# The effect of metabolic syndrome on cancer mortality among blacks and whites in the US 

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## Background

* A black-white disparity in total cancer death has been pervasive in the US despite the general decline in cancer mortality rates due to reduced tobacco smoking, more widespread cancer screening and testing, and improved therapies
* One factor that might contribute to the racial differences in cancer death that has not been fully evaluated is metabolic syndrome (MS).
* MS is highly prevalent among US adults, especially in racial minorities, and it is associated with increased risks of many cancers and cancer mortality
* Previous studies exploring MS or its components and cancer outcomes often aggregated racial groups, thereby masking possible differences in association by race.


## Methods

* We used data from adult participants from NHANES III (1988-1994) ( $\mathrm{N}=18,001$ ).
* We ascertained cancer death from NHANES III mortality follow-up study, which linked with the National Death Index and provides follow-up from survey baseline through December 2006.
$*$ MS was defined as $\geq 3$ of 5 risk factors: elevated triglycerides ( $\geq 150 \mathrm{mg} / \mathrm{dL}$ ), impaired fasting blood glucose ( $\geq 100 \mathrm{mg} / \mathrm{dL}$ ), increased waist circumference ( $\geq 88 \mathrm{~cm}$ for women and $\geq 102 \mathrm{~cm}$ for men), elevated blood pressure (BP) ( $\geq 130 \mathrm{mmHg}$ systolic BP or $\geq 85$ mmHg diastolic BP) and reduced HDL ( $<50 \mathrm{mg} / \mathrm{dL}$ ).
* The interaction between race and MS and its components against total cancer mortality was tested. We used Cox proportional hazards regression to estimate hazard ratios (HR) and 95\% confidence intervals for cancer mortality in relation to MS, MS individual component, and MS category ( $\leq 1,2,3$ or $\geq 4$ components) in whites and blacks.


## Purpose

* This study sought to examine the association between MS and its individual components and total cancer mortality in whites vs. blacks, separately
Table 1. Characteristics of the study population by race, NHANES III

|  | Non-Hispanic White [ $\mathrm{n}=11081, \%$ (SE)] | Non-Hispanic Black [ $\mathrm{n}=4722$, \% (SE)] |
| :---: | :---: | :---: |
| Age |  |  |
| $\geq 20$ to <30 | 20.90 (0.94) | 26.25 (0.87) |
| $\geq 30$ to <40 | 23.81 (0.86) | 26.76 (0.80) |
| $\geq 40$ to <50 | 20.62 (0.79) | 20.57 (0.87) |
| $\geq 50$ to <60 | 12.79 (0.47) | 10.77 (0.60) |
| $\geq 60$ to <70 | 11.64 (0.55) | 8.71 (0.61) |
| $\geq 70$ | 10.22 (0.69) | 6.93 (0.60) |
| Gender |  |  |
| Male | 48.84 (0.43) | 44.90 (0.91) |
| Female | 51.15 (0.43) | 55.09 (0.94) |
| Poverty to income ratio |  |  |
| $\leq 1$ (under poverty) | 9.42 (0.70) | 26.52 (1.73) |
| $>1$ to <3 | 38.32 (1.15) | 43.67 (1.29) |
| $\geq 3$ | 52.27 (1.36) | 29.80 (1.37) |
| Smoking |  |  |
| Never | 44.08 (0.91) | 50.37 (1.09) |
| Former | 27.52 (0.63) | 16.10 (0.68) |
| Current | 28.41 (0.92) | 33.52 (1.04) |
| Insurance |  |  |
| Yes | 88.27 (0.75) | 84.49 (1.48) |
| No | 11.73 (0.75) | 15.51 (1.48) |
| Metabolic syndrome (MS) |  |  |
| Yes ( $\geq 3$ components) | 23.11 (0.82) | 21.84 (0.58) |
| Central obesity | 39.64 (0.82) | 44.66 (1.09) |
| High blood pressure | 10.33 (0.45) | 17.77 (0.75) |
| Low HDL | 36.01 (1.21) | 26.86 (0.87) |
| High triglycerides | 36.03 (1.14) | 26.84 (0.81) |
| Impaired fasting glucose | 28.30 (1.45) | 33.53 (0.84) |
| Number of MS components |  |  |
| $\leq 1$ | 54.71 (1.33) | 53.78 (0.70) |
| 2 | 22.18 (0.81) | 24.38 (0.69) |
| 3 | 15.16 (0.51) | 14.55 (0.45) |
| $\geq 4$ | 7.95 (0.47) | 7.29 (0.40) |

## Results

Table 2. Hazard ratios (HR) for cancer death by metabolic syndrome and MS components in non-Hispanic whites and non-Hispanic blacks, separately.

Non-Hispanic White
Non-Hispanic Black

|  | Non-Hispanic White |  | Non-Hispanic Black |  |
| :---: | :---: | :---: | :---: | :---: |
|  | Crude Model HR (95\% CI) | Adjusted Model HR (95\% CI) ${ }^{1}$ | Crude Model HR (95\% CI) | Adjusted Model HR (95\% CI) ${ }^{1}$ |
| Metabolic syndrome ( $\geq 3$ factors) |  |  |  |  |
| No | Ref. | Ref. | Ref. | Ref. |
| Yes | 2.22 (1.82-2.71)** | 1.19 (0.99-1.44) | 1.61 (1.24-2.11)** | 0.88 (0.67-1.16) |
| Central obesity |  |  |  |  |
| No | Ref. | Ref. | Ref. | Ref. |
| Yes | 2.26 (1.83-2.81)** | 1.29 (1.05-1.59)* | 1.18 (0.94-1.49) | 0.83 (0.62-1.04) |
| High blood pressure |  |  |  |  |
| No | Ref. | Ref. | Ref. | Ref. |
| Yes | 1.55 (1.22-1.97)* | 1.15 (0.91-7.91) | 2.15 (1.68-2.75)** | 1.41 (1.10-1.80)* |
| Low HDL |  |  |  |  |
| No | Ref. | Ref. | Ref. | Ref. |
| Yes | 1.36 (1.12-1.63)* | 1.26 (1.04-1.52)* | 0.94 (0.70-1.24) | 0.91 (0.69-1.20) |
| High triglycerides |  |  |  |  |
| No | Ref. | Ref. | Ref. | Ref. |
| Yes | 1.57 (1.28-1.93)** | 1.02 (0.84-1.23) | 1.47 (1.11-1.94)** | 0.91 (0.70-1.19) |
| Impaired fasting glucose |  |  |  |  |
| No | Ref. | Ref. | Ref. | Ref. |
| Yes | 2.40 (1.91-3.02)** | 1.45 (1.19-1.76)* | 1.66 (1.32-2.08)** | 1.03 (0.80-1.32) |
| No. of MS factors |  |  |  |  |
| $\leq 1$ | Ref. | Ref. | Ref. | Ref. |
| 2 | 2.00 (1.58-2.55)** | 1.36 (1.07-1.73)* | 1.30 (0.91-1.85) | 1.01 (0.70-1.46) |
| 3 | 2.57 (2.03-3.27)** | 1.35 (1.05-1.72)* | 1.61 (1.09-2.40)* | 1.02 (0.68-1.51) |
| $\geq 4$ | 3.38 (2.35-4.86)** | 1.60 (1.13-2.27)* | $2.09(1.36-3.21)^{*}$ | 1.00 (0.65-1.53) |
| P for trend -MS categories | <0.0001 | 0.01 | <0.0001 | 0.91 |
| **<0.005; *<0.05. <br> ${ }^{1}$ Adjusted model included Cox-proportional hazards re | gender, income, insura sion. | status, and smoking | status; sampling weigh | has been considere |

## Conclusions/Discussion

* We found the effect of metabolic risk factors on total cancer mortality differed by race. High BP was significantly associated with total cancer death in blacks while in whites, central obesity, low HLD and impaired fasting glucose were positively associated with cancer death. * The results highlight the importance of early detection and management of metabolic risks, esp. hypertension and prediabetes/diabetes, to prevent cancer death in blacks and whites, respectively.
* Active monitoring of BP throughout the period of cancer treatment is recommended for cancer patients, especially for black cancer patients with elevated BP. Awareness and management of the glycemic status should also be part of patients' cancer care plans, particularly for white cancer patients with prediabetes/diabetes.
* Further studies need to investigate 1) whether overall MS, or individual metabolic risk factors, is associated with death from certain types of cancer and 2) whether the association varies by race.

