

4-15-2016

Usefulness of Relative Hypochromia in Risk Stratification for Nonanemic Patients With Chronic Heart Failure

Muhammad Hammadah
Emory University

Marie Luise Brennan
Epinomics

Yuping Wu
Cleveland State University, y.wu88@csuohio.edu

Stanley L. Hazen
Cleveland State University, S.HAZEN@csuohio.edu

W.H. Wilson Tang
Heart and Vascular Institute

Follow this and additional works at: https://engagedscholarship.csuohio.edu/scimath_facpub

 Part of the [Mathematics Commons](#)

[How does access to this work benefit you? Let us know!](#)

Repository Citation

Hammadah, Muhammad; Brennan, Marie Luise; Wu, Yuping; Hazen, Stanley L.; and Tang, W.H. Wilson, "Usefulness of Relative Hypochromia in Risk Stratification for Nonanemic Patients With Chronic Heart Failure" (2016). *Mathematics Faculty Publications*. 247.
https://engagedscholarship.csuohio.edu/scimath_facpub/247

This Article is brought to you for free and open access by the Mathematics and Statistics Department at EngagedScholarship@CSU. It has been accepted for inclusion in Mathematics Faculty Publications by an authorized administrator of EngagedScholarship@CSU. For more information, please contact library.es@csuohio.edu.

Usefulness of Cardiac Biomarker Score for Risk Stratification in Stable Patients Undergoing Elective Cardiac Evaluation Across Glycemic Status

W.H. Wilson Tang, MD, Naveed Iqbal, MD, Yuping Wu, PhD, and Stanley L. Hazen, MD, PhD

Increasingly, cardiac biomarkers have provided important information in predicting short-term and long-term risk profiles in patients *with* acute coronary syndromes, particularly when used in combination.¹ Several clinically available cardiac biomarkers, including B-type natriuretic peptide (BNP),² myeloperoxidase (MPO),³ and high-sensitivity C-reactive protein (hsCRP),⁴ provide incremental prognostic value in patients with acute coronary syndromes, alone or in combination. Their ability to predict cardiovascular risk has been postulated, as they reflect underlying biomarkers of myocardial dysfunction, plaque vulnerability, and systemic inflammation, respectively.⁵ However, the clinical utility of such biomarkers simultaneously in a stable, nonacute patient cohort is less well established. We hypothesized that simultaneous assessment of these clinically available cardiac biomarkers to produce a risk score (composed of the sums of "positive" biomarkers on the basis of established cutoff values) would provide incremental prognostic insight into predicting future adverse cardiovascular outcomes. As there is an evolving understanding of patients with diabetes and prediabetes being at heightened cardiovascular risk, we further analyzed the prognostic utility of these cardiac biomarkers across the spectrum of glycemic control.

Methods

We prospectively evaluated **3,635** consecutively consenting subjects who underwent elective cardiac catheterization recruited from 2001 to 2006 without evidence of myocardial infarction (cardiac troponin I <**0.03** ng/ml). All participants gave written informed consent, and the institutional review board of the Cleveland Clinic approved the study protocol. The Framingham risk score was calculated for each subject on the basis of Adult Treatment Panel ID guidelines.⁶ An estimate of creatinine clearance was calculated using the Cockcroft-Gault equation. Coronary artery

Table 1
Baseline characteristics

Variable	Whole Cohort (n 3,635)	Diabetes Mellitus (n 1,014)	Prediabetes (n 1,529)	Nondiabetes (n 1,092)	p Value
Age (yrs)	63 ± 11	64 ± 10	63 ± 11	61 ± 12	<0.001
Men	65%	61%	70%	64%	<0.001
Hypertension	71%	78%	70%	62%	<0.001
History of myocardial infarction	33%	35%	32%	32%	<0.150
Systolic blood pressure (mm Hg)	133 (120–146)	134 (120–149)	132 (120–145)	132 (119–147)	<0.015
Low density lipoprotein cholesterol (mg/dl)	95 (78–116)	95 (77–115)	97 (80–118)	94 (76–116)	<0.012
High density lipoprotein cholesterol (mg/dl)	34 (28–41)	32 (27–39)	34 (28–42)	34 (29–42)	<0.001
Creatinine clearance (ml/min/1.73 m ²)	100 (76–126)	99 (74–128)	100 (77–126)	100 (79–126)	<0.512
Cigarette smoking	65%	64%	68%	62%	<0.002
Aspirin	73%	73%	74%	71%	<0.357
β blockers	61%	65%	62%	56%	<0.001
Angiotensin converting enzyme inhibitors or angiotensin receptor blockers	50%	60%	47%	41%	<0.001
Statins	59%	63%	59%	54%	<0.001
hsCRP (mg/L)	2.00 (0.91–4.47)	2.56 (1.13–5.93)	1.89 (0.86–3.95)	1.67 (0.83–4.00)	<0.001
BNP (pg/ml)	83 (34–200)	93 (40–240)	78 (32–177)	83 (32–198)	<0.001
MPO (pmol/L)	103 (70–195)	105 (74–186)	104 (69–201)	100 (68–194)	<0.199

Data are expressed as mean ± SD, as percentages, or as median (interquartile range).

Table 2
Cox proportional hazards analyses for individual cardiac biomarker and future major adverse cardiovascular events at 3 years

Variable	Univariate Model		Multivariate Model*	
	HR (95% CI)	p Value	HR (95% CI)	p Value
BNP >100 pg/ml	2.76 (2.25–3.39)	<0.001	2.10 (1.65–2.68)	<0.001
hsCRP >2 ng/L	2.10 (1.71–2.58)	<0.001	1.82 (1.46–2.28)	<0.001
MPO >322 pmol/L	1.43 (1.12–1.82)	<0.004	1.32 (1.02–1.71)	<0.036

* Adjusted for age, gender, low density and high density lipoprotein cholesterol, systolic blood pressure, former or current cigarette smoking, diabetes mellitus, history of myocardial infarction, and creatinine clearance.

disease was defined as any clinical history of myocardial infarction, percutaneous coronary intervention, coronary artery bypass surgery, or angiographic evidence of coronary artery disease (≥50% stenosis) in ≥1 major coronary artery. Glycemic status and clinical definition of diabetes mellitus, “prediabetes,” and nondiabetes were defined by the latest practice guidelines on the basis of fasting glucose and glycosylated hemoglobin (HbA_{1c}) levels (fasting glucose <100 mg/dl and HbA_{1c} <5.7% for normal subjects; fasting glucose ≥126 mg/dl or HbA_{1c} ≥6.5% or currently taking glucose-lowering medications for diabetes mellitus; neither normal nor diabetes mellitus for prediabetes).⁷ Adjudicated outcomes were ascertained over the ensuing 3 years for all subjects after enrollment. The prospective determination of clinical outcomes was made by the research personnel contacting participants independent of study investigators, with prespecified criteria and confirmation by review of documentation independent of the investigators. Major adverse cardiovascular events (MACEs) were defined as death, nonfatal myocardial infarction, or nonfatal cerebrovascular accident after enrollment. Blood samples were collected before the administration of heparin, placed on ice, aliquoted, and frozen at –80°F within 2 hours of collection.

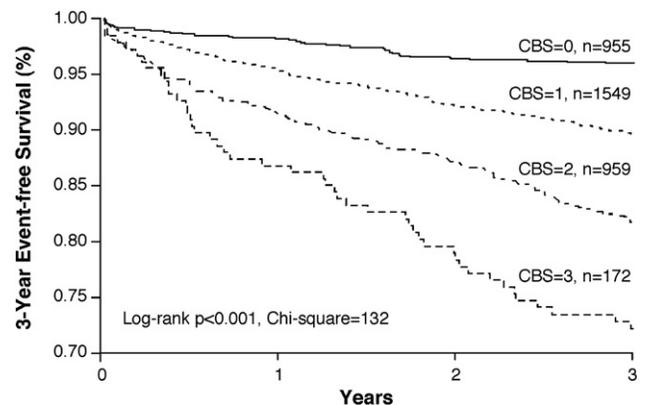


Figure 1. Kaplan Meier analysis of CBS predicting future MACEs at 3 year follow up.

All laboratory assays, including hsCRP, BNP, MPO, apolipoprotein A1, apolipoprotein B100, and creatinine, were performed using the Abbott ARCHITECT ci8200 platform (Abbott Diagnostics Inc., Abbott Park, Illinois). The intra- and interassay coefficients were 4% and 2.4% for hsCRP, 2.6% and 3.5% for BNP, and 6.2% and 4.1% for MPO, respectively.

Student’s *t* tests or Wilcoxon’s rank-sum tests for continuous variables and chi-square tests for categorical variables were used to examine the differences between the groups. A cardiac biomarker score (CBS) was given to each group on the basis of whether it had a positive value for each respective biomarker. We used cutoffs for each of the 3 biomarkers (BNP >100 pg/ml, hsCRP >2.0 ng/L, and MPO >322 pmol/L) on the basis of previous cutoffs used for the respective markers, as reported in previous studies.^{2,4,8} Each of the groups was categorized as 0, 1, 2, or 3 as a measure of how many biomarkers were deemed positive, which was defined as the CBS. Kaplan-Meier analysis with Cox proportional-hazards

Table 3
Cox proportional hazards analyses of cardiac biomarker score stratified by glycemic status

Variable	CBS			
	0	1	2	3
Whole cohort (n = 3,635)				
Unadjusted HR	1	2.59 (1.82–3.68)*	4.72 (3.33–6.69)*	7.61 (4.98–11.65)*
Adjusted HR, model 1 (95% CI)	1	2.27 (1.59–3.23)*	3.67 (2.58–5.24)*	6.11 (3.98–9.38)*
Adjusted HR, model 2 (95% CI)	1	2.27 (1.59–3.23)*	3.67 (2.58–5.24)*	6.11 (3.98–9.38)*
MACEs	39/955	159/1,549	173/959	47/172
Normal (n = 1,014)				
Unadjusted HR	1	1.78 (0.94–3.37)	3.23 (1.69–6.16)*	6.23 (2.9–13.37)*
Adjusted HR, model 1 (95% CI)	1	1.43 (0.74–2.75)	2.37 (1.21–4.64)*	4.73 (2.18–10.25)*
Adjusted HR, model 2 (95% CI)	1	1.36 (0.7–2.61)	2.23 (1.14–4.39)*	4.24 (1.96–9.18)*
MACEs	13/290	35/445	32/230	13/49
Prediabetes (n = 1,529)				
Unadjusted HR	1	2.74 (1.56–4.84)*	4.75 (2.7–8.38)*	10.27 (5.24–20.13)*
Adjusted HR, model 1 (95% CI)	1	2.4 (1.36–4.23)*	3.67 (2.06–6.54)*	7.87 (3.99–15.55)*
Adjusted HR, model 2 (95% CI)	1	2.37 (1.35–4.19)*	3.58 (2–6.41)*	7.62 (3.87–15.01)*
MACEs	15/427	61/654	61/379	20/69
Diabetes mellitus (n = 1,092)				
Unadjusted HR	1	3.17 (1.67–6)*	5.59 (2.98–10.49)*	6.16 (2.8–13.52)*
Adjusted HR, model 1 (95% CI)	1	3.06 (1.61–5.78)*	4.79 (2.55–9)*	5.59 (2.54–12.31)*
Adjusted HR, model 2 (95% CI)	1	3.05 (1.61–5.77)*	4.8 (2.55–9.01)*	5.61 (2.55–12.33)*
MACEs	11/238	63/450	80/350	14/54

Model 1 was adjusted for traditional risk factors, including age, gender, systolic blood pressure, low density lipoprotein cholesterol, high density lipoprotein cholesterol, and smoking. Model 2 was adjusted for traditional risk factors plus apolipoprotein B100/apolipoprotein A1 ratio.

* $p < 0.001$.

regression was used for time-to-event analysis to determine hazard ratios (HRs) and 95% confidence intervals (95% CIs) for MACEs. Unadjusted trends for all-cause mortality rates as well as rates of nonfatal myocardial infarction or stroke with increasing quartiles of MPO, hsCRP, and BNP were evaluated using the Cochran-Armitage test using a time-to-event approach. Adjustments were made for individual traditional cardiac risk factors (including age, gender, low-density and high-density lipoprotein cholesterol, systolic blood pressure, former or current cigarette smoking, diabetes mellitus, apolipoprotein B100/apolipoprotein A1 ratio, history of myocardial infarction, and creatinine clearance) to predict incident 3-year risk for MACEs. Net reclassification analysis was performed with the 2 Cox models adjusted for traditional risk factors. Cutoff values for net reclassification index estimation used a ratio of 6:3:1 for low-, medium-, and high-risk categories. All analyses were performed using R version 8.02 (R Foundation for Statistical Computing, Vienna, Austria), and p values < 0.05 were considered statistically significant.

Results

Table 1 describes the baseline characteristics of our study population and is stratified according to glycemic status. The median levels of hsCRP, BNP, and MPO were 2.00 pg/ml (interquartile range 0.91 to 4.47), 83 pg/ml (interquartile range 34 to 200), and 103 pmol/L (interquartile range 70 to 195), respectively. All 3 biomarkers were notably elevated in patients with diabetes compared to those with prediabetes or nondiabetics.

Table 2 represents the prognostic value of individual cardiac biomarkers in our study cohort. All 3 cardiac

biomarkers provided incremental risk prediction in our study cohort. After adjusting for traditional risk factors, including Framingham risk factors, log-transformed BNP, hsCRP, and MPO each remained independent predictors of incident MACEs at 3-year follow-up.

By summing the number of positive cardiac biomarkers, we developed a CBS that integrated the risk profile of our study cohort. As illustrated in Figure 1, the CBS provided incremental prognostic value, as displayed by Kaplan-Meier analysis. As listed in Table 3, the CBS, based on the sum total of positive biomarkers, provided independent prediction of future risk for incident MACEs at 3 years (HR 7.61, 95% CI 4.98 to 11.65, $p < 0.001$), even after adjustment for traditional risk factors (HR 6.11, 95% CI 3.98 to 9.38, $p < 0.001$), in addition to apolipoprotein B100/apolipoprotein A1 ratio (HR 6.11, 95% CI 3.98 to 9.38, $p < 0.001$) (Table 3). The ability of the CBS to provide incremental risk stratification can be seen in subgroups of patients with primary and secondary prevention, as well as those with maximal coronary artery stenoses $< 50\%$ and $\geq 50\%$ (Figure 2). A higher CBS predicted future risk for MACEs at 3 years regardless of age, gender, body mass index, diabetes mellitus, hypertension, renal insufficiency, or previous myocardial infarction ($p < 0.01$ for all). Use of the CBS on top of traditional risk factors was also shown to reclassify subjects (net reclassification index 12.86%, $p < 0.001$; integrated discrimination improvement 12.0%, $p < 0.001$; C-statistic 66.9% vs 71.1%, $p < 0.001$).

We further analyzed our study cohort according to glycemic status and different subgroups. The capability of the CBS to stratify patients' risk profiles within subjects with diabetes mellitus, prediabetes, or normal glycemic status (neither prediabetes nor diabetes) on the basis of practice

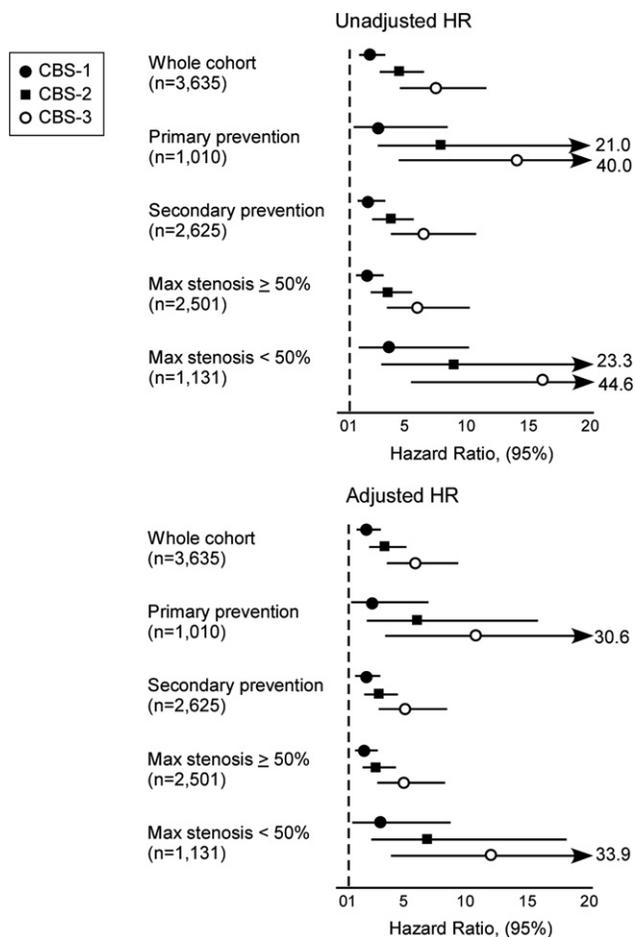


Figure 2. Forest plot of unadjusted and adjusted HRs for predicting future MACEs at 3 year follow up according to CBS according to subgroups (zero score as reference, adjustments as in Table 3, model 1).

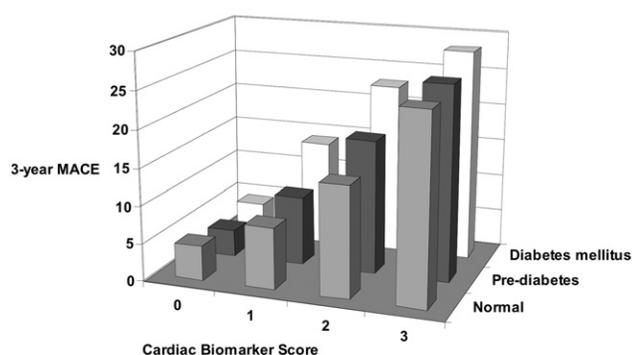


Figure 3. Event rates for future MACEs at 3 year follow up according to glycemic status.

guidelines⁷ remained robust (Table 3, Figure 3). In a similar manner, after adjustment for HbA_{1c}, the prognostic value of the CBS was preserved. Furthermore, the prognostic value of the CBS was similar regardless of age, gender, body mass index, diabetes mellitus, hypertension, renal insufficiency, or previous myocardial infarction ($p < 0.01$ for all).

Discussion

Although previous studies have examined similar multi-marker strategies for risk prediction, many of them used biomarkers that are not commonly used or available in the clinical practice settings. The key finding in this study is the incremental prognostic value of all 3 clinically available plasma cardiac biomarkers beyond standard evaluation of classic Framingham risk factors, renal function, and apolipoprotein B100/apolipoprotein A1 ratio in troponin-negative, stable cardiac patients who undergo coronary angiography. We further identified comparable prognostic value within subsets of patients with prediabetes or nondiabetes or those with no significantly obstructive coronary artery disease, further underscoring the potential for a multimarker approach in identifying vulnerable patients within cohorts that may allow targeted risk factor modifications and more aggressive preventive interventions.

HsCRP is the most common systemic inflammatory biomarker used in clinical practice, particularly in patients with diabetes mellitus and potential response to statin therapy.^{4,9,11} Elevated levels have also been associated with altered cardiac structure and function, as well as adverse long-term consequences.¹² In addition, hsCRP has been suggested to play a role in atherosclerosis and its complications, although genetic studies suggest that the association with adverse outcomes may not be causal.^{13,14} In contrast, MPO has been shown to directly promote the catalytic consumption of nitric oxide, leading to the development of endothelial dysfunction.^{15,17} MPO is a leukocyte-derived hemoprotein that has been linked in the development and subsequent instability of atherosclerotic plaques.^{18,19} Previous studies have shown MPO to have prognostic significance in subjects with unstable angina, as well as after acute myocardial infarction, acute heart failure, and chronic stable heart failure, as well as healthy middle-aged and elderly subjects.^{3,18,20,23} Recently, it was also found that MPO remained a statistically significant prognostic indicator of cardiovascular risk in a large stable CAD population.⁸ In contrast, the natriuretic peptide family is a group of endogenous peptides primarily produced in the heart that provide counter-regulatory effects on a wide range of organs to maintain perfusion and reduce overloading status of the vasculature.²⁴ BNP has recently been shown to be elevated in acute coronary syndromes without necessarily having myocardial infarction, and it may reflect not only the underlying impairment of left ventricular function but also the severity of the ischemic episode.²⁵ Altogether, this combination of biomarkers offers complementary mechanistic insights during cardiac evaluation in stable patients, although only a small subset of patients demonstrated positive results for all 3 biomarkers in our relatively stable patient cohort.

Our study further explored the impact of glycemic control on the prognostic value of cardiac biomarkers, particularly as the latest guidelines have highlighted a subset of “at-risk” patients that is thought to have heightened risk for developing diabetes mellitus and future cardiovascular risk.^{7,26} We observed a graded increase in the levels of each of our corresponding biomarkers as patients were determined to have nondiabetes, prediabetes, and diabetes. The

overall trend toward increased risk for MACEs was similar among all groups on the basis of their CBS. Similarly, the CBS provided significant prognostic value among subjects for whom no significant angiographic evidence of stenosis was discovered, who are most often considered to have lower risk.

The strength of this study is the considerable size of the patient population. This contemporary cohort of stable cardiac patients is representative of current clinical practice. The focus on the homogenous elective coronary angiography population and the availability and inclusion of only biomarkers cleared by the United States Food and Drug Administration for the analyses strengthen the study, as the present biomarkers, although clinically available for use, are not routinely measured. Including them in the analysis provides insight into a nonacute, troponin-negative population, which has yet to be thoroughly investigated. Incremental contributions of these cardiac biomarkers toward risk stratification above and beyond standard clinical and biochemical characteristics in this population have also not been thoroughly tested, particularly with rigorous statistical evaluation or covariate adjustments. Potential weaknesses of the study population arise because a clinical trial cohort of patients who underwent coronary angiography was used, and thus resulting in, particularly because they were already undergoing cardiac evaluation. Moreover, our data relate to prognostic rather than diagnostic applications of these biomarkers. We also did not have high-sensitivity troponin assays in this cohort, which was deemed troponin negative by currently approved troponin assays. Last, although our results were based on previously used cut points, they may overestimate the strengths of the risk relations. With different studies using different cutoff values for different populations, there is a need to identify clinically useful cut points on the basis of consensus of results.

Disclosures

Dr. Tang received research grant support from Abbott Laboratories, Abbott Park, Illinois. Dr. Hazen is a co-inventor on pending and issued patents held by the Cleveland Clinic relating to cardiovascular diagnostics. Dr. Hazen has received consulting fees from AstraZeneca Pharmaceuticals, Wilmington, Delaware; Cleveland HeartLab, Cleveland, Ohio; Eli Lilly & Company, Indianapolis, Indiana; Esperion Therapeutics, Plymouth, Michigan; LipoScience Inc., Raleigh, North Carolina; Merck & Company, Whitehouse Station, New Jersey, and Pfizer, Inc., New York, New York. Dr. Hazen has received research funding from Abbott Laboratories, Cleveland HeartLab, Esperion Therapeutics, and LipoScience Inc. Dr. Hazen has the right to receive royalty payments for inventions or discoveries related to cardiovascular diagnostics from Abbott Laboratories; Cleveland HeartLab; Frantz BioMarkers, LLC, Mentor, Ohio; LipoScience Inc.; and Siemens Healthcare, Erlangen, Germany.

1. Sabatine MS, Morrow DA, de Lemos JA, Gibson CM, Murphy SA, Rifai N, McCabe C, Antman EM, Cannon CP, Braunwald E. Multi marker approach to risk stratification in non ST elevation acute coronary syndromes: simultaneous assessment of troponin I, C reactive

protein, and B type natriuretic peptide. *Circulation* 2002;105:1760–1763.

2. de Lemos JA, Morrow DA, Bentley JH, Omland T, Sabatine MS, McCabe CH, Hall C, Cannon CP, Braunwald E. The prognostic value of B type natriuretic peptide in patients with acute coronary syndromes. *N Engl J Med* 2001;345:1014–1021.
3. Brennan ML, Penn MS, Van Lente F, Nambi V, Shishehbor MH, Aviles RJ, Goormastic M, Pepoy ML, McErean ES, Topol EJ, Nissen SE, Hazen SL. Prognostic value of myeloperoxidase in patients with chest pain. *N Engl J Med* 2003;349:1595–1604.
4. Ridker PM, Hennekens CH, Buring JE, Rifai N. C reactive protein and other markers of inflammation in the prediction of cardiovascular disease in women. *N Engl J Med* 2000;342:836–843.
5. Morrow DA, Braunwald E. Future of biomarkers in acute coronary syndromes: moving toward a multimarker strategy. *Circulation* 2003;108:250–252.
6. Executive summary of the third report of the National Cholesterol Education Program (NCEP) Expert Panel on Detection, Evaluation, and Treatment of High Blood Cholesterol in Adults (Adult Treatment Panel III). *JAMA* 2001;285:2486–2497.
7. Standards of medical care in diabetes 2012. *Diabetes Care* 2012;35(suppl):S11–S63.
8. Tang WH, Wu Y, Nicholls SJ, Hazen SL. Plasma myeloperoxidase predicts incident cardiovascular risks in stable patients undergoing medical management for coronary artery disease. *Clin Chem* 2011;57:33–39.
9. Ridker PM, Danielson E, Fonseca FA, Genest J, Gotto AM Jr, Kastelein JJ, Koenig W, Libby P, Lorenzatti AJ, MacFadyen JG, Nordestgaard BG, Shepherd J, Willerson JT, Glynn RJ. Rosuvastatin to prevent vascular events in men and women with elevated C reactive protein. *N Engl J Med* 2008;359:2195–2207.
10. Ridker PM, Rifai N, Clearfield M, Downs JR, Weis SE, Miles JS, Gotto AM Jr. Measurement of C reactive protein for the targeting of statin therapy in the primary prevention of acute coronary events. *N Engl J Med* 2001;344:1959–1965.
11. Ridker PM, Rifai N, Rose L, Buring JE, Cook NR. Comparison of C reactive protein and low density lipoprotein cholesterol levels in the prediction of first cardiovascular events. *N Engl J Med* 2002;347:1557–1565.
12. Tang WH, Shrestha K, Van Lente F, Troughton RW, Martin MG, Borowski AG, Jasper S, Klein AL. Usefulness of C reactive protein and left ventricular diastolic performance for prognosis in patients with left ventricular systolic heart failure. *Am J Cardiol* 2008;101:370–373.
13. Anand SS, Yusuf S. C reactive protein is a bystander of cardiovascular disease. *Eur Heart J* 2010;31:2092–2096.
14. Elliott P, Chambers JC, Zhang W, Clarke R, Hopewell JC, Peden JF, Erdmann J, Braund P, Engert JC, Bennett D, Coin L, Ashby D, Tzoulaki I, Brown IJ, Mt Isa S, McCarthy MI, Peltonen L, Freimer NB, Farrall M, Ruokonen A, Hamsten A, Lim N, Froguel P, Waterworth DM, Vollenweider P, Waeber G, Jarvelin MR, Mooser V, Scott J, Hall AS, Schunkert H, Anand SS, Collins R, Samani NJ, Watkins H, Kooper JS. Genetic loci associated with C reactive protein levels and risk of coronary heart disease. *JAMA* 2009;302:37–48.
15. Hazen SL, Zhang R, Shen Z, Wu W, Podrez EA, MacPherson JC, Schmitt D, Mitra SN, Mukhopadhyay C, Chen Y, Cohen PA, Hoff HF, Abu Soud HM. Formation of nitric oxide derived oxidants by myeloperoxidase in monocytes: pathways for monocyte mediated protein nitration and lipid peroxidation In vivo. *Circ Res* 1999;85:950–958.
16. Schmitt D, Shen Z, Zhang R, Colles SM, Wu W, Salomon RG, Chen Y, Chisolm GM, Hazen SL. Leukocytes utilize myeloperoxidase generated nitrating intermediates as physiological catalysts for the generation of biologically active oxidized lipids and sterols in serum. *Biochemistry* 1999;38:16904–16915.
17. Vita JA, Brennan ML, Gokce N, Mann SA, Goormastic M, Shishehbor MH, Penn MS, Keaney JF Jr, Hazen SL. Serum myeloperoxidase levels independently predict endothelial dysfunction in humans. *Circulation* 2004;110:1134–1139.
18. Nicholls SJ, Hazen SL. Myeloperoxidase and cardiovascular disease. *Arterioscler Thromb Vasc Biol* 2005;25:1102–1111.
19. Nicholls SJ, Hazen SL. Myeloperoxidase, modified lipoproteins, and atherogenesis. *J Lipid Res* 2009;50(suppl):S346–S351.
20. Tang WH, Shrestha K, Troughton RW, Borowski AG, Klein AL. Integrating plasma high sensitivity C reactive protein and myeloperoxidase for risk prediction in chronic systolic heart failure. *Congest Heart Fail* 2011;17:105–109.

21. Nicholls SJ, Tang WH, Brennan D, Brennan ML, Mann S, Nissen SE, Hazen SL. Risk prediction with serial myeloperoxidase monitoring in patients with acute chest pain. *Clin Chem* 2011;57:1762–1770.
22. Reichlin T, Socrates T, Egli P, Potocki M, Breidthardt T, Arenja N, Meissner J, Noveanu M, Reiter M, Twerenbold R, Schaub N, Buser A, Mueller C. Use of myeloperoxidase for risk stratification in acute heart failure. *Clin Chem* 2010;56:944–951.
23. Naruko T, Furukawa A, Yunoki K, Komatsu R, Nakagawa M, Matsumura Y, Shirai N, Sugioka K, Takagi M, Hozumi T, Itoh A, Haze K, Yoshiyama M, Becker AE, Ueda M. Increased expression and plasma levels of myeloperoxidase are closely related to the presence of angiographically detected complex lesion morphology in unstable angina. *Heart* 2010;96:1716–1722.
24. Tang WH. B type natriuretic peptide: a critical review. *Congest Heart Fail* 2007;13:48–52.
25. Fonarow GC, Peacock WF, Phillips CO, Givertz MM, Lopatin M, ADHERE Scientific Advisory Committee and Investigators. Admission B type natriuretic peptide levels and in hospital mortality in acute decompensated heart failure. *J Am Coll Cardiol* 2007;49:1943–1950.
26. Tabak AG, Herder C, Rathmann W, Brunner EJ, Kivimaki M. Prediabetes: a high risk state for diabetes development. *Lancet* 2012;379:2279–2290.