

## Association of Variability and Hypertensive Peaks in 24-h Blood Pressure with Cardiovascular Risk and Mortality

Jesus D. Melgarejo

*The University of Texas Rio Grande Valley, [Jesus.Melgarejo@utrgv.edu](mailto:Jesus.Melgarejo@utrgv.edu)*

Luis J. Mena

*Polytechnic University of Sinaloa*

Kristina P. Vatcheva

*The University of Texas Rio Grande Valley, [kristina.vatcheva@utrgv.edu](mailto:kristina.vatcheva@utrgv.edu)*

Jose A. Garcia

*The University of Texas Rio Grande Valley*

Dhrumil Patil

*Katholieke Universiteit Leuven*

*See next page for additional authors*

Follow this and additional works at: <https://scholarworks.utrgv.edu/somrs>



Part of the [Medicine and Health Sciences Commons](#)

---

### Recommended Citation

Melgarejo, Jesus D.; Mena, Luis J.; Vatcheva, Kristina P.; Garcia, Jose A.; Patil, Dhrumil; Satizabal, Claudia; Chavez, Carlos A.; Pirela, Rosa V.; Calmon, Gustavo; Lee, Joseph H.; Terwilliger, Joseph D.; Seshadri, Sudha; and Maestre, Gladys E., "Association of Variability and Hypertensive Peaks in 24-h Blood Pressure with Cardiovascular Risk and Mortality" (2024). *Research Symposium*. 7.  
<https://scholarworks.utrgv.edu/somrs/2023/talks/7>

This Oral Presentation is brought to you for free and open access by ScholarWorks @ UTRGV. It has been accepted for inclusion in Research Symposium by an authorized administrator of ScholarWorks @ UTRGV. For more information, please contact [justin.white@utrgv.edu](mailto:justin.white@utrgv.edu), [william.flores01@utrgv.edu](mailto:william.flores01@utrgv.edu).

---

**Presenter Information (List ALL Authors)**

Jesus D. Melgarejo, Luis J. Mena, Kristina P. Vatcheva, Jose A. Garcia, Dhrumil Patil, Claudia Satizabal, Carlos A. Chavez, Rosa V. Pirela, Gustavo Calmon, Joseph H. Lee, Joseph D. Terwilliger, Sudha Seshadri, and Gladys E. Maestre

## Association of Variability and Hypertensive Peaks in 24-h Blood Pressure with Cardiovascular Risk and Mortality

Jesus D. Melgarejo, MD, PhD<sup>1,2,3</sup>; Luis J. Mena, PhD<sup>4</sup>; Kristina P. Vatcheva, PhD<sup>1,5</sup>;  
Jose A. Garcia, PhD<sup>6</sup>; Dhruvil Patil, MD<sup>7</sup>; Claudia L. Satizabal, PhD<sup>8,9</sup>; Carlos A. Chavez, MD<sup>3</sup>;  
Rosa V. Pirela, IE<sup>1,2</sup>; Egle Silva, MD<sup>10</sup>; Gustavo Calmon, MD<sup>10</sup>; Joseph H. Lee, PhD<sup>11,12,13</sup>;  
Joseph D. Terwilliger, PhD<sup>11,12,13,14</sup>; Sudha Seshadri, MD<sup>8,9</sup>; Gladys E. Maestre, MD, PhD<sup>1,2,3,6</sup>.

<sup>1</sup>Institute of Neuroscience, University of Texas Rio Grande Valley, Harlingen, TX; <sup>2</sup>Alzheimer's Disease Resource Center for Minority Aging Research, Harlingen, TX; <sup>3</sup>Laboratory of Neuroscience, University of Zulia, Maracaibo, Zulia, Venezuela; <sup>4</sup>School of Mathematical and Statistical Science, UTRGV, Brownsville, TX; <sup>5</sup>Polytechnic University of Sinaloa, Mazatlán, Sinaloa, Mexico; <sup>6</sup>Department of Human Genetics, University of Texas Rio Grande Valley, Brownsville, TX; <sup>7</sup>Cardiovascular and Hypertension Unit, Cardiovascular Department, KU Leuven, Leuven, Belgium; <sup>8</sup>Glenn Biggs Institute for Alzheimer's and Neurodegenerative Diseases, UT Health San Antonio, San Antonio, TX; <sup>9</sup>Department of Neurology, Boston University School of Medicine, Boston, MA; <sup>10</sup>Laboratory of Ambulatory Recordings, Cardiovascular Institute (IECLUZ), University of Zulia, Maracaibo, Venezuela; <sup>11</sup>Departments of Psychiatry and Genetics & Development, Columbia University, New York, NY; <sup>12</sup>Sergievsky Center & Department of Epidemiology, Columbia University Medical Center, New York, NY; <sup>13</sup>Taub Institute for Research on Alzheimer's Disease and the Aging Brain, Columbia University, New York, NY; <sup>14</sup>Division of Public Health Genomics, National Institute for Health and Welfare, Helsinki, Finland.

### Abstract

**BACKGROUND:** Blood pressure (BP) variability relates to cardiovascular (CV) diseases and one unexplored mechanism may involve hypertensive peaks caused by high BP variability.

**OBJECTIVES:** To test this hypothesis, we studied the association of cumulative hypertensive peaks (CHP) in 24-h systolic BP with CV risk.

**METHODS:** A total of 1212 participants from the Maracaibo Aging Study (mean age, 66; women, 67.2%) underwent 24-h ambulatory BP monitoring and were followed between 1998 and 2010. BP variability was the 24-h average real variability (ARV). CHP in systolic BP (expressed as %) was the number of systolic BP measures  $\geq 125$  mmHg (based on the ACC/AHA threshold) each participant experienced over 24-h divided by the number of recordings. The primary endpoint was a composite of fatal and nonfatal coronary, heart failure, and stroke events, while secondary endpoints were total and CV mortality, and fatal and nonfatal coronary and stroke endpoints. Statistics included adjusted Cox proportional models adjusted.

**RESULTS:** During a median follow-up of 8 years, 242 participants developed a composite of any CV endpoint, and 353 died (210 cardiovascular deaths), 129 had coronary and 57 stroke endpoints. An increment of +2 mmHg in 24-h ARV (HR [hazard ratio], 1.18; 95% confidence interval [CI], 1.05-1.33) or +5% in CHP (HR, 1.05; 95% CI, 1.02-1.07) increased CV risk. The inclusion of both indexes in the same Cox proportional models resulted in CHP, but not ARV ( $P=0.075$ ), associated with the primary endpoints ( $P=0.004$ ). For secondary endpoints, the association of ARV attenuated while CHP was similar.

**CONCLUSIONS:** In this population-based cohort study, CHP in 24-h systolic BP explains the association of high 24-h BP variability and CV risk. Clinical management of high 24-h BP variability is challenging but recognizing that an increased variability results in CHP seems a feasible alternative to address in CV prevention.