

## CURRICULUM VITAE

Updated date: 10/29/2019

### A. PERSONAL INFORMATION

American Name:	Nelson Chen, Ph.D.
Official Name:	Zhengjia Chen, Ph.D.
Business Address	Research Associate Professor, Department of Biostatistics and Bioinformatics Rollins School of Public Health & Biostatistics and Bioinformatics Shared Core Resource Winship Cancer Institute Emory University, 1518 Clifton Road, GCR 342 Atlanta, GA 30329
Business Phone	(404)-712-8278 (office)
Fax	(404)-778-5016
E-mail:	<a href="mailto:zchen38@emory.edu">zchen38@emory.edu</a>

### B. EDUCATION

Ph.D. in Biostatistics, University of Southern California, Los Angeles, CA, 2008  
Ph.D Dissertation: PHASE I CLINICAL TRIAL DESIGNS (Range and Trend of Expected Toxicity Level in Standard A+B Designs and an Extended Isotonic Design Treating Toxicity as a Quasi-Continuous Variable).

M.S. in Biometry, University of Southern California, Los Angeles, CA, 2001  
Master Thesis: Ethnic Differences in Mammographic Density.

M.S. in Biochemistry and Molecular Biology, Peking University, Beijing, P.R.China, 1998  
Masters Thesis: Genetic Engineering of Metallothionein in Cyanobacteria.

B.S. in Microbiology, Peking University, Beijing, P.R.China, 1995.  
Bachelor Thesis: Characterization of Metallothionein from Cyanobacteria.

### C. AWARDS OR HONORS.

2017: International Chinese Statistical Association (ICSA) Significant Contribution Award as Treasurer at the 2016 ICSA Applied Statistics Symposium.

2016: Award for Outstanding Service to Clinical and Translational Research Committee (CTRC) at Winship Cancer Institute

2013: Award for Dedication to High Quality Oncology Research and Commitment to Clinical and Translational Research Committee (CTRC) at Winship Cancer Institute

08/1999- 07/2001: University of Southern California  
Honor: Research Scholarships.

08/1998- 07/1999: University of Southern California  
Honor: Fellowship.

09/1995-07/1998: Peking University, Beijing, P.R.China  
Honors: Excellent M.S. thesis.

09/1990-07/1995: Peking University, Beijing, P.R.China  
Honors: Exemption of Entrance Examination to Graduate School

#### **D. PROFESSIONAL EXPERIENCES**

1/1/2015-Present  
Research Associate Professor,  
Department of Biostatistics and Bioinformatics  
Department of Hematology and Medical Oncology  
Department of Radiology and Imaging Sciences  
Emory University

8/1/2009-12/31/2014  
Research Assistant Professor,  
Department of Biostatistics and Bioinformatics  
Department of Hematology and Medical Oncology  
Department of Radiology and Imaging Sciences  
Emory University

08/2010- present:  
Co-Director of Biostatistics and Bioinformatics Core of Lung Cancer SPORE, Winship Cancer Institute, Emory University.

01/2019- present:  
Leading Biostatistician of Clinical and Translational Research Committee (CTRC) at Winship Cancer Institute, Emory University

01/2011- present:  
Leading Biostatistician of Phase I Unit of Winship Cancer Institute, Emory University

06/2018 - present:

Director of Biostatistics and Informatics Core of GI SPORE, Winship Cancer Institute, Emory University

01/2019- present:

Director of Biostatistics and Informatics Core of Head & Neck SPORE, Winship Cancer Institute, Emory University

08/2011-12/2018:

Co-Director of Biostatistics and Informatics Core of Head & Neck SPORE, Winship Cancer Institute, Emory University

08/2010- present:

Co-Director of Biostatistics and Bioinformatics Core of Lung Cancer PO1, Winship Cancer Institute, Emory University

5/2003 – 7/2009:

Statistician at Children's Oncology Group, CureSearch, Arcadia, California

1/2001 – 4/2003

Programmer/Analyst at University of Southern California, Los Angeles, California

9/1994 – 7/1998

Research Assistant at Peking University, Beijing, P.R.China

## **E. COMMITTEE MEMBERSHIPS**

- Executive Committee of International Chinese Statistical Association (ICSA) 2016 Applied Statistics Symposium in Atlanta.
- Curriculum Committee in Department of Biostatistics & Bioinformatics, Emory University. 2010 to present.
- Collaborative Committee in Department of Biostatistics & Bioinformatics, Emory University. 2013 to present.
- Clinical and Translational Research Committee (CTRC) at Winship Cancer Institute, Emory University. 2010 to present.
- Full member of the Discovery & Developmental Therapeutics (DDT) program at Winship Cancer Institute, Emory University. 2009 to present.

## **F. PROFESSIONAL APPOINTMENTS and ACTIVITIES**

### **National Grant Review**

2019 – June: Member, Clinical Oncology (CONC) Study Section, Center for Scientific Review, Bethesda, MD.

February 2018 – present: Statistician of National Institute of Allergy and Infectious Diseases (NIAID) Host Directed Therapy for Tuberculosis and HIV Co-Infection.

January 2018: Grant Reviewer of Nevada IDEA Network of Biomedical Research Excellence (INBRE) program of NIH.

### **Local Grant Review**

2016 Member of review committee of Research Technologies Program at the Atlanta Clinical & Translational Science Institute (ACTSI).

2014 Member of Emory University Research Committee (URC), Atlanta, Georgia.

### **Professional Activities**

Organizer and Chair of an invited session entitled “Novel Statistical Methods for Big Health Data” at ICSA 2019 International Conference in Hangzhou, China.

Organizer and Chair of an invited session entitled “Utilization of Big Data in Precision Medicine” at ICSA 2019 International Conference in Hangzhou, China.

Organizer and Chair of an invited session entitled “Adaptive Methods and Regulation for Clinical Trials” at 2016 ICSA Applied Statistics Symposium in Atlanta.

Chair of an invited session entitled “Semiparametric statistical methods for complex data” at 2016 ICSA Applied Statistics Symposium in Atlanta.

## **G. TEACHING EXPERIENCE**

### **Lecturer at Emory University and Winship Cancer Institute.**

Course Number	Course Title	Credit Hours	Number of students enrolled
2019 Summer Semester	BIOS 500S: Phase I Clinical Trials in Emory SIBS 2019	NA	NA
2019 Spring Semester	Bois 520 Clinical Trial Methodology (Sole Instructor)	2	37
2018 Summer Semester	BIOS 500S: Phase I Clinical Trials in Emory SIBS 2018	NA	NA
2018 Spring Semester	Statistics in oncology overview	NA	About 15
2017 Summer Semester	BIOS 500S: Phase I Clinical Trials in Emory SIBS 2017	NA	NA
2017 Spring Semester	Bois 520 Clinical Trial Methodology (Sole Instructor)	2	36
2016 Fall Semester	Journal Club of Medical Fellows (Statistical lecturer on Phase I clinical trial design)	NA	About 25

2015 Spring Semester	Bois 520 Clinical Trial Methodology (Sole Instructor)	2	30
	Journal Club of Medical Fellows (Statistical lecturer on Phase I clinical trial design)	NA	About 25
2014 Spring Semester	Journal Club of Medical Fellows (Statistical lecturer on Phase I clinical trial design)	NA	About 20
2013 Spring Semester	Bois 520 Clinical Trial Methodology (Sole Instructor)	2	21
	Bois 520M Clinical Trial (A lecture on interim analysis)	2	About 20
	Journal Club of Medical Fellows (Statistical lecturer on Phase I clinical trial design)	NA	About 30
2012 Fall Semester	Fundamentals of Clinical Research (Statistical lecturer on clinical trial)	2	About 6
2011 Spring Semester	Bois 520 Clinical Trial Methodology (Sole Instructor)	2	22
	Journal Club of Medical Fellows (Statistical lecturer)	NA	About 20
2010 Fall Semester	Fundamentals of Clinical Research (Statistical lecturer on clinical trial)	2	About 7
	Journal Club of Medical Fellows (Statistical lecturer)	NA	About 20
2010 Spring Semester	Journal Club of Medical Fellows (Statistical lecturer)	NA	About 20
	Winship Elkin Lecture (A lecture on Phase I clinical trial design)	NA	About 40

Teaching Assistant at Division of Biostatistics, Department of Preventive Medicine, University of Southern California.

- 2009 Spring semester: PM511. Data Analysis.
- 2008 Fall semester: PM522. Introduction to the Theory of Statistics
- 2008 Spring semester: PM510. Principle of Biostatistics

## **G. Students Mentoring**

### \* PhD Dissertation Advisor

1. Yuzi Zhang, PhD, Dept. of Biostatistics and Bioinformatics, Emory University, 2018-Present. “Advanced Bayesian Approaches for Early Phase Clinical Trials”.

2. Ye Cui, PhD, (jointly advising with Dr. Runyan Luo, PhD), Dept. of Mathematics and Statistics, Georgia State University, 2010-2013 (Graduated). “Advanced Designs of Cancer Phase I and Phase II Clinical Trials”.

MPH Thesis Advisor

1. Yi Lasanajak, MS, Dept. of Biostatistics and Bioinformatics, Emory University, 2012-2013. (graduated). “Evaluation of Significant Biomarkers Associated with Progression Free Survival and Overall Survival in Thyroid Cancer Patients”.
2. Zheng Li, MS, Dept. of Biostatistics and Bioinformatics, Emory University, 2012-2014. (graduated). “Dose Escalation with Over-dose and Under-dose Controls for Phase I/II Clinical Trials”. **Won the Shepard Award for the best thesis of the Department of Biostatistics and Bioinformatics 2014.**
3. Run Zhang, MS, Dept. of Biostatistics and Bioinformatics, Emory University, 2014-2016. (graduated). “Escalation with overdose control using normalized equivalent toxicity score and patient’s characteristics to define personalized maximum tolerated dose in phase I clinical trial”. **Won the Shepard Award for the best thesis of the Department of Biostatistics and Bioinformatics 2016.**
4. Zheyu Hu, MS, Dept. of Biostatistics and Bioinformatics, Emory University, 2014-2016. (graduated). “Racial disparities in multiple myeloma”.
5. Yufan Chen, MS, Dept. of Biostatistics and Bioinformatics, Emory University, 2016-2017. (graduated). “Survival Analysis of Patients with Acute Myeloid Leukemia”.
6. Yuzi Zhang, MS, Dept. of Biostatistics and Bioinformatics, Emory University, 2016-2018. (graduated). “Adaptive Bayesian Phase I Clinical Trial Designs for Estimating the Maximum Tolerated Doses for Two Drugs while Fully Utilizing all Toxicity Information”.
7. Chaejin Kim, MS, Dept. of Biostatistics and Bioinformatics, Emory University, 2016-2018. (graduated). “Partitioning Around Medoids Clustering in Follicular Lymphoma Patients: Comparison with FLIPI and FLIPI2 in PFS”.
8. Tianyu Gao, MS, Dept. of Biostatistics and Bioinformatics, Emory University, 2017-2018. (graduated). “ Safety of Administering Pegfilgrastim on the Same Day of Continuously infused 5-Fluorouracil (5-FU) for Patients With Gastrointestinal (GI) Malignancies: A Retrospective Study”.
9. Tianyi Xu, MS, Dept. of Biostatistics and Bioinformatics, Emory University, 2017-2018. (graduated). “Impacts and Improvement in Delineation of the lumpectomy Cavity Boost Volume by using 3-D Implantable Tissue Markers”.
10. Youyun Zheng, BS, Dept. of Computer Science, Emory University, 2016-2018. (**graduated with the highest honor**). “Interactive Software for Dose Recommendation and Simulation of Phase I Cancer Clinical Trial Using EWOC-NETS Design”.
11. Bokai Zhao, MS, Dept. of Biostatistics and Bioinformatics, Emory University, 2018-2019. (graduated). “Investigation of Multiple Biomarkers in Predicting the Disease Free Survival and Overall Survival Among Head and Neck Cancer Patients”.

12. Chenyue Yang, MS, Dept. of Biostatistics and Bioinformatics, Emory University, 2018-2019. (graduated). “Effect Of 5-FU Bolus on Survival in Patients with Metastatic Colorectal Cancer Treated with Combination Chemotherapy”.
13. Jiapeng Shuang, MS, Dept. of Biostatistics and Bioinformatics, Emory University, 2018-2019. (graduated). “Real-world Outcomes of Immune Checkpoint Inhibitors (ICI) in Lung Cancer Patients”.
14. Tao Liang, MS, Dept. of Biostatistics and Bioinformatics, Emory University, 2018-2019. (graduated). “Impact of uncommon genomic alterations on outcomes in metastatic colorectal cancer patients”.
15. Yawen Wang, MS, Dept. of Biostatistics and Bioinformatics, Emory University, 2018-2019. (graduated). “Association between breast cosmesis and two common radiotherapy regimens”.
16. Zeyuan Wang, MS, Dept. of Biostatistics and Bioinformatics, Emory University, 2018-2019. (graduated). “Effect of Lipid-Lowering Therapy on Epicardial Adipose Tissue Radiodensity in Hyperlipidemic Post-Menopausal Women”.

#### MPH Student Thesis Reader

1. Shaoman Yin, MS, Department of Biostatistics, Emory University, 2012-2013. (graduated). “Analyzing Durability and Efficacy of Long-lasting Insecticide-treated Bed Nets: A Longitudinal Monitoring Study at Western Kenya”.

#### Internship Provided

1. Zhibo Wang, MS, Dept. of Mathematics and Statistics, Georgia State University, 2010-2012. “Development of Statistical Software for Phase I Clinical Trials”.
2. Jing Sun, MS, Dept. of Computer Science, Emory University, 2010. “Development of Statistical Software for Phase I Clinical Trials”.
3. Anran Liu, MS, Dept. of Biostatistics and Bioinformatics, Emory University, 2014. “Escalation with overdose control using normalized equivalent toxicity score and patient’s characteristics to define personalized maximum tolerated dose in phase I clinical trial”.

#### Research Assistantship Provided

1. Zheng Li, MS, Dept. of Biostatistics and Bioinformatics, Emory University, 2012-2014. “Advanced Designs of Cancer Phase I Clinical Trials”.

#### Supervisor of Students for Practicum

1. Xiaojing Wang, MS, Dept. of Biostatistics and Bioinformatics, Emory University, 2012-2013.
2. Songli Xu, MS, Dept. of Biostatistics and Bioinformatics, Emory University, 2009-2010.
3. Chang Liu, MS, Dept. of Biostatistics and Bioinformatics, Emory University, 2012-2013.
4. Xiaowei Liu, MS, Dept. of Actual Science, Georgia State University, 2013-2014.
5. Anqi Pan, BS, Dept. of Biostatistics and Bioinformatics, Emory University, 2015-2016. “Prevalence of Zinc Deficiency in Patients with Upper GI Cancer on Chemotherapy”.

6. Ishaan Amit Dave, BS, Dept. of Biostatistics and Bioinformatics, Emory University, 2016. “EWOC-NETS for combination of two drugs in cancer Phase I clinical trial”.
7. Yuzi Zhang, BS, Dept. of Biostatistics and Bioinformatics, Emory University, 2016-Present.
8. Chaejin Kim, BS, Dept. of Biostatistics and Bioinformatics, Emory University, 2016-Present.
9. Wanqi Chen, BS, Dept. of Biostatistics and Bioinformatics, Emory University, 2018-Present.
10. Weixing Huang, BS, Dept. of Biostatistics and Bioinformatics, Emory University, 2018-Present.

## **H. EDITOR AND REVIEWER OF THE JOURNALS**

### Editors:

1. Guest Editor of Journal of Probability and Statistics
2. Editor of New Journal of Science
3. Associate Editor of Clinical Cancer Research Letter
4. Member of Editorial Board of Journal of Bioinformatics, Computational and Systems Biology

### Reviewer:

#### Statistical Journals:

1. Biometrics.
2. Biostatistics.
3. Statistics in Medicine.
4. Statistics in Biopharmaceutical Research.
5. Contemporary Clinical Trial.
6. Pharmaceutical Statistics
7. Journal of Biopharmaceutical Statistics
8. Journal of Bioinformatics, Computational and Systems Biology

#### Other Scientific Journals:

1. Journal of Clinical Oncology.
2. Cancer.
3. Pediatric Blood and Cancer.
4. Ethnicity and Disease.
5. Neuro-Oncology.
6. International Journal of Epidemiology.
7. Plos One.

## **I. MEMBERSHIP OF PROFESSIONAL ASSOCIATIONS**

1. 2002 to present, Member of American Statistical Association (ASA)
2. 2009 to present, Member of International Chinese Statistics Association (ICSA).
3. 2003 to present, Member of Children Oncology Group (COG).
4. 2008 to present, Member of The Society for Clinical Trials (SCT).
5. 2002 to present, Member of American Association of Cancer Research (AACR).



## J. GRANTS

### ACTIVE GRANTS

4 P30 CA138292-08S1 (W. Curran) 04/07/09-03/31/22  
NIH/NCI \$1,400,000  
Emory Winship Cancer Institute Cancer Center Support Grant  
This Cancer Center Support Grant provides NCI support for the Administration, Research Programs, Shared Resources, Clinical trials and developmental funds for the Winship Cancer Institute of Emory University.  
Role: Co-Director of Biostatistics and Biomedical Informatics Core  
Effort: 20%

1 P50 CA217691-01A1 (Ramalingam) 06/01/19-08/31/24  
NIH/NCI \$1,474,358 (Bios Core 3)  
Emory University Lung Cancer SPORE  
This SPORE grant is to improve response to immunotherapy in NSCLC and develop novel targeted agents for resistant NSCLC.  
Role: Co-Director of Biostatistics and Biomedical Informatics Core (Bios Core 3)  
Effort: 20%

HHSN261201800037C (Chen, G.) 07/01/18- 06/30/19  
OCEAN NANOTECH \$427,529  
Quantum Dot FRET Immunoassay to Quantify Intramolecular Epitopes of Analytes in Tissue Sections  
The objective of this project is to develop a quantum dot (QD)-based immunoassay to quantify activated growth signal transducers using proximity technology of Fluorescence Resonance Energy Transfer (FRET). We hypothesize that a ratio of activated signal transducers usually as phosphorylated proteins to their total protein levels in cancer cells can be determined by FRET signal. Quantification of activated growth signal transducer is critical for us to understand cancer cell progression and evaluate anti-cancer therapies which aim to block growth signaling pathway and induce apoptosis.  
Role: Co-Investigator  
Effort: 3%

5 R01 CA140515-09 (Chen, J.) 08/01/15-07/31/20  
NIH/NCI \$252,098  
Metabolic Rewiring by Oncogenic BRAF V600E Links Ketogenesis Pathway to BRAF-MEK1 Signaling  
In this proposal, we plan to explore the molecular mechanisms by which oncogenic BRAF V600E “rewires” metabolic and cell signaling networks and signals through transcription factor Oct-1 to promote the HMGCL-acetoacetate axis that selectively enhances BRAF V600E-dependent tumor development.  
Role: Co-Investigator  
Effort: 5%

1 R01 CA201340-01 (Marcus) 12/16/15-08/30/20  
NIH/NCI \$228,750

Defining Early Escape Strategies in LKB1 Mutant Lung Cancer

The goal of this project is to define how LKB1 loss impacts lung cancer metastasis. Therefore, we hypothesize that LKB1-mutant cells use EMT to initially invade through the basement membrane, which is followed by defective FAK-based adhesion signaling that allows cells to navigate the collagen microenvironment during metastasis.

Role: Co-Investigator

Effort: 5%

1 R01 CA194027-01A1 (Marcus) 12/04/15-11/30/20  
NIH/NCI \$323,257

Developing a Pharmacologic Approach to Treat LKB1 Mutant NSCLC

The objectives of this proposal are i) to determine if pharmacologic FAK inhibition can specifically suppress the metastasis of Lkb1-mutant lung tumors in vivo and ii) define which clinically observed LKB1 mutations cause pFAK 397 activation and create a tumor vulnerability targetable by pharmacologic FAK inhibition.

Role: Co-Investigator

Effort: 4%

PC160820P1 (Liu) 09/01/17-08/31/20  
US DOD \$157,346

Multimodality Imaging Platform for Neurovascular Bundle Sparing Prostate Radiotherapy to Preserve Sexual Function

The ultimate goal is to develop MRI-guided "Neurovascular Bundle Sparing (NVB) prostate radiotherapy, advance imaging technology to assess treatment-related erectile dysfunction and thereby improve sexual outcomes and quality of life in prostate-cancer survivors.

Role: Co-Investigator

Effort: 5%

1R01 CA207768-01A1 (Kang) 7/1/2017 - 6/30/2022  
NIH/NCI \$250,000

Role of MAST1 kinase in cisplatin resistant cancers

Our central hypothesis is that MAST1 provides cisplatin-resistant proliferative signals through its potential substrate and effectors in cancer cells. Thus, MAST1 signaling represents a promising anti-cancer target in combination with cisplatin.

Role: Co-Investigator

Effort: 5%

5 R01 CA203388 -02 (Mao) 04/01/17-03/31/21  
NIH/NCI \$275,949

2DR MR Correlational Spectroscopy Platform for Molecular and Genetic Characterizations of Glioma

This research is to develop a clinically implemented a 2D COSY platform for non-invasive genetically classifying and characterizing gliomas in patients using an onco- metabolite R(-)-

2-hydroxyglutarate, (2-HG) as a biomarker of heterozygous mutations in the isocitrate dehydrogenase (IDH1 and IDH2) genes.

Role: Co-Investigator

Effort: 5%

1R01CA223220-01A1 (Sun) 07/01/18-06/30/23

NIH \$275,133

Modulation of death receptor 4 in EGFR-targeted cancer therapy

This proposal is to understand the mechanism and biological significance of death receptor 4 downregulation during EGFR-targeted cancer therapy.

Role: Co-Investigator

Effort: 5%

1R01CA228414 (Lesinski) 04/01/18-03/31/22

NIH \$333,238

Targeted MEK inhibition to enhance immunotherapy in cholangiocarcinoma

This proposal is to investigate whether MEKi will synergize with PD-L1 blockade to elicit anti-tumor immune responses via altering cytokine and chemokine signatures that promote CD8<sup>+</sup> T cell infiltration and survival and decreased immunosuppressive cells in the TME.

Role: Co-Investigator

Effort: 5%

7 R01CA208253-03 (Lesinski) 09/15/17-07/31/22

NIH/NCI \$228,511

Enhancing immune therapy in pancreatic cancer by targeting IL-6

This proposal will enhance our understanding of how the stroma influences carcinogenesis and immune suppression in PDAC.

Role: Co-Investigator

Effort: 5%

W81XWH-17-1-0186 (Kang) 09/01/17-08/31/19

US/DoD \$200,000

Signaling and targeting of glutamate dehydrogenase 1 in metastatic non-small cell lung carcinoma

The goal of the proposed research is to decipher the mechanism of metabolic enzyme GDH1-dependent regulation of metastasis and to validate GDH1 as a novel therapeutic target in the treatment of metastatic lung carcinoma using GDH1 small molecule inhibitor R162.

Role: Co-Investigator

Effort: 5%

1R01CA215718-01A1 (Yang) 02/01/18-01/31/23

NIH/NCI \$356,850

Multiparametric MRI-guided Prostate HDR Brachytherapy with Focal Tumor Boost.

The proposed personalized mp-MRI-guided HDR brachytherapy based on accurate prostate segmentation and registration technologies has two clinical benefits: (1) to improve long-term survival and (2) to reduce the side effects.

Role: Co-Investigator  
Effort: 5%

1R01HL143350-02 (Garcia) 07/01/18-06/30/22  
NIH/NHLBI \$442,618

Quantification of myocardial blood flow using Dynamic PEC/CAT fused imagery to determine physiological significance of specific coronary lesions  
The aim of this work is to develop software tools to fuse coronary anatomy data obtained from CT coronary angiography with dynamic PET data (combination of anatomic and physiologic information) to noninvasively measure absolute myocardial blood flow, flow reserve and relative flow reserve across specific coronary lesions.

Role: Co-Investigator  
Effort: 5%

1 R01 CA175316-01A1 (Kang) 04/01/19-02/28/24  
NIH/NCI \$207,500

Transcription-dependent and -independent Signaling of RSK2 in Cancer Metastasis  
In this proposal, we will examine the role of RSK2 as a signal integrator in metastatic cells by phosphorylating and regulating multiple protein factors to provide anti-apoptosis, pro-invasive and pro-metastatic signals in cancer cells, and validate RSK2 signaling pathways in combination with others as an alternative therapeutic target in treatment of metastatic cancers using a novel RSK inhibitor FMK-MEA.

Role: Co-Investigator  
Effort: 5%

5 R01 CA203388 -02 (Mao) 04/01/17-03/31/21  
NIH/NCI \$275,949

2DR MR Correlational Spectroscopy Platform for Molecular and Genetic Characterizations of Glioma.

This research is to develop a clinically implemented a 2D COSY platform for non-invasive genetically classifying and characterizing gliomas in patients using an onco-metabolite R(-)-2-hydroxyglutarate, (2-HG) as a biomarker of heterozygous mutations in the isocitrate dehydrogenase (IDH1 and IDH2) genes.

Role: Co-Investigator  
Effort: 5%

PC180599 (Olson) 08/01/2019 – 07/31/2022  
US DOD \$600,000

Targeting Rb loss using BET inhibition to reprogram the prostate tumor microenvironment and enhance the efficacy of immunotherapy.

We seek to determine if Rb loss decreases T cell infiltration into the tumor microenvironment and can be reprogrammed using BET inhibition. We hypothesize that Rb-deficient cancer is associated with an immunosuppressive tumor microenvironment and can be selectively targeted via BET inhibition.

Role: Co-Investigator  
Effort: 5%

R01 CA 226992 (Jani/Schuster) 05/01/2019-04/30/2024.  
NIH NCI \$3.4M  
Advanced PET-CT Directed Post-Prostatectomy Radiotherapy to Enhance Prostate Cancer Outcomes  
The results of this trial will be significant for the use of advanced molecular imaging to guide post-prostatectomy radiotherapy.  
Role: Co-Investigator  
Effort: 5%

%48889 (Harvey III) 01/01/19-12/31/19  
Hematology/Oncology Pharmacy Association \$49,312  
Phase II Evaluation of the Effect of 2 Versus 6 Hour Oxaliplatin Infusions on Neuropathy.  
This research is to support a hematology/oncology pharmacy practice research project with subsequent publication that aligns closely with the HOPA strategic plan to demonstrate the value of hematology/ oncology pharmacy practice.  
Role: Co-Investigator  
Effort: 5%

1 R01 CA228406-01A1 (Lesinski) 04/01/2019-03/30/2024  
NIH \$253,094  
Targeting immune stroma interactions in pancreatic cancer.  
Testing the hypothesis that the tumor microenvironment represents a major barrier promoting immune suppression in pancreatic cancer, and that targeting Hsp90 can modulate components of the tumor microenvironment such as cytokines and signaling in T cells to enhance the efficacy of anti-PD-1 immunotherapy.  
Role: Co-Investigator  
Effort: 5%

### **COMPLETED GRANTS**

1 R01 CA175316-01A1 (Kang) 04/01/14-02/28/19  
NIH/NCI \$207,500  
Transcription-dependent and -independent Signaling of RSK2 in Cancer Metastasis  
In this proposal, we will examine the role of RSK2 as a signal integrator in metastatic cells by phosphorylating and regulating multiple protein factors to provide anti-apoptosis, pro-invasive and pro-metastatic signals in cancer cells, and validate RSK2 signaling pathways in combination with others as an alternative therapeutic target in treatment of metastatic cancers using a novel RSK inhibitor FMK-MEA.  
Role: Co-Investigator  
Effort: 5%

1 R21 MH108928-01 (Goodman) 01/01/16-12/31/18  
NIH/NIMH \$150,000  
PET Imaging Agents for 5-HT<sub>2C</sub> Receptors  
This project aims to satisfy this unmet need to develop carbon-11 5-HT<sub>2C</sub>-specific radioligand antagonists with characteristics appropriate for in vivo imaging studies by PET.

Role: Co-Investigator  
Effort: 5%

2017 Seed Grant (Shaib) 11/15/17-11/14/18  
Hirshberg Foundation For Pancreatic Cancer \$40,000  
Effect of Treatment of Immune Pathways in Pancreatic Adenocarcinoma: A Comparative  
This is a tissue study of Pancreas cancer resected from patients. 50 patients received  
treatment prior to surgery and 50 patients did not receive treatments. We are comparing  
immune marker differences in these 2 groups.  
Role: Co-Investigator  
Effort: 2%

5 R01 CA 169188-04 (Jani) 09/01/12-06/30/18  
NIH NCI \$328,840  
Advance Molecular Imaging with FACBC PET-CT to Improve Post prostatectomy  
Radiotherapy  
The results of this trial will be significant for the use of advanced molecular imaging to guide  
post-prostatectomy radiotherapy.  
Role: Co-Investigator  
Effort: 5%

2 R01 GM069971-09 (Morgan) 02/01/04-05/31/18  
NIH/NIGMS \$316,710  
Nitric Oxide Regulation of Human CYP Enzymes  
We propose to use proteomic methods to define the scope of NO-mediated degradation in  
human hepatocytes. We will characterize the proteolytic enzymes involved in the NO-  
mediated degradation of CYP2B6 and 2C9, and elucidate the mechanism whereby NO  
regulates these processes in an enzyme-specific manner. This project will help us to  
understand how nitric oxide formed in the liver during inflammation contributes to this  
change, and so will allow us to predict what patients will need to have their drug doses  
adjusted to avoid this problem.  
Role: Co-Investigator  
Effort: 3%

4 R01 CA169937-04 (Mao) 04/15/13-03/31/18  
NIH \$289,970  
MR Investigation of IDH Mutation and Its Marker 2-HG in Brain Tumor Patients  
We propose to develop a novel, non-invasive and clinically applicable(MRS) method for  
investigating a newly discovered onco-metabolite, R(-)-2-hydroxyglutarate (2-HG), that  
accompanies the progression of gliomas.  
Role: Co-Investigator  
Effort: 6%

640011-0615-04 (Qayed) 07/01/14-06/30/17  
Children's Health Care of Atlanta (CHOA) \$92,104

A Phase I Study of Mesenchymal Stromal Cells for the Treatment of Acute and Chronic Graft versus Host Disease

The long-term goal of this project is to develop autologous mesenchymal stromal cells (MSC) as a cellular therapeutic to treat graft versus host disease (GVHD).

Role: Co-Investigator

Effort: 1%

(Sechopoulos)

08/16/13-08/30/17

Susan G. Komen Foundation

\$186,440

Achieving Personalized Radiation Dose Estimates in Breast Cancer Screening

The objective of this research project is to develop, optimize, validate and test the methods necessary to obtain an accurate estimate of the personalized radiation dose each patient receives during breast cancer screening.

Role: Co-Investigator

Effort: 3%

1 R01 CA157754-01A1 (Nahta)

06/07/12-04/30/17

NIH/NCI

\$207,500

Mechanisms of Herceptin Resistance

The long-term goal of this application is to identify new drug targets and predictors of Herceptin resistance in order to improve the survival of patients with HER2-overexpressing breast cancer.

Role: Co-Investigator

Effort: 3%

1 R01 CA160522-01A1 (Sun)

04/01/12-03/31/17

NIH

\$207,500

Therapeutic potential of mTOR kinase inhibitors in lung cancer

This proposal will evaluate the therapeutic potential of novel mTOR kinase inhibitors either alone or in combination with other agents against lung cancer and determine the impact of genetic alterations on cell responses to these inhibitors.

Role: Co-Investigator

Effort: 5%

5 U01 CA168930-03 (Cummings)

08/23/12-06/30/17

NIH/NCI

\$269,449

The Tumor Antigens Tn and SialylTn in Human Colorectal Carcinoma

The goal of this study is to propose, test and validate a new assay for colorectal carcinoma based on the Tn and SialylTn Antigens. Although there exist assays for early detection of other carcinomas, there is none for colorectal carcinoma and this would be the first of its kind.

Role: Co-Investigator

Effort: 7%

(Cox)

07/01/15-06/30/16

Radiological Society of North America

\$40,000

MRI Liver Surface Nodularity Score: A New Noninvasive Biomarker for Chronic Viral

Role: Co-Investigator  
Effort: 5%

1 R41 CA186498-01A1 (Kucuk) 09/01/15-08/31/16  
NIH/NCI \$31,027  
A Dietary Supplement as Adjunct Therapy in Castration-Resistant Prostate Cancer.  
Role: Co-Investigator  
Effort: 20%

1 R43 CA183312-01 (Chen) 05/01/14-04/30/15  
Ocean Nanotech/NIH/NCI \$32,051  
Assessment of Microvessel Density and Tumor Proximity for Prognosis of Cancer  
The objective of this project is to develop a quantum dot (QD)-based imaging technology for assessment of microvessel density and tumor proximity in order to provide an accurate prognosis for cancer patients.  
Role: Co-Investigator  
Effort: 2%

(Goldsmith) 09/01/14-08/31/15  
Children's Health Care of Atlanta (CHOA)  
PHORG: A Phase I study using simvastatin in combination with conventional chemo  
Phase I trial of simvastatin in combination with topotecan/cyclophosphamide or vincristine/doxorubicin/dexrazoxane for refractory and/or relapsed solid tumors of childhood to define toxicity and to evaluate cholesterol levels and IL6/STAT3 pathway change.  
Role: Co-Investigator  
Effort: 3.5%

5 R33 CA161873-02 (Chen & Hoyt) 07/27/11-06/30/15  
NIH/NCI \$287,275  
Developing a Platform for Prediction of Metastasis Using Multiplexed QD-imaging  
This project proposes to develop a platform which combines imaging and quantification of multiplexed immunostaining plus bioinformatics for the prediction of lymph node metastases from the primary tumor of squamous cell carcinoma of the head and neck.  
Role: Co-Investigator  
Effort: 2%

5 P50 CA128301-04 (Meltzer) 09/28/07-08/31/14  
NIH/NCI  
Emory Molecular and Translational Imaging Center  
The long term goal of this research is to determine if PET with anti-[18F] FACBC will lead to improved patient care in the diagnosis and staging of prostate carcinoma and to elucidate the mechanism of its uptake within malignant cells. A secondary goal is to translate this work to facilitate the use of intensity-modulated-radiation-therapy (IMRT) for treatment of prostate carcinoma.  
Role: Co-Investigator  
Effort: 11%

(Chen) 09/28/12-09/27/14



Ocean Nanotech

MultiBiomarker Imaging Using Quantum Dots for Prediction of HPV-Associated Cancer

The objective of this project is to utilize QD technology for the prediction of HPV associated human cancer. Our team has extensive expertise in studies of HPV and cancer biomarkers.

We also have experience in using QD for imaging and quantification of biomarker expression in patient's tumor tissue specimens.

Role: Co-Investigator

Effort: 5%

CSOM230CUS18T (Owonikoko)

09/01/10-09/30/14

Novartis Pharmaceuticals Corporation

CSOM230CUS12T: A 3-Arm Randomized Phase II Trial Evaluating Single Agent and Combined Efficacy of Pasireotide and Everolimus in Adult Patients with Radiiodine-Refractory Differentiated and Medullary Thyroid Cancer

The overall goal of this project is to evaluate single agent and combined efficacy of Pasireotide and Everolimus in adult patients with radiiodine-refractory differentiated and medullary thyroid cancer.

Role: Co-Investigator

Effort: 5%

MORAB-009-003 (Owonikoko)

05/24/11-05/24/15

Morphotek

A Study of the Pattern of Expression of Alpha-Folate Receptors, Mesothelin, and Endosialin in Pleural Mesotheliomas

The overall goal of this project is to study the Pattern of Expression of Alpha-Folate Receptors, Mesothelin, and Endosialin in Pleural Mesotheliomas.

Role: Co-Investigator

Effort: 1%

CCOP SEED GRANT (Higgins)

03/01/13-02/28/14

Radiation Therapy Oncology Group

A Pilot Study of Soy Isoflavone in Combination with Radiation Therapy and Cisplatin in Locally Advanced Squamous Cell Carcinoma of the Head and Neck

We hypothesize that soy isoflavone supplementation administered to patients with SCCHN undergoing concurrent modality therapy may decrease radiation-induced toxicity. We are asking for support for a pilot project examining the feasibility of supplementation of soy Isoflavones to patients with SCCHN undergoing concurrent chemoradiation.

Role: Co-Investigator

Effort: 1%

5 K23 EB013221-02(Tridandapani)

06/01/11-05/31/14

NIH

An Innovative Ultrasound-Based Prospective-Gating Technique for Cardiac CT

The objective of the proposed research is to merge the unique capabilities of ultrasound (US) and computed tomography (CT) by using real-time US information to provide a gating signal in order to obtain motion-free CT images of coronary arteries.

Role: Co-Investigator  
Effort: 5%

5 P01 CA116676-03(Khuri)

06/20/06-05/31/11

NIH

Lung P01

Targeting Cell Signaling in Lung Cancer to Enhance Therapeutic Efficacy The overall goal is to develop effective clinical approaches to prevent, treat, and cure lung cancer.

Role: Co-Director of Biostatistics and Biomedical Informatics Core

Effort: 25%

5 R01 CA112643-05(Shin)

08/18/06-07/31/13

NIH

Second Primary Tumor Prevention with EGFR TKI'S and COX-2 Inhibitors in Head and Neck Cancer

The overall goal of this project is to study second primary tumor prevention using EGFR TKIs and COX-2 Inhibitors in head and neck cancer.

Role: Co-Investigator

Effort: 5%

5 P50 CA128613-05(Shin)

08/23/07-06/30/12

NIH

SPORE in Head and Neck

The goal of this proposal is to improve prevention and treatment of head and neck cancer with emphasis on new discoveries and rapid translation to patients who are suffering disability and morbidity caused by this disease.

Role: Co-Director of Biostatistics and Informatics Core

Effort: 25%

5 R01 CA116804-05(Van Meir)

09/01/07-07/31/12

NIH/NCI

Targeting Glioblastoma Using Novel Small Molecule HIF-1 Pathway Inhibitors

The overall goal of this project is to see if small molecules that inhibit the HIF-1 pathway will inhibit the growth of glioblastoma, either singly or in combination with other agents.

Role: Co-Investigator

Effort: 4%

R01 CA131294-01A2(Sharma)

07/01/09 - 12/31/10

Title of project: Role of Adipocytokines, Leptin and Adiponectin in Breast Carcinogenesis

Funding agency: NIH

Role: Co-Investigator.

Effort: 3%

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### Meeting Abstracts

1. **Chen Z**. Adaptive Approach for Estimating MTDs of Two Drugs by Fully Utilizing All Toxicity Data. The Southern Regional Council on Statistics (SRCOS) 55nd Summer Research Conference. 2019, Session V: New Advances in Clinical Trial.
2. Sahin IH, Chen W, Sonbol MB, Das S, **Chen Z**, Akce M, Alese OB, Shaib WL, Ahn DH, Ciombor KK, Borad MJ, Berlin J, Bekaii-Saab TS, Draper A, El-Rayes BF, Wu C. Analysis of age, tumor-sidedness, and mismatch repair (MMR) genes with response to immune checkpoint inhibitors (ICIs) in MMR-deficient (dMMR) colorectal cancer (CRC) patients (pts): A multi-institutional study. 2019 ASCO Annual Meeting. Abstract No: e15029.
3. Sahin IH, Chen W, **Chen Z**, Akce M, Alese OB, Shaib WL, El-Rayes BF, Wu C. Impact of genomic alterations (GAs) on outcomes and their distribution by age groups in metastatic colorectal cancer (mCRC) patients (pts). 2019 ASCO Gastrointestinal Cancers Symposium. Abstract ID#: 240471.
4. Mao J, Strouse CS, Goldman M, Chen W, **Chen Z**, Maurer MJ, Calzada O, Churnetski M, Flowers C, Cerhan JR, Link BK, Thompson CA, Cohen JB. Impact on survival of surveillance imaging after first remission in follicular lymphoma. 2019 ASCO Annual Meeting. Abstract No: 7536.
5. Patel U, Meisel JL, Barbee MS, Frinzi K, Sakach E, **Chen Z**. The clinical utility of strict laboratory monitoring of CDK 4/6 inhibitors in metastatic breast cancer patients. 2019 ASCO Annual Meeting. Abstract No: 1062.
6. Owonikoko TK, Higgins KA, **Chen Z**, Zhang C, Pillai RN, Steuer CE, Saba NF, Pakkala S, Shin DM, Zhang G, Wang S, Hossain MS, Beardslee T, Engelhart A, Revenig J, Khuri F, Curran WJ, Lonial S, Waller EK, Ramalingam SS. A randomized phase II study of tremelimumab and durvalumab with or without radiation for patients with relapsed small cell lung cancer (SCLC). 2019 ASCO Annual Meeting. Abstract No: 8515.
7. Nazha B, **Chen Z**, Goyal S, Engelhart A, Carlisle JW, Beardslee T, Gill H, Odikadze L, Liu Y, Mishra MK, Behera M, Ramalingam SS, Owonikoko TK. Evaluating the role of race in



outcome of advanced non-small cell lung cancer (NSCLC) patients treated with immune checkpoint inhibitor (ICI): Our institutional experience. 2019 ASCO Annual Meeting. Abstract No: 9042.

8. Dhere V, Sudmeier L, Buchwald Z, Tian S, Jiang X, Zhang C, **Chen Z**, Eaton B, Shu HG, Crocker I, Curran W, Zhong J. Hypofractionated Radiation for Benign Meningiomas and Vestibular Schwannomas. 2019 ASTRO's 61st Annual Meeting. Poster #: 2218.
9. Sudmeier LJ, AbugideirivM, Tian S, Goyal S, **Chen Z**, Eaton BR, Khan MK, Esiashvili N. Reduced Lung Dose is Associated with Decreased Incidence of Pulmonary Toxicity after Total Body Irradiation in Pediatric Patients. 2019 ASTRO's 61st Annual Meeting.
10. Abiodun-Ojo OA, Jani AB, Akintayo AA, Alemozaffar M, Akin-Akintayo OO, Odewole O, Tade FI, Nieh PT, Master VA, Patel P, Shelton J, Kucuk O, **Chen Z**, Hershatter B, Fielder B, Halkar RK, Schuster DM. [<sup>18</sup>F]-fluciclovine positivity rate is not affected by androgen deprivation therapy (ADT) in recurrent prostate cancer post-prostatectomy. Society of Nuclear Medicine and Molecular Imaging (SNMMI) 2019 Annual Meeting.
11. Abiodun-Ojo OA, Jani AB, Akintayo AA, Alemozaffar M, Akin-Akintayo OO, Tade FI, Master VA, Patel P, Shelton J, Kucuk O, **Chen Z**, Hershatter B, Schreibmann E, Fielder B, Halkar RK, Schuster DM. Findings on 18F-fluciclovine PET/CT with failure-free survival of salvage radiotherapy in post-prostatectomy patients with biochemical recurrence. RSNA 2019 annual meeting. Abstract No: 19014549.
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23. Wade J, Little J, Zhang C, **Chen Z**, Meisel J, Hanley KZ. Pathologic characteristics of node positive invasive breast carcinomas associated with extranodal extension. USCAP 2018 Annual Meeting. Abstract No: 137.
24. Abiodun-Ojo OA, Jani AB, Akintayo AA, Alemozaffar M, Akin-Akintayo OO, Odewole O, Tade FI, Nieh PT, Master VA, Patel P, Shelton J, Kucuk O, **Chen Z**, Hershatter B, Fielder B, Halkar RK, Goodman MM, Schuster DM. Change in salvage radiotherapy management based on fluciclovine (<sup>18</sup>F) PET/CT guidance in post-prostatectomy recurrent prostate cancer. RSNA 2018 annual meeting. Abstract No: 18010759.
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27. Abugideiri M, Press RH, Zhang C, Thomas M, Tian S, Jhaveri J, R. J. Cassidy III RJ, Zaenger D, Morgan T, Madden N, Parks J, Buchwald ZS, Morrison D, **Chen Z**, Robertson Y, Phillips R, Landry JC, and Godette KD. ASTRO 2018 Annual Meeting. Abstract No: .
28. Shaib WL, Sayegh L, Alese OB, Maithel S, Cardona K, Sarmiento J, Belalcazar A, Ip A, Qu Y, Akce M, Zhang C, Wu C, **Chen Z**, El-Rayes BF. Resection of pancreatic cancer following induction chemotherapy. 2018 Gastrointestinal Cancers Symposium. Abstract No: 203305.
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149. Nicholson SH, Zhou T, **Chen Z**, Moran K, Sklar C, Zeltzer L, Neglia J. Quality of Life in Survivors of Childhood and Adolescent Acute Myelogenous Leukemia (AML) Does Not Differ by Treatment (Bone Marrow Transplant (BMT) vs. Chemotherapy). A Report from the Children's Oncology Group (COG). American Society of Hematology (ASH) 2005 Annual Meeting. Blood 2005 106:701.
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153. Ozkaynak F, Krailo M, **Chen N**, Feusner J. Randomized comparison of antibiotics alone and with granulocyte colony-stimulating factor (G-CSF) in children with chemotherapy-induced febrile neutropenia: a report from the Children's Oncology Group. *Pediatric Blood & Cancer*; 42(6):1769a, 2004. Abstract No: 1769.
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### **Book Chapters**



Pappo A, Krailo M, **Chen Z**, Reaman G. The Rare Tumor Initiative of the Children's Oncology Group: Lessons Learned in the First 6 Years and Their Effect on Future Plans. *ASCO Educational Book*; 2009.

#### L. INVITED TALKS

1. SRCOS 2019 Annual Meeting, Kentucky.  
Date: 6/5/2019.  
Topic "Advanced Phase I Clinical Trial Designs".
2. Department of Biostatistics. Washington University. St. Louis, Missouri.  
Date: 6/2/2017.  
Teach a workshop entitled "Commons and Innovative Designs for Phase I Clinical Trial".
3. Department of Biostatistics and Bioinformatics. Georgetown University. Washing DC.  
Date: 2/27/2015.  
Topic: Bayesian optimal designs for cancer Phase I clinical trials.
4. Department of Biostatistics. Washington University. St. Louis, Missouri.  
Date: 3/16/2012.  
Topic: Dose Escalation with Overdose Control using a Quasi-Continuous Toxicity Score in Cancer Phase I Clinical Trials.
5. Workshop of Department of Mathematics and Statistics. Georgia State University. Atlanta, Georgia.  
Date: 5/6/2012.  
Topic: Dose Escalation with Overdose Control using a Quasi-Continuous Toxicity Score in Cancer Phase I Clinical Trials.
6. Department of Mathematics and Statistics. Georgia State University. Atlanta, Georgia.  
Date: 10/19/2010.  
Topic: A Novel Toxicity Scoring System Treating Toxicity Response as a Quasi-Continuous Variable in Phase I Clinical Trials.
7. Winship Cancer Institute. Emory University. Atlanta, Georgia.  
Date: 5/28/2010.  
Topic: A Novel Toxicity Scoring System Treating Toxicity Response as a Quasi-Continuous Variable in Phase I Clinical Trials.
8. Department of Biostatistics and Bioinformatics. Emory University. Atlanta, Georgia.  
Date: 6/14/2009.  
Topic: An Extended Isotonic Phase I Design Treating Toxicity Response as a Quasi-Continuous Variable.
9. Department of Biostatistics. SUNY at Buffalo. Buffalo, New York.  
Date: 6/1/2009.

Topic: An Extended Isotonic Phase I Design Treating Toxicity Response as a Quasi-Continuous Variable.

## M. CLINICAL SOFTWARE DEVELOPED

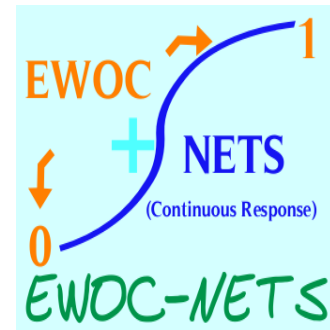
### 1. ID-NETS (Isotonic Design using Normalized Equivalence Toxicity Score).

ID-NETS is a novel semi-parametric Phase I design which is developed by integration of original isotonic design and my NETS system. ID-NETS can substantially improve the accuracy of MTD determination and trial efficiency by fully utilizing all toxicities of each patient and treating toxicity response as a quasi-continuous variable instead of a binary indicator of DLT. ID-NETS is also very robust because it only assumes a monotonically increasing dose toxicity relationship instead of any parametric model. ID-NETS is particularly appropriate for clinical trials which have no reliable parametric model for dose toxicity relationship and the target toxicity level is outside the range of 18-33% of DLT. ID-NETS<sup>®</sup>™ is an interactive and user-friendly statistical software to implement the design ID-NETS in Phase I clinical trials.



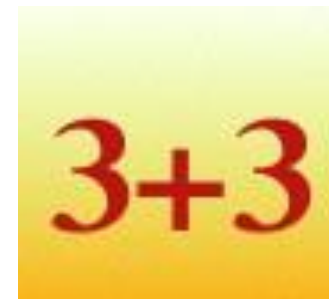
### 2. EWOC-NETS (Escalation with Overdose Control using Normalized Equivalence Toxicity Score).

EWOC-NETS is a novel Bayesian adaptive design which is a combination of EWOC and my NETS system. EWOC-NETS not only retains all the advantages of EWOC, but also treats toxicity response as a quasi-continuous variable instead of a binary indicator of DLT, fully utilizes all toxicity information, and improves the accuracy of MTD estimation and the efficiency of Phase I trials. EWOC-NETS<sup>®</sup>™ is an interactive and user-friendly statistical software to implement the design EWOC-NETS in Phase I clinical trials.



### 3. Standard 3+3 Design simulator

Standard 3+3 with or without dose de-escalation designs are still most widely used in cancer Phase I clinical trials for their simplicity and robustness. In clinical practice and grant applications, it is necessary to provide the operating characteristics of Phase I clinical trials at the planning stage. Therefore I have developed an interactive and user-friendly software called Standard-3+3-design<sup>®</sup>™ to estimate the operating characteristics of Phase I clinical trials using Standard 3+3 designs.



4. Development of R code for application of Escalation with over dose and under dose control (EWOC) design in cancer Phase I/II clinical trial. (R code available per request).
5. Development of R code for application of Escalation with Over Dose with Normalized Equivalent Toxicity Score (EWOC-NETS) design in cancer Phase I/II clinical trial. (R code available per request).
6. Extension of the R code to estimate personalized Maximum Tolerated Dose (MTD) in cancer Phase I/II clinical trial using the design: Escalation with Over Dose with Normalized Equivalent Toxicity Score (EWOC-NETS). (R code available per request)
7. Development of R code for calculation of the operating characteristics of Phase I clinical trials using Standard 3+3 designs. (R code available per request).
8. Development of an interactive statistical software for calculation of the operating characteristics of Phase I clinical trials using standard 3+3 design. (Using R shiny).
9. Development of an interactive statistical software for Phase I clinical trials using EWOC-NETS design. (Using R shiny).

## **N. COMPUTATION SKILLS**

### **Statistical:**

- Statistical Language: SAS (More than 12 years of experience), STATA (More than 4 years of experience), EPILOG More than (8 years of experience), SURVEY More than (3 years of experience), S-plus, SPSS, R language, etc.
- Databases: Access, Oracle, FoxPro, SQL, PL/SQL, MySQL.
- Application Software: Microsoft Word, Excel, Powerpoint, Sigma plot.

### **Computer Science:**

- Programming languages: C/C++, Fortran.
- Platforms: Unix, Windows 95/98/2000/NT, MacOS, Linux, MS-DOS.