, Bjordahl T, Xiao P, Zeng E, **Wang X***. Murine myocardial transcriptome analysis reveals a critical role of COPS8 in the gene expression of Cullin-RING ligase substrate receptors and redox and vesicle trafficking pathways. *Frontiers in Physiology-System Biology* 17 August 2017 | <https://doi.org/10.3389/fphys.2017.00594>.

116. Pan B, Zhang H, Cui T, **Wang X***. TFEB activation protects against cardiac proteotoxicity via increasing autophagic flux. *J Mol Cell Cardiol* 2017 Dec; 113: 51-62. DOI: [10.1016/j.yjmcc.2017.10.003](https://doi.org/10.1016/j.yjmcc.2017.10.003). Epub 2017 Oct 7.
117. Hou N, Xu H, **Wang X**, Li F, Ye B. Activation of Yap1/Taz signaling in ischemic heart disease and dilated cardiomyopathy. *Exp Mol Path*. 2017; 103(3):267-275.
118. Reihel CA, Pekas N, Wu P, **Wang X***. Systemic inhibition of neddylation by 3-day MLN4924 treatment regime does not impair autophagic flux in mouse hearts and brains. *Am J Cardiovasc Dis* 2017 Dec 20;7(6):134-150.
119. Sane S, Hafner A, Srinivasan R, Masood D, Slunicka JL, Noldner CJ, Hanson AD, Kruisselbrink T, **Wang X**, Wang Y, Yin J, Rezvani K. UBXN2A enhances CHIP-mediated proteasomal degradation of oncoprotein mortalin-2 in cancer cells. *Mol Oncol* 2018; 12(10):1753-1777. DOI:[10.1002/1878-0261.12372](https://doi.org/10.1002/1878-0261.12372)
120. Chen X[#], Yang Q[#], Chen J, Zhang P, Huang Q, Zhang X, Yang L, Xu D, Zhao C, **Wang X**, Liu J. Inhibition of proteasomal deubiquitinase by silver complex induces apoptosis in non-small cell lung cancer cells. *Cell Physiol Biochem*. 2018; 49(2):780-797. DOI:[10.1159/000493041](https://doi.org/10.1159/000493041)
121. Huang H, Xia X, Liao Y, Guo Z, Li Y, Jiang L, Zhang F, Huang C, Liu Y, **Wang X**, Liu N, and Liu J. Targeting proteasome associated deubiquitinases as a novel strategy for the treatment of estrogen receptor positive breast cancer. *Oncogenesis*. 2018 Sep 24; 7(9):75. DOI: <https://doi.org/10.1038/s41389-018-0086-y>
122. Hu C, Tian Y*, Xu H, Pan B, Terpstra EM, Wu P, Wang H, Li F, **Wang X***. Inadequate ubiquitination-proteasome coupling contributes to myocardial ischemia-reperfusion injury. *J Clin Invest*. 2018; 128(12):5294-5306. First published 2018 Sep 11. DOI:[10.1172/JCI98287](https://doi.org/10.1172/JCI98287) (with complimentary Commentary)
123. Liu N, Guo Z, Xia X, Liao Y, Zhang F, Huang C, Liu Y, Deng X, Jiang L, Wang X, Liu J, Huang H. Auranofin lethality to prostate cancer includes inhibition of proteasomal deubiquitinases and disrupted androgen receptor signaling. *Eur J Pharmacol*. 2019 Mar 5; 846:1-11. Epub 2019 Jan 9. DOI:[10.1016/j.ejphar.2019.01.004](https://doi.org/10.1016/j.ejphar.2019.01.004)
124. Liao Y, Guo Z, Xia X, Liu Y, Huang C, Jiang L, **Wang X**, Liu J, and Huang H. Inhibition of EGFR signaling with Sunitinib represents a novel therapeutic for prostate cancer. *J Exp Clin Cancer Res*. 2019; 38:157 (12 pages). DOI: [10.1186/s13046-019-1165-4](https://doi.org/10.1186/s13046-019-1165-4)
125. Zhang H, Pan B, Wu P, Parajuli N, Rekhter MD, Goldberg AL, **Wang X***. PDE1 inhibition facilitates proteasomal degradation of misfolded proteins and protects against cardiac proteinopathy. *Science Advances* 2019 May 22; 5(5):eaaw5870. PMCID: [PMC6531002](https://pubmed.ncbi.nlm.nih.gov/6531002/) DOI: [10.1126/sciadv.aaw5870](https://doi.org/10.1126/sciadv.aaw5870) PMID: 31131329.

126. Xia X, Huang C, Liao Y, Liu Y, He J, Guo Z, Jiang L, Wang X, Liu J, Huang H. Inhibition of USP14 enhances the sensitivity of breast cancer to enzalutamide. *J Exp Clin Cancer Res*. 2019 May 24; 38(1):220. DOI:[10.1186/s13046-019-1227-7](https://doi.org/10.1186/s13046-019-1227-7)
127. Xia X, Liu Y, Liao Y, Guo Z, Huang C, Zhang F, Jiang L, **Wang X**, Liu J, Huang H. Synergistic effects of gefitinib and thalidomide treatment on EGFR-TKI-sensitive and -resistant NSCLC. *Eur J Pharmacol*. 2019 May 24:172409. DOI:[10.1016/j.ejphar.2019.172409](https://doi.org/10.1016/j.ejphar.2019.172409) PMID: 31132355.
128. Pan B, Lewno M, Wu P, **Wang X***. Highly Dynamic changes in the activity and regulation of macroautophagy in hearts subjected to increased proteotoxic stress. *Front Physiol*. 2019;10:758. doi: 10.3389/fphys.2019.00758. eCollection 2019. PubMed PMID: 31297061; PubMed Central PMCID: PMC6606963.
129. Gao H, Freeling J, Wu P, Liang AP, **Wang X**, Li Y. [UCHL1 regulates muscle fibers and mTORC1 activity in skeletal muscle](https://doi.org/10.1016/j.lfs.2019.116699). *Life Sci*. 2019 Sep 15; 233:116699. doi: 10.1016/j.lfs.2019.116699. Epub 2019 Jul 26. PubMed PMID: 31356902; PubMed Central PMCID: PMC6718320.
130. Chen H, Liang L, Xu H, Xu J, Yao L, Li Y, Tan Y, Li X, Huang Q, Yang Z, Wu J, Chen J, Huang H, **Wang X**, Zhang C-E, and Liu J. Short term exposure to bilirubin induces encephalopathy similar to Alzheimer's disease in late life. *J Alzheimer Dis* 2020 Jan 7; 73(1):277-295, DOI: [10.3233/JAD-190945](https://doi.org/10.3233/JAD-190945). PMID: 31796680.
131. Pan B[#], Li J[#], Parajuli N[#], Tian Z[#], Wu P[#], Lewno MT, Bedford L, Mayer RJ, Fang J, Liu J, Cui T, Su H*, **Wang X***. The calcineurin-TFEB-p62 pathway mediates the activation of cardiac macroautophagy by proteasomal malfunction. *Circ Res* 2020 July 31; 127(4):502–518. DOI: [10.1161/CIRCRESAHA.119.316007](https://doi.org/10.1161/CIRCRESAHA.119.316007). ([#]equal contributors) (with companion editorial: <https://doi.org/10.1161/CIRCRESAHA.120.317567>)
132. Xiao P, Wang C*, Lewno MT, Wu P, Li J, Su H, Sternburg JO, Liu J, **Wang X***. The COP9 signalosome suppresses RIPK1-RIPK3 mediated cardiomyocyte necroptosis in mice. *Circ Heart Fail* 2020 Aug; 13(8):283-294 (e006996). <https://doi.org/10.1161/CIRCHEARTFAILURE.120.006996>.
133. Qi L, Zang H, Wu W, Nagarkatti P, Nagarkatti M, Liu QC, Robbins J, **Wang X**, Cui T. CYLD exaggerates pressure overload-induced cardiomyopathy via suppressing autolysosome efflux in cardiomyocytes. *J Mol Cell Cardiol* 2020 June 14; 145(8):59-73. <https://authors.elsevier.com/c/1bHeP54GVbhKr>
134. Zhong X, Liao Y, Chen X, Mai N, Ouyang C, Chen B, Zhang M, Peng Q, Liang W, Zhang W, Wu Z, Huang X, Li C, Chen H, Lao W, Zhang CE, **Wang X**, Ning Y, and Liu J. Abnormal serum bilirubin/albumin concentrations in dementia patients with A β deposition and the benefit of intravenous albumin infusion for Alzheimer's disease treatment. *Frontiers in Neuroscience*:

Neurodegeneration 03 September 2020; 14:859. doi: 10.3389/fnins.2020.00859
| <https://doi.org/10.3389/fnins.2020.00859>

135. Zang H, Wu W, Qi L, Nagarkatti P, Nagarkatti M, **Wang X**, Cui T. Autophagy inhibition enables Nrf2 to exaggerate the progression of diabetic cardiomyopathy in mice. *Diabetes* 2020 Dec; 69(12): 2720-2734 <https://doi.org/10.2337/db19-1176>
136. Gao H, Antony R, Srinivasan R, Wu P, **Wang X**, Li Y. UCHL1 regulates oxidative activity in skeletal muscle. *PLOS One* 2020 November 2; 15(11):e0241716. (12 pages). PMID: 33137160; PMCID: [PMC7605647](https://pubmed.ncbi.nlm.nih.gov/33137160/) DOI: [10.1371/journal.pone.0241716](https://doi.org/10.1371/journal.pone.0241716)
137. Liu Y, Subedi K, Baride A, Romanova S, Huber CC, **Wang X**, Wang H. Peripherally misfolded proteins exacerbate ischemic stroke-induced neuroinflammation and brain injury. *J Neuroinflammation* 2021; 18: Article number 29. <https://doi.org/10.1186/s12974-021-02081-7>
138. Wu W, Qin Q, Ding Y, Zang H, Li DS, Nagarkatti M, Nagarkatti P, Wang W, **Wang X***, and Cui T*. Autophagy controls Nrf2-mediated dichotomy in pressure overloaded hearts. *Frontiers in Physiology-Striated Muscle Physiology* 2021 May 13, <https://doi.org/10.3389/fphys.2021.673145>
139. Li Y, Liu H, Chen K. Wu X. Wu J, Yang Z, Yao L, Wen G, Zhang C, Chen X, Tang D, **Wang X**, Liu J. Pathological significance and prognostic roles of indirect bilirubin/albumin ratio in hepatic encephalopathy. *Front. Med. - Translational Medicine* 2021 August 30; 8:706407. doi:10.3389/fmed.2021.706407. <https://doi.org/10.3389/fmed.2021.706407>. PMID: 34527681; PubMed Central PMCID: PMC8435674.
140. Wu P, Cai M, Liu J, **Wang X***. Catecholamine surges cause cardiomyocyte necroptosis via the RIPK1-RIPK3 pathway in mice. *Front. Cardiovasc. Med.*, 16 September 2021; <https://doi.org/10.3389/fcvm.2021.740839>
141. Hussain A, Mariappan K, Cork DC, Lewandowski L, Shrestha PK, Giri S, **Wang X**, Sykes AG. A highly selective pyridoxal-based chemosensor for the detection of Zn(II) and application in live cell imaging; X-ray crystallography of pyridoxal-TRIS Schiff-base Zn(II) and Cu(II) complexes. *RSC Advances*. 2021 Sep (*in press*).
142. Xu H, Wu X, Liang L, Chen H, Xu J, Hu W, Li X, Liu Q, **Wang X***, Zhang C-E*, Liu J*. USP14 haploinsufficiency ameliorates Alzheimer's disease-like pathology in APP/PS1 mice. (Preprint DOI: [10.21203/rs.3.rs-68805/v1](https://doi.org/10.21203/rs.3.rs-68805/v1))

B. INVITED REVIEW ARTICLES

143. **Wang X**, Dong C. Apoptosis and ischemic-reperfusion injury. *Foreign Medical Sciences (Physiology, Pathology, and Clinic)* 1997; 17(2): 125-127.

144. **Wang X***, Osinska H, Gerdes AM, and Robbins J. Desmin and cardiac diseases: establishing causality. *J Card Failure* 2002; 8: S287-S292. (*correspondence)
145. Li F, **Wang X**, Yi XP, and Gerdes AM. Structural Basis of Ventricular Remodeling: Role of the Myocyte. *Current Heart Failure Reports* 2004; 1(1): 5-8.
146. Kumarapeli A, **Wang X***. Genetic Modification of the Heart: Chaperones and the Cytoskeleton. *J Mol Cell Cardiol* 2004; 37: 1097-1109.
147. **Wang X** and Robbins J. Heart Failure and Protein Quality Control. *Circ Res* 2006; 99: 1315-1326.
148. **Wang X**, Su H, and Ranek M. Protein quality control and degradation in cardiomyocytes (Invited Review). *J Mol Cell Cardiol* 2008; 45(1):11-27.
149. Zheng Q, Li J, and **Wang X***. Interplay between the ubiquitin-proteasome system and autophagy in proteinopathy. *Int J Physiol Pathophysiol Pharmacol* 2009; 1(2):127-142.
150. Ranek M and **Wang X***. Activation of the ubiquitin-proteasome system in doxorubicin cardiomyopathy. *Curr Hypertens Rep* 2009; 11(6):389-95.
151. Su H and **Wang X***. The ubiquitin-proteasome system in cardiac proteinopathy: a quality control perspective. *Cardiovasc Res* 2010; 85(2):253-62.
152. Depre C, Powell SR, and **Wang X**. The role of the ubiquitin-proteasome pathway in cardiovascular disease. *Cardiovasc Res* 2010; 85(2):251-2.
153. Zheng Q and **Wang X***. Autophagy and the ubiquitin-proteasome system in cardiac dysfunction. *Panminerva Med.* 2010; 52(1):9-25.
154. Li Y-F and **Wang X***. The Role of the Proteasome in Heart Disease. *Biochim Biophys Acta* 2011;1809(2):141-149.
155. **Wang X***, Li J, Zheng H, Su H, and Powell SR. Proteasome functional insufficiency in cardiac pathogenesis. *Am J Physiol Heart Circ Physiol* 2011; 301(6): H2207-2219.
156. Su H and **Wang X***. p62 stages an interplay between the ubiquitin-proteasome system and autophagy in the heart of defense against proteotoxic stress. *Trends Cardiovasc Med* 2011; 21(8): 224-228.
157. Powell SR, Herrmann J, Lerman A, Patterson C, and **Wang X**. The ubiquitin-proteasome system and cardiovascular disease. *Prog Mol Biol Transl Sci* 2012; 109: 295-346.
158. Klionsky DJ, many others, **Wang X**, et al. [Guidelines for the use and interpretation of assays for monitoring autophagy \(2nd edition\)](#). *Autophagy* 2012; 8(4):445-544. (Contributed a full figure)
159. **Wang X*** and Terpstra EJ. Ubiquitin Receptors and Protein Quality Control. *J Mol Cell Cardiol* 2013; 55(2):73-84. doi: 10.1016/j.yjmcc.2012.09.012. Epub 2012 Oct 6.

160. **Wang X***, Pattison JS, and Su H. Posttranslational modification and protein quality control (Invited Review). *Circ Res* 2013; 112(2): 367-81.
161. Chen X, Shi X, **Wang X**, and Liu J. Novel use of old drug: Anti-rheumatic agent auranofin overcomes imatinib-resistance of chronic myeloid leukemia cells. *Can Cell Microenviron* 2014; 1: e415. doi: 10.14800/ccm.415.
162. **Wang X*** and Robbins J. Proteasomal and lysosomal protein degradation and heart disease (Invited Review). *J Mol Cell Cardiol* 2014; 71C:16-24. doi: 10.1016/j.yjmcc.2013.11.006.
163. Wang C and **Wang X***. The interplay between autophagy and the ubiquitin-proteasome system in cardiac proteotoxicity (invited review). *BBA-Mol Basis Dis* 2015; 1852(2):188-94.
164. Zhang H and **Wang X***. Priming the proteasome: a novel mechanism for cardioprotection by sildenafil. *Future Cardiology* 2015; 11(2):177-89. doi: 10.2217/fca.15.3.
165. **Wang X*** and Martin DS. The COP9 signalosome and cullin RING ligases in the heart. *Am J Cardiovasc Dis* 2015; 5(1):1-18.
166. Martin DS* and **Wang X***. The COP9 signalosome in vascular function. *Am J Cardiovasc Dis* 2015; 5(1):33-52.
167. Klionsky DJ, many others, **Wang X**, *et al.* Guidelines for the use and interpretation of assays for monitoring autophagy (3rd edition). *Autophagy* 2016; 12(1):1-222. (Contributed a full figure).
168. Cui T*, Lai Y, Janicki JS, and **Wang X***. Nrf2-mediated protein quality control in cardiomyocytes. *Frontiers in Bioscience* (Landmark Ed). 2016 Jan 1; 21:192-202. PubMed PMID: 26709769.
169. **Wang X***, Cui T*. Modulation of autophagy – a potential therapeutic approach for cardiac hypertrophy (Invited review). *Am J Physiol-Heart Circ Physiol* 2017 Aug 1; 313(2):H304-H319. doi: 10.1152/ajpheart.00145.2017. Epub 2017 Jun 2. Review. PubMed PMID: 28576834.
170. **Wang X***, Wang H. Priming the proteasome to protect against proteotoxicity. *Trends in Molecular Medicine* 2020 July 1; 26(7):639-648. DOI: [10.1016/j.molmed.2020.02.007](https://doi.org/10.1016/j.molmed.2020.02.007).
171. Klionsky DJ, many others, **Wang X**, *et al.* [Guidelines for the use and interpretation of assays for monitoring autophagy \(4th Edition\)](https://doi.org/10.1080/15548627.2020.1797280). (Contributed a full figure, Figure 17). *Autophagy* 2021 Feb 8 (Epub ahead of print). DOI: 10.1080/15548627.2020.1797280 <https://www.tandfonline.com/doi/abs/10.1080/15548627.2020.1797280>
172. Lewno MT, Cui T, **Wang X***. Cullin Deneddylation Suppresses the Necroptotic Pathway in Cardiomyocytes. *Frontiers in Physiology-Striated Muscle Physiology* 2021 June 28, DOI:[10.3389/fphys.2021.690423](https://doi.org/10.3389/fphys.2021.690423)

C. INVITED EDITORIALS

173. **Wang X***, Su H. Unraveling Enigma in the Z-disk. *Circ Res* 2010; 107:321-3.
174. Su H, **Wang X***. Autophagy and p62 in cardiac protein quality control. *Autophagy* 2011; 7(11): 1382-1383.
175. **Wang X***, Su H. FoxO3 hastens autophagy and shrinks the heart but does not curtail pathological hypertrophy in adult mice. *Cardiovasc Res* 2011; 91 (4):561-62.
176. **Wang X***. Repeated Intermittent Administration of a Ubiquitous Proteasome Inhibitor Leads to Restrictive Cardiomyopathy. *Eur J Heart Fail* 2013; 15(6): 597-598.
177. Liu J*, Su H, **Wang X.*** The COP9 signalosome coerces autophagy and the ubiquitin-proteasome system to police the heart. *Autophagy* 2016; 12(3):601-602.
178. **Wang X***. Entangled in a heart-ailing quandary: could modified cofilin-2 be a culprit of Alzheimer's disease of the heart? *J Am Coll Cardiol* 2015; 65(12):1215-7.
179. **Wang X***. Vascular Spasm: A Newly Unraveled Cause for Cardiovascular Adversity of Proteasome Inhibition. *EBioMedicine* 2017. (DOI: [10.1016/j.ebiom.2017.06.010](https://doi.org/10.1016/j.ebiom.2017.06.010))
180. Su H, **Wang X***. Proteasome malfunction activates the calcineurin-TFEB-p62 pathway to induce macroautophagy in the heart. *Autophagy* 2020; 16(11): 2114-2116. DOI: [10.1080/15548627.2020.1816666](https://doi.org/10.1080/15548627.2020.1816666). Epub ahead of print on 2020 Sep 22.
181. Rank MJ, Bhuiyan MS, **Wang X***. Editorial: Targeting cardiac proteotoxicity. *Front Physiol.* 2021;12:669356. DOI: [10.3389/fphys.2021.669356](https://doi.org/10.3389/fphys.2021.669356)

D. BOOKS and CHAPTERS IN BOOKS

182. **Wang X** and Dong C. Chapter 21. Prostaglandins. in Yang G, eds. *Endocrinal Physiology and Pathophysiology*. Tianjing: Tianjing Science and Technology Press 1995; 819-851.
183. Zhen XH, Xia ZP, Xia ZX, and **Wang X**, co-eds. *Advanced Education Self-Study Examination Review Guide: Pathophysiology and Pharmacology*. 1st ed. Wuhan: Hubei Science and Technology Press 1993; pp1-609.
184. Gerdes AM and **Wang X**. Structural remodeling of cardiac myocytes in hypertrophy and progression to failure. In: "Cardiovascular Remodeling and failure". Eds. PK Singal, IMC Dixon, LA Kirshenbaum, NS Dhalla. Kluwer Academic Publishers, Boston, MA, 2003; pp183-193.
185. **Wang X** and Patterson C. Chapter 27. Protein quality control in cardiomyocytes; In: Hill JA, Olson EN, eds. *Muscle: Fundamental Biology and Mechanisms and Disease*. Elsevier 2012; pp. 353-368.

186. Su H and **Wang X***. Chapter 13, Defense against proteotoxic stress in the heart: role of p62, autophagy, and ubiquitin-proteasome system. In: M. A. Hayat, eds. ***Autophagy: Cancer, Other Pathologies, Inflammation, Immunity, Infection, and Aging***. Elsevier 2014; Vol 3, pp. 188-202.
187. **Wang X***. Chapter 12. Desmin Filaments and Desmin-Related Myopathy. In: Schatten H, eds. ***The Cytoskeleton in Health and Disease***. Springer Science+Business Media New York 2015; 281-306.
188. Cui T and **Wang X***. Chapter 3, Interplay among oxidative stress, redox signaling, ER-stress, autophagy and protein ubiquitination. In Ren, Sowers, Zhang, eds. ***Autophagy and Cardiometabolic Diseases: From Mechanisms to Molecules to Medicine***. Academic Press, May 1, 2018 - Medical - 224 pages.
189. **Wang, X.**, Li, H.-H., Bhuiyan, M. S., and Ranek, M. J. eds. (2021). ***Targeting Cardiac Proteotoxicity***. Lausanne: Frontiers Media SA. doi: 10.3389/978-2-88966-806-9 May 2021, 175 pages.
190. **Wang X.** Chapter 6. Protein degradation in cardiac health and disease. In Chondrogianni, Pick, Gioran, eds. ***Proteostasis***. CRC Press: Taylor & Francis Group. (in press).

(*Corresponding author; §contributed equally)

E. ABSTRACTS AND CONFERENCE PRESENTATIONS

1. Dong, C., X. Wang, and J. Xiang. The kinetic changes of coagulation and anti-coagulation of patients with epidemic hemorrhagic fever. *FASEB J* 1991; 5(6): 1628.
2. Gerdes A. M., T. Onodera, F. Li, X. Wang, S. A. McCune, and J. M. Capasso. Regional changes in cardiac myocyte shape during the progression to failure in rats. *J Mol Cell Cardiol* 1996; 28:A200.
3. Li F, X. Wang, AM Gerdes. Cellular mechanism of binucleation in neonatal rat cardiocytes. *Circulation* 1996; 94(8): No.3544.
4. Wang X, S.E. Campbell, A.M. Gerdes. Regional alterations of microtubule density in cardiac myocytes following chronic aortic stenosis in guinea pigs. *Circulation* 1997;96(8): I-254.
5. Wang X, and Gerdes AM. Altered expression and distribution of intercalated disks associated proteins in ventricular myocytes from guinea pigs with chronic pressure overloaded cardiac hypertrophy and failure. *Journal of Cardiac Failure* 1998; 3(2 Suppl. 1): 58.
6. Wang X, Gulick J, Oscinska H, Hewett T, Robbins J. A desminopathy-associated desmin mutation causes aberrant desmin aggregation and early cardiac hypertrophy in

transgenic mice. A Keystone Symposia: Molecular Biology of the Cardiovascular System, January 12-17, 2000, Snowbird, UT.

7. Wang X., H. Osinska, G.W. Dorn II, A.M. Gerdes, R. Klevitsky, J. Gulick, J. Robbins. In vivo modeling desmin-related cardiomyopathies with transgenics. *Journal of Cardiac Failure* 2000; 5(Young Investigator Award 2nd prize at the 4th Annual Scientific Meeting of the Heart Failure Society of America, September 10-13, 2000, Boca Raton, FL.
8. Schwartzbauer, G.T., J.D. Molckintin, M.A. Sussman, X. Wang, J. Robbins. Differential protein expression among four models of cardiomyopathy identified by high throughput screening of cardiac polysome fractions. *Circulation* 2000; 102(18): II-33.
9. Wang X., H. Osinska, R. Klevitsky, A. Sambe, J. Robbins. R120G-alpha-B-crystallin causes desmin-related cardiomyopathy in transgenic mice. *Circulation* 2000; 102(18): II-201.
10. Wang X., R. Klevitsky, T.E. Hewett, A. M. Gerdes, T. R. Kimball, J. Robbins. Aberrant desmin aggregation disrupts desmin filament networks and leads to early concentric cardiac hypertrophy in transgenic mice. *Circulation* 2000; 102(18): II-290.
11. Wang X., H. Osinska, G.W. Dorn II, R. Klevitsky, J. Gulick, J. Robbins. A transgenic mouse model of desmin related cardiomyopathy. *Circulation* 2000; 102(18): II-218.
12. Wang X., R. Klevitsky, J. Robbins. J. Robbins. Loss of function of alpha B-crystallin enhances the pathogenesis of a desmin mutation. *FASEB J* 2001; 15(5): A1158.
13. Wang X, Klevitsky R, Robbins J. α B-Crystallin is essential to keeping desmin from aggregating adversely. *J Mol Cell Cardiol* 2001; 33(6): A129.
14. Hahn HS, Lin G, Wu G, Barrett T, Wang X, Robbins J, Lorenz JN, Mochly-Rosen D, Dorn II GW. α PKC Inhibition leading to a desmin related cardiomyopathy: analysis with mutant attenuated promoters. *Circulation* 2001; Publishing ID: 402
15. Wang X, Klevitsky R, Grupp IL, Robbins J. Cardiac α -B-crystallin Is protective against ischemia/reperfusion injury and essential in cardiac preconditioning: genetic studies. *Circulation* 2001; Publishing ID: 485
16. Wang X, Huang W, Andersen SM, Gerdes AM, Robbins J. Expression of R120G- α B-crystallin impairs proteolytic function of the proteasomes. *J Mol Cell Cardiol* 2002; 34(7):A15.
17. Saffitz JE, Green KG, Wang X, Robbins J. Down-regulation of intercellular junction proteins in a mouse model of desmin-related cardiomyopathy. *Circulation* 2002; 106(19):II-206.
18. Dong X, Huang W, Glasford JW, Harden NR, Li F, Gerdes AM, Wang X. Dynamic monitoring of the proteolytic function of the ubiquitin-proteasome system in cultured cardiomyocytes. *J Mol Cell Cardiol* 2003; 35 (6):A50. The 25th Annual Meeting ISHR-North American Section, 6/28-7/1/2003, Mystic, CT.

19. Huang W, Glasford JW, Harden NR, Dong X, Gerdes AM, Robbins J, Wang X. Remodeling of the intercalated disks in desmin-related cardiomyopathy caused by an alpha B-crystallin mutation in mice. *J Card Fail* 2003; 9(5): S27.
20. Chen QH, Huang W, Dong X, Glasford JW, Li F, Gerdes AM, Robbins J, Wang X. Modulation of the ubiquitin-proteasome system by R120G- α B-crystallin in vivo and in vitro. *Circulation* 2003; 108(17): IV-177.
21. Wang X, Kumarapeli AR, Glasford J, Liu J, Chen QH, Horak K, Zheng H, Dong X. In vivo monitoring dynamic changes in the proteolytic function of the ubiquitin-proteasome system. (A late-breaking abstract presented in 1st Annual Symposium of the AHA Council on Basic Cardiovascular Sciences. July 14-18, 2004, Stevenson, WA.)
22. Liu J, Chen QH, Wang X. Impairment of the ubiquitin-proteasome system in cardiac myocytes by reactive oxygen species revealed by a surrogate substrate. *Circ Res* 2004, 94(12):data supplement on abstracts of 1st Annual Symposium of the AHA Council on Basic Cardiovascular Sciences. July 14-18, 2004, Stevenson, WA.
23. Kumarapeli AR, Liu J, Glasford JW, Chen QH, Horak K, Dong X, Wang X. Myocardial ischemia/reperfusion impairs the ubiquitin-proteasome system: role of reactive oxygen species. *Circulation* 2004; 110(17): III-237. Oral presentation at AHA Scientific Sessions 2004, New Orleans, LA, November 7-10, 2004.
24. Wang X, Liu J, Chen QH, Dong X, Wawrousek EF, Li F, Horak K. Impairment of the ubiquitin-proteasome system by misfolded cytoplasmic protein in cardiac myocytes. *Circulation* 2004; 110 (17):III-160. Oral presentation at AHA Scientific Sessions 2004, New Orleans, LA, November 7-10, 2004.
25. Wang X, Chen Q, Liu J, Kumarapeli ARK, Zheng H, Horak K, Glasford JW, Gerdes AM, Li F. A novel functional indicator mouse model reveals impairment of the ubiquitin-proteasome system by misfolded proteins in the heart. The Inaugural Annual Symposium on Protein Folding Disorders. San Diego, CA, Jan 11-13, 2005.
26. Liu J, Tang M, Chen Q, Wang X. H₂O₂ inhibits proteolytic function of the ubiquitin-proteasome system by activating caspases. *Experimental Biology/IUPS 2005 Late Breaking Abstracts*, LB15. San Diego, CA, March 31-April 5, 2005.
27. Liu J, Chen Q, Huang W, Horak K, Zheng H, Mestrlil R, Wang X. Aberrant protein aggregation impairs the ubiquitin-proteasome system by affecting the entry of ubiquitinated proteins into the 20S proteasomes: A novel pathogenic pathway in cardiac remodeling and failure. Second Annual Symposium of the AHA Council on Basic Cardiovascular Sciences. July 24-27, 2005, Keystone, CO.
28. Li F, Zhou J, Yi XP, Graber, K, Qu J, Huber L, Wang X. Remodeling of the intercalated disk in cardiac specific β -catenin knockout mice. Second Annual Symposium of the AHA Council on Basic Cardiovascular Sciences. July 24-27, 2005, Keystone, CO.
29. Zhou J, Yi XP, Huber L, Qu J, Graber K, Wang X, Gerdes AM, Li F. Serine phosphorylation and nuclear redistribution of FAK and FRANK in cardiac myocytes.

Second Annual Symposium of the AHA Council on Basic Cardiovascular Sciences.
July 24-27, 2005, Keystone, CO.

30. Liu J-B, Chen Q, Wang X. Aberrant protein aggregation impairs protein degradation in mouse heart. AHA Research Symposium, November 12, 2005, Dallas, TX.
31. Chen Q, Liu J-B, Horak KM, Zheng H, Li J, Tang M, Su H, Kumarapeli AR, Li F, Gerdes AM, Wang X. Cytoplasmic aberrant protein aggregation impairs protein degradation in the heart by compromising the entry of ubiquitinated proteins into the 20S proteasomes. *Circulation* 2005 Oct 25; 112(17): II-188. AHA Scientific Sessions, November 12-16, 2005, Dallas, TX.
32. Zheng H, Horak KM, Robbins J, Wang X. Genetic inhibition of 20S proteasome in the heart. Late-breaking abstract in *Experimental Biology* 2006, April 1-5, 2006, San Francisco, CA
33. Kumarapeli ARK, Wang X. Alpha B-crystallin modulates pressure overload cardiac hypertrophy. Late-breaking abstract in *Experimental Biology* 2006, April 1-5, 2006, San Francisco, CA
34. Zheng H, Su H, Horak KM, Wang X. The ubiquitin-proteasome system in cardiac remodeling and failure. *J Mol Cell Cardiol* 2006; 41(4):748. The 28th Meeting of the North American Section of ISHR, June 12-15 2006, Toronto, Canada.
35. Su H, Menon S, Horak KM, Li J, Li F, Wei N, Wang X. Postnatal cardiomyocyte-restricted knockout of a COP9 signalosome gene compromises proteolytic function of the ubiquitin proteasome system and causes congestive heart failure in mice. Late-breaking basic science abstracts of the AHA Scientific Sessions 2006, November 12-15, 2006, Chicago, IL.
36. Liu JB, Zheng H, Tang M, Wang X. Doxorubicin activates ubiquitin-proteasome system mediated proteolysis by acting on both ubiquitination apparatuses and the proteasome. *FESAB J* 2007:A1024; Abstract 808.4 in the Abstract Book PART II of *Experimental Biology* 2007, April 28-May 2, Washington, DC.
37. Su H, Li JB, Osinska H, Menon S, Li F, Robbins J, Wei N, Wang X. Genetic inhibition of cullin based ubiquitin ligase dynamics in adult mouse hearts suffices to cause heart failure. *FESAB J* 2007: A870; Abstract 747.11 in the Abstract Book PART II of *Experimental Biology* 2007, April 28-May 2, Washington, DC.
38. Lei D-X, Li F, Menon S, Su H, Horak KM, Wei N, Wang X. Ablation of a COP9 signalosome gene in liver causes cirrhosis and dysplastic changes. 2007 Abstract Book of Keystone Symposium on Ubiquitin and Signal, Feb 4-9, 2007, Big Sky, Montana.
39. Zheng Q, Mizushima N, Wang X. Proteasome malfunction increases autophagosomes in cardiomyocytes and mouse hearts. *FASEB J*. 2008 22:605.8 *Experimental Biology Meeting* 2008

40. Zheng H, Li M, Horak KM, Huang W, Sanbe A, Robbins J, Li F, Wang X. Moderate cardiomyocyte-restricted proteasome inhibition exacerbates diastolic malfunction in desmin-related cardiomyopathy mice. *J Mol Cell Cardiol* 2008; 44:S50.
41. Su H, Li J, Osinska H, Menon S, Horak KM, Li F, Molkenin JD, Robbins J, Wei N, Wang X. Cardiac ablation of CSN8 perturbs proteasomal proteolysis, activates autophagy, and causes fatal heart failure in adult mice. *J Mol Cell Cardiol* 2008; 44:S51.
42. Li J, Horak KM, Sanbe A, Jeffrey Robbins J, Wang X. A transgenic mouse model of benign enhancement of cardiac proteasomal function. *J Mol Cell Cardiol* 2008; 44:S53.
43. Ryu Y-C, Wang X. Hyper-phosphorylation of proteasome subunits increases proteasome peptidase activities in desminopathy mouse hearts. *J Mol Cell Cardiol* 2008; 44:S60.
44. Huabo Su, Jie Li, Suchithra Menon, Kathleen Horak, Faqian Li, Ning Wei, Xuejun Wang. Perinatal Cardiomyocyte-Restricted Knockout of the COP9 Signalosome Subunit 8 Gene Compromises Proteasome Proteolytic Function, Triggers Necrosis, and Causes Heart Failure in Mice. *Circ Res* 2008; 103:e35-e70.
45. Kumarapeli AR, Tang M, Zheng H, Horak KM, Li M, Molkenin JD, Wang X. Protein Quality Control Inadequacy Activates NFAT Signaling in Cardiomyocytes. *Circulation* 2008; 118: S_393. Oral presentation, AHA Scientific Sessions 2008; Nov 8-12, New Orleans, LA.
46. Hanqiao Zheng; Mingxin Tang; Asangi R. K. Kumarapeli; Kathleen M. Horak; Xuejun Wang. Doxycycline Attenuates Protein Aggregation in Cardiomyocytes and Improves Survival of a Mouse Model of Cardiac Conformational Disease. Poster presentation at ISHR North American Section Meeting: New Discoveries for Prevention and Treatment of Heart Disease May 26-29, 2009 Baltimore, MD.
47. Ranek MJ, Wang X. Activation of muscarinic receptor 2 stimulates proteasome function in cardiomyocytes. *Experimental Biology Meeting* April 24-28, 2010 Anaheim, CA.
48. Ranek MJ, Wang X. Stimulation of adrenergic receptors regulates proteasome function in cardiomyocytes. *Experimental Biology Meeting* April 24-28, 2010 Anaheim, CA.
49. Davis F, Predmore J, Wang P, Li J, Su H, Converso K, Allen A, Jones R, Powell SR, Wang X, Day SM. Activation of the ubiquitin proteasome system after myocardial infarction. Presented at the AHA Scientific Sessions, November 2010, Chicago, IL. *Circulation* 2010; 122(21): A18730.
50. Li J, Horak KM, Su H, Sanbe A, Robbins J, Wang X. Enhancement of Proteasomal Function Protects Against Proteinopathy and Myocardial Ischemia-Reperfusion Injury in Mice. AHA Basic Cardiovascular Sciences 2011 Scientific Sessions, New Orleans, LA July 18-21, 2011
51. Tian Z, Zheng H, Wang X. Genetically Induced Moderate Inhibition of the Proteasome in Cardiomyocytes Exacerbates Myocardial Ischemia-Reperfusion Injury in Mice (Poster 271). AHA Basic Cardiovascular Sciences 2011 Scientific Sessions, New Orleans, LA July 18-21, 2011

52. Wang C, Tian Z, Zheng Q, Su H, Li J, Wang X. Interplay Between the Ubiquitin-Proteasome System and Autophagy in the Heart. AHA Basic Cardiovascular Sciences 2011 Scientific Sessions, New Orleans, LA July 18-21, 2011
53. Su H, Tian Z, Wang C, Said S, Ranek MJ, Wang X. Impaired Autophagosome Removal in Cardiomyocytes Triggers Programmed Necrosis in Mouse Hearts. AHA Basic Cardiovascular Sciences 2011 Scientific Sessions, New Orleans, LA July 18-21, 2011
54. Zheng Q, Su H, Ranek MJ, Wang X. The Role of p62 in Cardiac Protein Quality Control. AHA Basic Cardiovascular Sciences 2011 Scientific Sessions, New Orleans, LA July 18-21, 2011
55. Su H, Li F, Ranek MJ, Wei N, Wang X. The COP9 Signalosome Regulates Autophagy. AHA Basic Cardiovascular Sciences 2011 Scientific Sessions, New Orleans, LA July 18-21, 2011
56. Ranek MJ, Wang X. Protein Kinase G Regulates the UPS in Cardiomyocytes. Presented at the Protein Degradation Pathways in Health and Diseases, San Diego, CA, USA, January 2012.
57. Ranek MJ, Martin D, Kost C, Wang X. Activation of Protein Kinase G Enhances Proteasome-Mediated Degradation of Misfolded Proteins. Presented at the AHA Basic Cardiovascular Sciences Scientific Sessions, New Orleans, LA, July 22-24, 2012.
58. Ranek MJ, Terpstra EJM, Li J, Wang X. Protein kinase G regulates proteasome-mediated degradation of misfolded proteins. the AHA Scientific Sessions, Los Angeles, CA. November 3-8, 2012
59. Su H., Li J., Wei N., Wang X. The COP9 signalosome subunit 8 hypomorphism impairs deneddylation and exacerbates desmin-related cardiomyopathy. Poster presentation at Experimental Biology 2013, April 20-24, Boston, MA, USA.
60. Ranek MJ, Terpstra EJM, Li J, Kass DA, Wang X. Protein kinase G positively regulates proteasome-mediated degradation of misfolded proteins. Presented at the AHA Basic Cardiovascular Sciences Scientific Sessions, Las Vegas, Nevada, July 22-25, 2013.
61. Terpstra EJM, Ranek MJ, Callegari E, Wang X. The Proteasome is a Target of Protein Kinase G. Presented at the AHA Scientific Sessions, Dallas TX, November 16-20, 2013.
62. Day S M, Yob J, Davis F, Wang P, Converso K L, Wang X, Powell, S R. Selective Inhibition of the Immunoproteasome Attenuates Adverse Left Ventricular Remodeling, Improves Cardiac Function, and Prevents Heart Failure After Myocardial Infarction. Presented at the AHA Scientific Sessions, Dallas TX, November 16-20, 2013.
63. Ranek MJ, Terpstra EJM, Li J, Kass DA, Wang X. Protein kinase G positively regulates proteasome-mediated degradation of misfolded proteins. Presented at the AHA Scientific Sessions, Dallas TX, November 16-20, 2013.

64. Tian Z, Wang C, Hu C, Tian Y, Liu J, Wang X. Autophagic-Lysosomal Inhibition Compromises Ubiquitin-1 Proteasome System Performance in a p62 Dependent Manner in Cardiomyocytes. Presented at the 35th International Society for Heart Research North American Section Meeting: "NOVEL STRATEGIES TO COMBAT HEART FAILURE", Miami Beach, Florida, May 12 -15, 2014
65. Hu, C., Wang, X., Wang, H., Tian, Y. Inadequate Coupling between Ubiquitination and the Proteasome is a Major Pathogenic Factor of Myocardial Ischemia/Reperfusion Injury. Presented at the The First Annual ACRE/APS Symposium, Las Vegas, NV. (2014, July 13).
66. Wang, X., Hu, C., Tian, Y., Wang, H. Inadequate Coupling between Ubiquitination and the Proteasome is a Major Pathogenic Factor of Myocardial Ischemia/Reperfusion Injury. Presented at the Basic Cardiovascular Sciences 2014 Scientific Sessions: Pathway to Cardiovascular Therapeutics, Las Vegas, NV. (2014, July 16).
67. Wang, C., Jahn, A., Su, H., Wang, X. Duo-impairment of the Ubiquitin-Proteasome System and Autophagy by Ablation of COP9 Signalosome Subunit 8 Activates a Programmed Necrosis Pathway Mediated by RIP1-RIP3 Kinases but not Cyclophilin D-regulated Mitochondrial Membrane Permeability . Presented at the Basic Cardiovascular Sciences 2014 Scientific Sessions: Pathway to Cardiovascular Therapeutics, Las Vegas, NV. (2014, July 16).
68. Su, H., Li, J., Zhang, H., Wei, N., Wang, X. The COP9 signalosome controls the degradation of cytosolic misfolded proteins and protects against cardiac proteotoxicity. Presented at the 36th International Society for Heart Research North American Section (ISHR-NAS) Meeting, Seattle, WA. (June 8, 2015)
69. Li, J., Ma, W., Li, H., Hou, N. Wang, X., Kim I-M., Li, F., Su, H. NEDD8 ultimate buster-1 long (NUB1L) protein regulates atypical neddylation and protects against myocardial ischemia-reperfusion injury. Presented at the AHA Basic Cardiovascular Sciences 2015 Scientific Sessions: Pathway to Cardiovascular Therapeutics, New Orleans, LA. July 13-16, 2015.
70. Wang, X., Terpstra, E.J., Callegari, E., Hu, C., Zhang, H. Wang, X. Proteasome priming by protein kinase G protects against myocardial ischemia-reperfusion injury. Presented at the AHA Basic Cardiovascular Sciences 2015 Scientific Sessions: Pathway to Cardiovascular Therapeutics, New Orleans, LA. July 13-16, 2015.
71. Wang, X., Wang, C., Terpstra, E.J., Wang, Y., Wang, X. Activation of the p38 branch of mitogen activated protein kinase pathway stimulates proteasome proteolytic function. Presented at the AHA Basic Cardiovascular Sciences 2015 Scientific Sessions: Pathway to Cardiovascular Therapeutics, New Orleans, LA. July 13-16, 2015.
72. Wang C, Li J, Su H, Xiao P, Wang X. COPS8 inhibits cardiomyocyte necroptosis in mouse hearts via suppressing the RIPK1-RIPK3 pathway. Presented at American Heart Association (AHA) Scientific Sessions, Orlando, FL, November 9; 2015.
73. Zhang, Hanming, Wang, Xuejun. PDE1 inhibition improves cardiac protein quality control. Presented at the 3rd ACRE/APS symposium, Phoenix, AZ. July 17, 2016.

74. Wu, Penglong, Zhu, Wei, Li, Jinbo, Wang, Xuejun. Cathepsin D haploinsufficiency exacerbates post-MI cardiac remodeling and malfunction by impairing autophagosome removal. Presented at the 3rd ACRE/APS symposium, Phoenix, AZ. July 17, 2016.
75. Zhang, Hanming, Wang, Xuejun. PDE1 inhibition primes the proteasome. Presented at the AHA Basic Cardiovascular Sciences (BCVS) 2016 Scientific Sessions: Pathways to Cardiovascular Therapeutics, Phoenix, AZ. July 18, 2016.
76. Wu, Penglong, Zhu, Wei, Li, Jinbo, **Wang, Xuejun**. Cathepsin D haploinsufficiency exacerbates post-MI cardiac remodeling and malfunction by impairing autophagosome removal. Presented at the AHA Basic Cardiovascular Sciences 2016 Scientific Sessions: Pathways to Cardiovascular Therapeutics, Phoenix, AZ, July 19, 2016.
77. Abdullah, Ammara; Eyeter, Kathleen M; Bjordahl, Travis; Xiao, Peng; Zeng, Erliang; **Wang, Xuejun**. Cardiac transcriptome analysis reveals a critical role for the COP9 signalosome in transcriptional regulation of the substrate receptors of cullin-RING ligases in mice. Presented at The 12th International Conference on Pathways, Networks, and Systems Medicine, Aegean Conference, Crete, Greece, June 29 ~ Jul 4, 2017.
78. **Zhang, Hanming**; Wang, Xuejun. Inhibition of phosphodiesterase 1 confers striking therapeutic benefit to HFpEF in mice. Presented at the AHA Basic Cardiovascular Sciences 2017 Scientific Sessions. (New Investigator Travel Award), Portland, OR, July 10-13, 2017.
79. **Zhang, Hanming**, Rekhter, Mark D., Wang, Xuejun. Inhibition of Type 1 Phosphodiesterase Confers Therapeutic Benefit to Proteinopathy-based HFpEF in Mice. Presented at the 2018 Experimental Biology Meeting, San Diego, CA. (April 25, 2018)
80. **Wu, Penglong**, Wang, Yibin, Wang, Xuejun. EXCESSIVE BETA-ADRENERGIC RECEPTOR STIMULATION INDUCES CARDIOMYOCYTE NECROPTOSIS VIA A RIP3-DEPENDENT PATHWAY. Presented at the 2018 Experimental Biology Meeting, San Diego, CA. (April 22, 2018)
81. **Wang, Xuejun**, Wu, Penglong, Parajuli, Nirmal, Pan, Bo, Lewno, Megan, Liu, Jinbao. Proteasome phosphorylation and activation by PKA protects against cardiac remodeling in mice subjected to myocardial infarction. Presented at the 2019 Experimental Biology meeting, Orlando, FL. (April 9, 2019)
82. **Wu, Penglong**, Wang, Yibin, Liu, Jinbao, Wang, Xuejun. Necroptosis Resulting from Activation of a RIP3-dependent Pathway Contributes to Cardiomyocyte Death Induced by Isoproterenol. Presented at the 2019 Experimental Biology meeting, Orlando, FL. (April 8, 2019)
83. **Wu, Penglong**, Li, Yi-Fan, Liu, Jinbao, Wang, Xuejun. Post-MI Cardiac Remodeling and Malfunction in Mice Are Exacerbated by Cardiomyocyte-restricted Ablation of the Uchl1 Gene. Presented at the 2019 Experimental Biology meeting, Orlando, FL. (April 7, 2019)
84. Penglong Wu, Bo Pan, Megan Lewno, Nirmal Parajuli, **Xuejun Wang**. In vivo genetic interrogations establish unequivocally the pathophysiological significance of proteasome phosphoregulation by protein kinase A. *J Mol Cell Cardiol*. March 2020;

140:6. DOI: <https://doi.org/10.1016/j.yjmcc.2019.11.010> Presented at the 23rd World Congress of International Society for Heart Research (ISHR) held in Beijing, China (June 4, 2019).

85. Hanming Zhang, Bo Pan, Penglong Wu, Nirmal Parajuli, Mark D. Rekhter, Alfred L Goldberg, **Xuejun Wang**. Dual Activation of PKA and PKG by PDE1 Inhibition Facilitates Proteasomal Degradation of Misfolded Proteins and Protects Against Proteinopathy-Based HFpEF. Oral abstract presentation at Session 1A “HFpEF: Unraveling the Gordian Knot” of 2019 AHA BCVS Scientific Sessions (July 29, 2019; Boston, MA).
86. Penglong Wu, Nirmal Parajuli, Megan Lewno, Jinbao Liu, **Xuejun Wang**. Proteasome priming by cyclic AMP signaling protects stressed hearts in mice. Presented at American Heart Association (AHA) Scientific Sessions, Philadelphia, PA, November 16-18, 2019.
87. Hanming Zhang, Bo Pan, Penglong Wu, Nirmal Parajuli, Mark D. Rekhter, Alfred L Goldberg, **Xuejun Wang**. Dual Activation of PKA and PKG by PDE1 Inhibition Facilitates Proteasomal Degradation of Misfolded Proteins and Protects Against Proteinopathy-Based HFpEF. Presented at the Session entitled “Best of AHA Specialty Conferences: BCVS 2019 of 2019”, AHA Scientific Sessions, Philadelphia, PA, November 17, 2019.
88. **Penglong Wu**, Nirmal Parajuli, Megan Lewno, Liuqing Yang, Jinbao Liu, Xuejun Wang. RPN6-Ser14 Phosphorylation Is Responsible for Proteasome Activation by PKA and Protects against Pathological Cardiac Hypertrophy and Malfunction in Mice. *FASEB J* 17 April 2020; 34(S1): 03399. <https://doi.org/10.1096/fasebj.2020.34.s1.03399> (with this work, Dr. Wu won the Runner-up of APS-Cardiovascular Section Outstanding Postdoctoral Trainee Awards).
89. **Megan Lewno**, Xuejun Wang. Phenotypic Differences Among Mice with Induced Cardiomyocyte-Restricted Ablation of Cops5, Cops8, or Both. Presented at *Iowa Physiological Society (IPS) and Midlands Society of Physiological Sciences (MSPS) Scientific Sessions 2020 (virtual)*, October 30-31, 2020.
90. **Samiksha Giri**, Chao Suo, Megan T. Lewno, Douglas S. Martin, Xuejun Wang. Defining molecular mechanism promoting neointimal hyperplasia by CSN8 hypomorphism. Presented at *Iowa Physiological Society (IPS) and Midlands Society of Physiological Sciences (MSPS) Scientific Sessions 2020 (virtual)*, October 30-31, 2020.
91. **Liuqing Yang**, Nirmal Parajuli, Jack O. Sternburg, Xuejun Wang. Ser14-Psm11/Rpn6 phosphorylation is required for activation of the 26S proteasome by PKA but is dispensable for cardiac responses to increased proteotoxic stress. Presented at *Iowa Physiological Society (IPS) and Midlands Society of Physiological Sciences (MSPS) Scientific Sessions 2020 (virtual)*, October 30-31, 2020.
92. **Mingqi Cai**, Xuejun Wang. Soluble guanylate cyclase activation increases proteasome activities and facilitates degradation of misfolded proteins in cardiomyocytes. Presented at *Iowa Physiological Society (IPS) and Midlands Society of Physiological Sciences (MSPS) Scientific Sessions 2020 (virtual)*, October 30-31, 2020.

93. **Samiksha Giri**, Chao Suo, Douglas S. Martin, Xuejun Wang. Defining Molecular Mechanism Promoting Neointimal Hyperplasia by CSN8 Hypomorphism. *FASEB J.* 14 May 2021; 35(S1):04143. <https://doi.org/10.1096/fasebj.2021.35.S1.04143> (with this work, PhD student Samiksha Giri won an APS-Cardiovascular Section Research Recognition Award).
94. **Megan Lewno**, Xuejun Wang. Phenotypic Differences Among Mice with Induced Cardiomyocyte-Restricted Ablation of Cops5, Cops8, or Both. *FASEB J.* 14 May 2021; 35(S1):05216. <https://doi.org/10.1096/fasebj.2021.35.S1.05216>
95. **Mingqi Cai**, Xuejun Wang. Soluble guanylate cyclase activation increases proteasome activities and protects against proteotoxicity in cardiomyocytes. *FASEB J.* 14 May 2021; 35(S1): 05087. <https://doi.org/10.1096/fasebj.2021.35.S1.05087>

GRANT/CONTRACT SUPPORT

Current Grant Support

- | | | |
|---|-------------------------|---|
| 1 R01 HL153614-02
NIH/NHLBI | Wang | 8/1/20-6/30/24
\$250,000 |
| Cardiac Pathophysiology of Proteasome Phosphoregulation
The goal of this project is to determine in vivo (patho)physiological significance of phosphoregulation of the 26S proteasome by PKA.
Role: PI | | |
| 2 R01 HL072166-16
NIH/NHLBI | Wang | 7/1/03 – 6/30/24
\$343,921 |
| PKG and PKA Duo-Activation to Treat Cardiac Proteotoxicity
The goal of this project is to exploit proteasome activation by PKG and PKA duo-activation to treat cardiac proteinopathy.
Role: PI | | |
| R01 HL131667-04
NIH/NHLBI | Cui, Wang (Contact) | 4/1/17 - 3/31/22
(NCE) |
| The NRF2-p62 Axis in the Cross-Talk between Proteasomal and Lysosomal Degradation
The goal of this project is to test the role of the NRF2-p62 pathway in the impact of autophagy insufficiency on UPS performance.
Role: Duo-PI (contact PI) | | |
| 20TPA35490091
American Heart Association (AHA) | Wang | 1/1/21-12/31/23
\$90,909 |
| JAK1-STAT Pathway Promotes Cardiac Proteotoxicity
This AHA transformative project award will support a research project to investigate the role of the JAK1-STAT signaling pathway in the pathogenesis of cardiac proteinopathy.
Role: PI | | |
| 1 R41 HL152919-01A1
NIH/NHLBI | Shaffer (Contact); Wang | 2/16/21 -1/31/22
\$320,198 (\$251,774 to Wang lab) |

VAL-0914 Decreases PAO to Protect Against Cardiac Proteinopathies

The goal of this STTR project is to test a novel proprietary compound (VAL-0914) in a mouse model of cardiac proteinopathy.

Role: Duo-PI

Pending Grants:

1R01AG072510-01A1 Wang H (Contact), Wang X 4/1/2022 – 3/31/2027
NIH/NIA \$375,000

Priming the proteasome to protect against aging and Alzheimer's disease.

Role: Duo-PI

3 R01 HL153614-02S1 Wang 7/1/2022 – 6/30/2023
NIH/NIA \$250,000

AD Supplement to Cardiac Pathophysiology of Proteasome Phosphoregulation

Role: PI

3 P20GM103443-21S1 Goodman (PI)/Wang X, Wang H (PL) 9/1/2022 – 8/31/2023
NIH/NIA \$250,000

AD Supplement to SD BRIN

Role: one of the 2 Project Leaders for the AD Supplement

Previous Grant Support

AHA Postdoctoral Fellowship Wang (PI) 7/1/2000 – 6/30/2002
American Heart Association (AHA) Ohio Affiliate \$70,000 (total direct)
Dissecting Desmin-related Cardiomyopathy with Transgenesis
Role: PI
(Terminated upon assistant professor appointment on October 1, 2001)

AHA 0235099N Wang (PI) 7/1/2002 – 6/30/2007
American Heart Association (AHA) National Center SDG \$260,000 (total direct)
Ubiquitin-Proteasome System in the Pathogenesis of Crystallinopathy
Role: PI

1P20RR17662-019003 Wang (PI) 9/20/2002 – 6/30/2007
NIH/NCRR \$100,000 (annual direct)
Mechanisms of Cardiovascular Remodeling: Molecular Biology Core
Role: Core Director

1P20RR17662-010001 Wang (PI) 9/20/2002 – 6/30/2007
NIH/NCRR \$200,000 (annual direct)
Mechanisms of Cardiovascular Remodeling: Project 1-Ubiquitin-Proteasome System
Dysfunction in Cardiac Remodeling and Failure
Role: PI of Project 1.

Reference # 0740025N Wang (PI) 1/1/2007- 12/31/2012
AHA Established Investigator Award \$500,000 (total cost)
Inadequate Protein Quality Control in Heart Failure

Role: PI

R01HL068936-04~05 Powell (contact), Wang 12/1/2008-6/30/2014
 NIH/NHLBI \$280,000 (annual direct)
 Proteasome, protein oxidation, and cardiomyocyte function
 Role: Duo-PI, PI #2

AHA 0510069Z Kumarapeli (PI) 1/1/2005-12/31/2006
 AHA Predoctoral Fellowship \$52,000 (total cost)
 Alpha B-crystallin modulates cardiac hypertrophic response to mechanical overload.
 Role: Sponsor and Primary Mentor

AHA 0620032Z H. Zheng (PI) 1/1/2006-12/31/2007
 AHA Postdoctoral Fellowship Grant \$100,000 (total cost)
 Genetic Inhibition of Proteasomal Function in the Heart.
 Role: Sponsor and Primary Mentor

Postdoctoral fellowship Su (PI) 7/1/2006 – 6/30/2008
 AHA Greater Midwest Affiliate \$100,000 (total cost)
 Physiological Significance of COP9 Signalosome in Adult Hearts.
 Role: Sponsor and Primary Mentor

Reference # 0815571G Q. Zheng (PI) 7/1/2008-6/30/2010
 AHA Predoctoral Fellowship \$52,000 (total cost)
 Autophagy is activated by and compensates for proteasome malfunction in desminopathy
 Role: Sponsor and Primary Mentor

11PRE5730009 Ranek (PI) 1/1/2011-12/31/2012
 AHA Predoctoral Fellowship \$52,000 (total cost)
 The Role of PKG in the Stimulation of the UPS by the Muscarinic 2 Receptor
 Role: Sponsor and Primary Mentor

11SDG6960011 Su (PI) 07/01/2011-06/30/2015
 AHA Scientist Development Grant (National Center) \$280,000 (total direct)
 The Ubiquitin Proteasome System in Diabetic Cardiomyopathy
 Role: Collaborator and Mentor

1 F32 HL122045-01 Terpstra (PI) 07/01/2014 – 6/30/2017
 NIH National Research Service Award (Postdoc Fellowship) \$162,000 (total direct)
 Direct proteasomal enhancement contributes to PKG-triggered cardioprotection
 Role: Sponsor and Primary Mentor;
(Terminated on 8/2015 upon the PI's matriculation to medical school)

16UFEL29640003 Reihe (PI) 06/01/2016 – 8/31/2016
 AHA Undergraduate Student Research Program \$4000 (scholarship)
 Effect of Neddylation Inhibition on Autophagic Flux in Cardiomyocytes
 Role: Sponsor and Primary Mentor

16PRE27790059 Zhang (PI) Wang (Sponsor) 01/01/2016 – 12/31/2017

AHA Predoctoral Fellowship PDE1 inhibition improves cardiac protein quality control Role: Sponsor and Primary Mentor		\$52,000 (total direct)
R01 HL085629-01~13 NIH/NHLBI The COP9 Signalosome in the Heart Role: PI	Wang	7/1/06 - 6/30/21 \$4,684,110 (total cost)

CONTRIBUTION TO SCIENCE

- Discovery of UPS functional insufficiency in mouse models of heart disease using innovative tools generated in my lab. UPS-mediated protein degradation, initially discovered in 1980s, had not begun to attract a broader spectrum of cell biologists until mid-1990s. The attention to the UPS from general biomedical scientists outside the cell biology field was likely drawn by the award of a 2004 Nobel Prize to 3 scientists for their contributions to the discovery of the UPS. As a pioneer for studying UPS dysfunction in cardiac pathogenesis, my lab has been working on the UPS in the heart since 2001. By then, research into health and disease of most organs/systems, including the heart, had emphasized primarily changes in gene expression at the transcription or, in some cases, protein synthesis, although changes in protein degradation could have equal or even greater impact on the level and functioning of a protein. A major hurdle then was lacking tools to monitor UPS function in vivo. We first developed stable cell lines, adenoviruses, and stable transgenic (tg) mouse lines expressing a modified GFP (GFPu or GFPdgn) that is a proven surrogate substrate of the UPS, allowing monitoring the dynamics of UPS performance in situ and in vivo. These new tools have been distributed to researchers around the world and also enabled my lab to demonstrate in intact animals for the first time in the world that increases in misfolded proteins and resultant aberrant protein aggregation impair UPS proteolytic function and cause proteasome functional insufficiency (PFI). Similarly, we were also the first to document cardiac UPS functional insufficiency in acute ischemia/reperfusion (I/R) injury, chronic pressure overload, and diabetic cardiomyopathy. These publications have prompted investigations into cardiac pathogenic role of UPS dysfunction.

 - Chen Q, Liu JB, Horak KM, Zheng H, Kumarapeli AR, Li J, Li F, Gerdes AM, Wawrousek EF, **Wang X**. Intrascoplasmic amyloidosis impairs proteolytic function of proteasomes in cardiomyocytes by compromising substrate uptake. *Circ Res*. 2005; 97(10):1018-26. PMID: [16210548](#).
 - Kumarapeli AR, Horak KM, Glasford JW, Li J, Chen Q, Liu J, Zheng H, **Wang X**. A novel transgenic mouse model reveals deregulation of the ubiquitin-proteasome system in the heart by doxorubicin. *FASEB J*. 2005; 19(14):2051-3. PMID: [16188962](#).
 - Liu J, Chen Q, Huang W, Horak KM, Zheng H, Mestril R, **Wang X**. Impairment of the ubiquitin-proteasome system in desminopathy mouse hearts. *FASEB J*. 2006; 20(2):362-4. PMID: [16371426](#).
 - Li J, Ma W, Yue G, Tang Y, Kim IM, Weintraub NL, **Wang X**, Su H. Cardiac proteasome functional insufficiency plays a pathogenic role in diabetic cardiomyopathy. *J Mol Cell Cardiol*. 2017; 102:53-60. PMID: [PMC5316366](#).
- Demonstration of the necessity of proteasome functional insufficiency (PFI) and inadequate ubiquitination-proteasome coupling in pathogenesis. We have established that PFI plays an essential pathogenic role in proteinopathy and I/R injury, as well as pressure overload right heart

failure and diabetic cardiomyopathy (by collaboration). Although UPS malfunction had been proposed first by neuroscientists to play a role in neurodegeneration the necessity of PFI in pathogenesis was not demonstrated until 2011 when we published that genetic enhancement of the proteasome protects against proteinopathy and I/R injury in mice. Our discovery that PA28 α overexpression enhances proteasome function, which identifies the 1st measure to achieve proteasome gain-of-function, allowing us to generate the first animal model with forced proteasome function enhancement. We also created the first cardiomyocyte-restricted proteasome inhibition (CR-PsmI) and ubiquitination-proteasome uncoupling mouse models. These unique genetic models are essential to, and have remarkably facilitated, defining the pathophysiological significance of cardiac UPS malfunction. Using these tools, we have established a major pathogenic role for PFI in I/R injury, pressure overloaded cardiac maladaptive remodeling, and diabetic cardiomyopathy, three major causes of heart failure. These studies also strongly indicative of improving proteasome function as a potentially novel therapeutic strategy for a large subset of heart diseases, providing compelling rationale for studies on the regulation of the UPS in the heart.

- a. Hu C, Tian Y, Xu H, Pan B, Terpstra EM, Wu P, Wang H, Li F, Liu J, **Wang X**. Inadequate ubiquitination-proteasome coupling contributes to myocardial ischemia-reperfusion injury. *J Clin Invest*. 2018; 128(12):5294-06. PMCID: [PMC6264645](#).
- b. Li J, Ma W, Yue G, Tang Y, Kim IM, Weintraub NL, **Wang X**, Su H. Cardiac proteasome functional insufficiency plays a pathogenic role in diabetic cardiomyopathy. *J Mol Cell Cardiol*. 2017; 102:53-60. PMCID: [PMC5316366](#).
- c. Tian Z, Zheng H, Li J, Li Y, Su H, **Wang X**. Genetically induced moderate inhibition of the proteasome in cardiomyocytes exacerbates myocardial ischemia-reperfusion injury in mice. *Circ Res*. 2012; 111(5): 532-42. PMCID: [PMC3426260](#).
- d. Li J, Horak KM, Su H, Sanbe A, Robbins J, **Wang X**. Enhancement of proteasomal function protects against cardiac proteinopathy and ischemia/reperfusion injury in mice. *J Clin Invest*. 2011; 121(9): 3689-700. PMCID: [PMC3163952](#).

3. Discovery of novel mechanisms by which proteasome function is regulated and identification of new strategies to prime or activate the proteasome. We discovered that cGMP-dependent kinase (PKG) positively regulate proteasome function in cardiomyocytes, PKG activation by either genetic or pharmacological (e.g., PDE5 inhibition) means promotes proteasome-dependent degradation of a surrogate and a bona fide misfolded protein in cardiomyocytes, and PDE5 inhibition by sildenafil reduces misfolded protein abundance and aggregation and slows down disease progression in a well-established mouse model of cardiac proteinopathy, which represents the first demonstration that proteasome function can be pharmacologically enhanced to treat disease. Our discovery that muscarinic receptor 2 activation enhances cardiac proteasomal function in a PKG dependent manner was the first to unveil the physiological requirement of kinase-elicited proteasome activation. More recently, we demonstrated duo-activation of PKA and PKG by PDE1 inhibition confers striking therapeutic benefit in a mouse model of proteinopathy-based HFpEF. These findings also demonstrate the feasibility to use pharmacological method to enhance proteasomal degradation of misfolded proteins and thereby improve PQC in the heart.

- a. Huang H, Zhang X, Li S, Liu N, Lian W, McDowell E, Zhou P, Zhao C, Guo H, Zhang C, Yang C, Wen G, Dong X, Lu L, Ma N, Dong W, Dou QP, **Wang X**, Liu J. Physiological levels of ATP negatively regulate proteasome function. *Cell Res*. 2010; 20(12):1372-85. PMCID: [PMC2996470](#).
- b. Ranek MJ, Terpstra EJ, Li J, Kass DA, **Wang X**. Protein kinase g positively regulates proteasome-mediated degradation of misfolded proteins. *Circulation*. 2013; 128(4):365-76. PMCID: [PMC3761383](#).

- c. Zhang H, Pan B, Wu P, Parajuli N, Rekhter MD, Goldberg AL, **Wang X**. PDE1 inhibition facilitates proteasomal degradation of misfolded proteins and protects against cardiac proteinopathy. *Sci Adv*. 2019; 5(5):eaaw5870. PMID: [PMC6531002](https://pubmed.ncbi.nlm.nih.gov/31111111/).
 - d. **Wang X** and Wang H. Priming the proteasome to protect against proteotoxicity. *Trends in Molecular Medicine*. 25 March 2020 (Online First); DOI:<https://doi.org/10.1016/j.molmed.2020.02.007>. (PMC in progress).
4. Elucidation of physiological roles of the COP9 signalosome (CSN) in the heart and liver. Ub ligases or E3s confer substrate specificity and catalyze the Ub ligation to substrates, the final and rate-limiting step in ubiquitination. The cullin-RING ligases (CRLs) are the largest family of Ub E3s, known to regulate a variety of cellular processes including cell cycle control. CRLs are activated by cullin neddylation, inhibition of which via MLN4924 is in clinical trials for treating cancers. The CSN holo-complex consisting of 8 unique protein subunits (COPS1 thru COPS8) functions as a deneddylase for cullins, indispensable to CRLs catalytic dynamics. The CSN was rarely studied for its role in a terminally differentiated organ of vertebrates. Through cardiac targeting the Cops8 gene, we demonstrate that COPS8/CSN is required for cardiomyocyte survival and functioning, COPS8/CSN regulates not only the UPS but also the autophagic-lysosomal system (ALP), the latter was not known before our discovery that COPS8/CSN is required for autophagosome fusion with lysosomes. More recently, our studies reveal for the first time that CRLs contribute to degradation of misfolded cytosolic proteins and that Cops8/CSN suppresses the RIPK1-RIPK3 mediated cardiomyocyte necroptosis in mice.
- a. Xiao P, Wang C, Li J, Su H, Yang L, Wu P, Lewno MT, Liu J, Wang X. COP9 signalosome suppresses RIPK1-RIPK3-mediated cardiomyocyte necroptosis in mice. *Circ Heart Fail* 2020; 13(8): e006996. [PMCID: PMC7438278](https://pubmed.ncbi.nlm.nih.gov/32111111/).
 - b. Su H, Li J, Zhang H, Ma W, Wei N, Liu J, **Wang X**. COP9 signalosome controls the degradation of cytosolic misfolded proteins and protects against cardiac proteotoxicity. *Circ Res*. 2015; 117(11):956-66. PMID: [PMC4636927](https://pubmed.ncbi.nlm.nih.gov/26111111/).
 - c. Su H, Li F, Ranek MJ, Wei N, **Wang X**. COP9 signalosome regulates autophagosome maturation. *Circulation*. 2011; 124(19):2117-28. PMID: [PMC3211066](https://pubmed.ncbi.nlm.nih.gov/21111111/).
 - d. Su H, Li J, Menon S, Liu J, Kumarapeli AR, Wei N, **Wang X**. Perturbation of cullin deneddylation via conditional Csn8 ablation impairs the ubiquitin-proteasome system and causes cardiomyocyte necrosis and dilated cardiomyopathy in mice. *Circ Res*. 2011; 108(1):40-50. PMID: [PMC3017673](https://pubmed.ncbi.nlm.nih.gov/20111111/).
5. Dissecting the crosstalk between cardiac UPS and ALP pathways in cardiac PQC. We have elucidated a pivotal role of the calcineurin-TFEB-p62/SQSTM1 pathway in this crosstalk. We have shown that proteasomal malfunction activates autophagy in the heart while ALP inhibition hinders the degradation of ubiquitinated proteins by the proteasome in a p62-dependent manner, contributing to the dark side of Nrf2 activation in both hypertensive and diabetic cardiomyopathies. We are the first to report the upregulation of p62 at both transcript and protein levels in proteinopathic hearts which are known to have UPS insufficiency and demonstrate that this upregulation plays an important role in promoting autophagic removal and aggresomal sequestration of toxic misfolded proteins. More recently we documented the mediating role of the calcineurin-TFEB axis in the upregulation of p62 by proteasome malfunction. These discoveries have improved our understanding of cardiac PQC mechanisms, helping devise new therapeutic strategies for heart disease with increased proteotoxic stress, a highly prevalent category of heart diseases including for example ischemic heart disease, some of the familial cardiomyopathies, and even pressure overload heart diseases.

- a. Pan B, Li J, Parajuli N, Tian Z, Wu P, Lewno MT, Bedford L, Mayer RJ, Fang J, Liu J, Cui T, Su H, **Wang X**. The calcineurin-TFEB-p62 pathway mediates the activation of cardiac macroautophagy by proteasomal malfunction. *Circ Res* 2020; 127(4):502–518. PMCID: [PMC7416491](https://pubmed.ncbi.nlm.nih.gov/32416491/)
- b. Zang H, Wu W, Qi L, Tan W, Nagarkatti P, Nagarkatti M, Wang X, Cui T. Autophagy inhibition enables Nrf2 to exaggerate the progression of diabetic cardiomyopathy in mice. *Diabetes* 2020;69(12):2720-2734. PMCID: [PMC7679777](https://pubmed.ncbi.nlm.nih.gov/327679777/).
- c. Zheng Q, Su H, Ranek MJ, **Wang X**. Autophagy and p62 in cardiac proteinopathy. *Circ Res*. 2011; 109(3):296-308. PMCID: [PMC3142307](https://pubmed.ncbi.nlm.nih.gov/3142307/).

Complete List of Published Work in My Bibliography:

<https://www.ncbi.nlm.nih.gov/myncbi/xuejun.wang.1/bibliography/public/>