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## Selenium May Help Explain Racial Differences in HT

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NEW ORLEANS — Reduced serum selenium is an independent predictor of hypertension, according to an analysis of data from the third National Health and Nutrition Examination Surveys.

The findings from this and other studies, that serum selenium concentrations are reduced in African Americans, compared with whites, may in part explain the increased incidence of hypertension in African Americans, Dr. Chizobam Ani reported in a poster at a meeting sponsored by the International Society on Hypertension in Blacks.

Serum selenium is an essential component in substances shown to mediate the incidence of cardiovascular disease, such as glutathione peroxidase and homocysteine. In 9,881 nonpregnant adults aged 40 years and older who participated in the third National Health and Nutrition Examination Surveys (NHANES III), significant differences in the concentrations of serum selenium were noted between African Americans and whites at the highest and lowest quartile concentrations (see graphic), reported Dr. Ani of Charles Drew University of Medicine and Science, Los Angeles.

On bivariate analysis, there was a significant association between serum selenium concentration and the prevalence of hypertension and other cardiovascular disease, including peripheral vascular disease, myocardial infarction, and congestive heart failure. An analysis that controlled for known predictors of cardiovascular disease, including family history, diabetes, renal disease, and sociodemographic variables, showed a significant relationship between serum selenium and the prevalence of hypertension (odds ratios 1.30), as well as a significant interaction effect between ethnicity and serum selenium among individuals with hypertension (odds ratio 1.10).

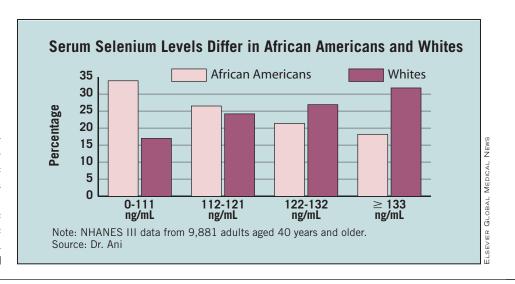
These findings are important because African Americans have higher rates of hypertension and mortality from heart disease and stroke than do whites and Hispanics in the United States, and because African American men have three times the risk of sudden death as do white men.

"Inquiry into biomarkers [that may be] predictors of differential risk and incidence, particularly at the population level, may provide useful explanatory insight regarding the differential burden on cardiovascular disease among African Americans," Dr. Ani wrote.

Based on the emerging understanding of the role of serum selenium in hypertension and cardiovascular disease, and the differing concentrations in African Americans and whites, Dr. Ani and his colleagues theorized that high serum concentrations of selenium might predict reduced levels of oxidate stress and vascular injury in certain ethnic groups that correlates with the incidence of cardiovascular diseases.

The current findings of a statistically significant interaction between serum sele-

nium concentration and ethnicity in individuals with hypertension appear to support this theory of "differential oxidative protection for cardiovascular injury" in African Americans, compared with whites, he said in an interview, adding that the findings are of particular interest because low serum selenium concentration is a modifiable risk factor.







STROKE

Proven protection demonstrated in:

broad range of patients

- CAPRIE: Reduced the risk of combined end point of MI, ischemic stroke, or vascular death in recent MI, recent stroke, or established PAD\*
- CURE: Reduced the risk of combined end point of MI, stroke, or CV death in UA/NSTEMI<sup>†</sup>
- COMMIT: Reduced the risk of all-cause mortality and combined end point of death, reinfarction, or stroke in STEMI<sup>‡</sup>
- CLARITY: Reduced the odds of combined end point of occluded infarct-related artery on predischarge angiogram; or death or recurrent MI before angiography in STEMI<sup>‡</sup>

## Indications

 $Plavix^{@} \ (clopidogrel\ bisulfate)\ is\ indicated\ for\ the\ reduction\ of\ atherothrombotic\ events\ as\ follows:$ 

## Recent MI, Recent Stroke, or Established Peripheral Arterial Disease

\*For patients with a history of recent myocardial infarction (MI), recent stroke, or established peripheral arterial disease, PLAVIX has been shown to reduce the rate of a combined end point of new ischemic stroke (fatal or not), new MI (fatal or not), and other vascular death.

## **Acute Coronary Syndrome**

<sup>†</sup>For patients with non–ST-segment elevation acute coronary syndrome (unstable angina/non–Q-wave MI), including patients who are to be managed medically and those who are to be managed with percutaneous coronary intervention (with or without stent) or CABG, PLAVIX has been shown to decrease the rate of a combined end point of cardiovascular death, MI, or stroke as well as the rate of a combined end point of cardiovascular death, MI, stroke, or refractory ischemia.

<sup>‡</sup>For patients with ST-segment elevation acute myocardial infarction, PLAVIX has been shown to reduce the rate of death from any cause and the rate of a combined end point of death, reinfarction, or stroke. This benefit is not known to pertain to patients who receive primary angioplasty.