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### Education & Training

- 2005 Ph.D. Anthropology (concentration in human statistical genetics of aging and complex disease), Department of Anthropology, Binghamton University, State University of New York.  
Dissertation: Genotype  $\times$  Age Interaction, and the Insulin-like Growth Factor I Axis in the San Antonio Family Heart Study: A Study in Human Senescence
- 2001 M.A. Anthropology (concentration in human population biology and infectious disease), Department of Anthropology, Binghamton University, State University of New York  
Thesis: Models from Mathematical Epidemiology, Ecology and Evolution Meet the Human Treponematoses: Model Development and a Preliminary Analysis of Selected Studies
- 1995 B.A. Biology, University of Guam, Mangilao, Guam

### Research Focus

My research interests are in the statistical genetics of physiological aging with an emphasis on the metabolic syndrome diseases associated with aging. I have focused on developing genotype  $\times$  environment interaction (GEI) approaches to study the role of major environmental variables in the etiology of the cardiometabolic disease. I am also actively engaged in bridging the traditional multivariate approaches of statistical genetics with advances in network and systems biology. To this end, I have successfully developed statistical genetic models to incorporate the effects of large gene expression systems, as measured by transcriptomic data, and their interaction with environmental exposures in the phenotypic determination of complex disease traits. I am currently pursuing the extension of this modeling approach to incorporate

methyloomic measures of the same systems for the creation of multi-omic GEI models for the study of complex risk in relation to environmental exposures

Recently, I have become keenly interested in FVIII immunogenicity and inhibitor development, and in the development of appropriate statistical analytical approaches to studying this problem. I developed generalized linear mixed model (GLMM) approaches to analyze dendritic cell protein processing and presentation assay data from the FVIII Epitope Determination (FED) Study. For the Personalized Alternative Therapies for Hemophilia (PATH) Study I helped to develop another type of GLMM appropriate for the analysis of determinants of inhibitor development in related individuals. Both GLMMs, for the FED Study and the PATH Study, accounted for important sources of non-independence and heterogeneity, which clarified and increased our power for risk factor estimation and prediction.

## Publications

Arya R, Lopez-Alvarenga JC, Almeida M, Kumar S, Peralta J, Diaz-Badillo A, **Diego VP**, Resendez RG, Fowler SP, Jenkinson CP, Lehman D, Curran J, Lynch JL, Hale DE, DeFronzo RA, Mummidi S, Blangero J, Duggirala R. Exome-chip-wide Association Study of Biomarkers of Liver Function and Metabolic Dysfunction-Associated Fatty Liver Disease (MAFLD) in Mexican Americans. *Front Med (Lausanne)*. 2022. In Press.

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Jankowski W, Park Y, McGill J, Maraskovsky E, Hofmann M, **Diego VP**, Luu BW, Howard TE, Kellerman R, Key NS, Sauna ZE. Peptides identified on monocyte-derived dendritic cells: a marker for clinical immunogenicity to FVIII products. *Blood Adv*. 2019 May 14;3(9):1429-1440. doi: 10.1182/bloodadvances.2018030452. PubMed PMID: 31053570; PubMed Central PMCID: PMC6517663.

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**Diego VP**, Rainwater DL, Wang X-L, Cole SA, Curran JE, Johnson MP, Dyer TD, Williams JT, Comuzzie AG, MacCluer JW, Mahaney MC, Blangero. Genotype x Adiposity Interaction Linkage Analyses Reveal a Locus on Chromosome 1 for Lipoprotein-associated Phospholipase A<sub>2</sub>, a Marker of Inflammation and Oxidative Stress. *American Journal of Human Genetics* 80(1):168-77, 2007.

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Voruganti VS, Göring HH, **Diego VP**, Cai G, Mehta NR, Haack K, Cole SA, Butte NF, Comuzzie AG. Genome-wide scan for serum ghrelin detects linkage on chromosome 1p36 in Hispanic children: Results from the Viva La Familia Study. *Pediatric Research* 62(4):445-50, 2007.

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**Diego VP**, Göring HHH, Cole SA, Almasy L, Dyer TD, Blangero J, Duggirala R, Laston S, Wenger C, Cantu T, Dyke B, North K, Schurr T, Best LG, Devereux RB, Fabsitz RR, Howard BV, MacCluer JW. Fasting insulin and obesity-related phenotypes are linked to chromosome 2p: The Strong Heart Family Study. *Diabetes* 55(6):1874-1878, 2006.

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## Abstracts

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