Appendix 1 (as supplied by the authors): Criteria for assigning Edmonton obesity staging system score.

EOSS Score Assigned			
0	1	2	3*
< 5.6	5.6-6.9	7.0 or self-report of diabetes	
(mmol/L)		or self-report of treatment with	
		insulin or antidiabetic agents	
BP < 130/85 with no	For individuals classified	Self-report of hypertension or	
self-report of	as having diabetes or	treatment antihypertensive	
hypertension or	chronic kidney disease,	drugs. For individuals classified	
antihypertensive drug treatment. BP < 125/75 if diabetes	SBP 125-129.9 or DBP 75-	as having diabetes or chronic	
	79.9. Otherwise, SBP 130-	kidney disease*, BP 130/80.	
	139.9 or DBP 85-89.9	Otherwise BP 140/90.	
or chronic kidney			
disease present.			
< 3.4	3.4-4.0	4.1	
< 5.2	5.2-6.1	6.2	
1.6	1.0-1.6	< 1.0	
< 1.7	1.7-2.3	2.3	
No self-report of any	Elevated liver enzymes	Elevated liver enzymes and self-	
liver condition and normal liver enzymes	but no self-report of liver	report of liver disease	
	disease		
GFR 90	GFR 60-89.9	GFR 30-59.9	GFR < 30
No history of joint or	Occasional joint or back	Self-report of osteoarthritis	
back pain	pain		
No functional or	Functional impairment	ADL limitations	
ADL limitations	but no ADL limitations		
	< 5.6  BP < 130/85 with no self-report of hypertension or antihypertensive drug treatment. BP < 125/75 if diabetes or chronic kidney disease present.  < 3.4  < 5.2  1.6  < 1.7  No self-report of any liver condition and normal liver enzymes  GFR 90  No history of joint or back pain  No functional or	Self-report of hypertension or antihypertensive drug treatment. BP < 125/75 if diabetes or chronic kidney disease present.  SBP 125-129.9 or DBP 75-79.9. Otherwise, SBP 130-139.9 or DBP 85-89.9  SBP 125-129.9 or DBP 75-79.9. Otherwise, SBP 130-139.9 or DBP 85-89.9  SBP 125-129.9 or DBP 75-79.9. Otherwise, SBP 130-139.9 or DBP 85-89.9  SBP 125-129.9 or DBP 85-89.9  SBP 125-129.9 or DBP 75-79.9. Otherwise, SBP 130-139.9 or DBP 85-89.9  SBP 125-129.9 or DBP 75-79.9. Otherwise, SBP 130-139.9 or DBP 85-89.9  SBP 125-129.9 or DBP 75-79.9. Otherwise, SBP 130-139.9 or DBP 85-89.9  SBP 125-129.9 or DBP 75-79.9. Otherwise, SBP 130-139.9 or DBP 85-89.9  SBP 125-129.9 or DBP 75-79.9. Otherwise, SBP 130-139.9 or DBP 85-89.9  SBP 125-129.9 or DBP 75-79.9. Otherwise, SBP 130-139.9 or DBP 85-89.9  SBP 125-129.9 or DBP 75-79.9. Otherwise, SBP 130-139.9 or DBP 85-89.9  SBP 125-129.9 or DBP 75-79.9. Otherwise, SBP 130-139.9 or DBP 85-89.9  SBP 125-129.9 or DBP 75-79.9 or DBP 75-79.9 or DBP 85-89.9  SBP 125-129.9 or DBP 75-79.9 or DBP 75-79.9 or DBP 85-89.9  SBP 125-129.9 or DBP 75-79.9 or DBP 75-79.9 or DBP 85-89.9  SBP 125-129.9 or DBP 75-79.9 or DBP 75-79.9 or DBP 85-89.9  SBP 125-129.9 or DBP 75-79.9 or DBP 75-79.9 or DBP 85-89.9  SBP 125-129.9 or DBP 75-79.9 or DBP 75-79.9 or DBP 85-89.9  SBP 125-129.9 or DBP 75-79.9 or DBP 75-79.9 or DBP 85-89.9  SBP 125-129.9 or DBP 85-89.9  SBP 125-129.9 or DBP 75-79.9 or DBP 75-79.9 or DBP 85-89.9  SBP 125-129.9 or DBP 75-79.9 or DBP 85-89.9  SBP 125-129.9 or	<ul> <li>&lt; 5.6</li> <li>5.6-6.9</li> <li>7.0 or self-report of diabetes or self-report of treatment with insulin or antidiabetic agents</li> <li>BP &lt; 130/85 with no self-report of as having diabetes or chronic kidney disease, antihypertension or antihypertensive drug treatment. BP</li> <li>&lt; 125/75 if diabetes or chronic kidney disease present.</li> <li>&lt; 3.4</li> <li>3.4-4.0</li> <li>4.1</li> <li>&lt; 5.2</li> <li>5.2-6.1</li> <li>6.2</li> <li>1.0-1.6</li> <li>1.0-1.6</li> <li>&lt; 1.7-2.3</li> <li>2.3</li> <li>No self-report of any liver condition and normal liver enzymes</li> <li>GFR 90</li> <li>GFR 60-89.9</li> <li>GFR 30-59.9</li> <li>Self-report of diabetes or shaving diabetes or chronic kidney disease having diabetes or chronic kidney disease*, BP 130/80. Otherwise BP 140/90.</li> <li>Otherwise BP 140/90.</li> <li>4.1</li> <li>&lt; 1.0</li> <li>&lt; 1.0</li> <li>Self-report of any liver enzymes but no self-report of liver disease</li> <li>GFR 90</li> <li>GFR 60-89.9</li> <li>GFR 30-59.9</li> <li>No history of joint or back pain</li> <li>No functional or</li> <li>Functional impairment</li> <li>ADL limitations</li> </ul>

ADL = activities of daily living, GFR = glomerular filtration rate, HDL = high-density lipoprotein, LDL = low-density lipoprotein.

## **Definitions of obesity-related comorbidities**

Diabetes: Based on self-report, laboratory testing and medication use using currently recommended thresholds.<sup>1</sup>

Hypertension: Based on self-report, medication use and blood pressure measurements.<sup>2</sup>

**Dyslipidemia**: Total cholesterol, LDL-cholesterol, HDL-cholesterol, and triglycerides were categorized according to current recommendations.<sup>3</sup>

Appendix to: Padwal RS, Pajewski N, Allison DB, Sharma AM. Using the Edmonton obesity staging system to predict all-cause mortality in a population-representative cohort of people with overweight and obesity. *CMAJ* 2011. DOI:10.1503/cmaj110387.

<sup>\*</sup> GFR < 60 ml/min/1.73m<sup>2</sup>

<sup>\*\*</sup> Subjects with a history of angina, coronary heart disease, congestive heart failure and cerebrovascular disease were automatically assigned to Stage 3.

**Osteoarthritis:** Based on self-reported "osteoarthritis". Self-reported pain, aching or stiffness or lower back pain in the past year (in the absence of osteoarthritis) was coded as 'occasional joint or back pain'. Subjects reporting joint pain/aching/stiffness following an injury were not considered to have an obesity-related cause and were scored 0 for this category.

**Liver Disease:** Based on the presence or absence of elevated enzymes and/or self-reported liver disease (latter variable available for 1999-2004 cohort only). Non-alcoholic fatty liver disease (NAFLD) is the primary liver-related comorbidity associated with excess adiposity, accounting for 9% of chronic liver disease in the US,<sup>4</sup> but is not directly coded within NHANES. NAFLD, hepatitis B or C, and alcohol-related liver disease account for over 95% of all cases of chronic liver disease within the US and other causes comprise less than 2% of cases.<sup>4</sup> Subjects with a history of elevated liver enzymes or self-reported liver disease were assumed to have NAFLD and assigned an EOSS score unless evidence for an alternate cause was present. Subjects with hepatitis B surface antigen, hepatitis C antibody, a transferrin saturation > 50%, or an average alcohol consumption of at least one (women) or two (men) drinks per day,<sup>5</sup> were considered to have an alternate cause of liver disease and were assigned an EOSS score of 0 for this category.

**Kidney Disease:** Serum creatinine and estimated Glomerular Filtration Rate (eGFR) were determined according to previously described methods.<sup>67</sup>

**Physical Function:** Based on health interview questions examining functional limitation and activities of daily living (ADL). A physical functioning score was calculated for respondents who answered at least 4 of 6 functional limitation questions and at least 2 of 3 ADL questions. Respondents were considered to have a functional limitation if they reported using an assistive device to walk or difficulty with the following tasks: walking ¼ mile, walking up 10 steps without resting, stooping/crouching/kneeling, lifting or carrying 10 lb, walking between rooms on the same floor, or standing from an armless chair. Respondents reporting difficulty/inability with getting in and out of bed, eating, or dressing were considered to have an ADL limitation.

**Metabolic Syndrome:** The ATP III Clinical Criteria, defined as having 3 or more of the following, were used: 1. awist circumference >102 cm (men) or > 88 cm (women), 2. Triglycerides 1.7 mmol/L, 3. HDL cholesterol <1.0 mmol/L (men) or <1.3 mmol/L (women), 4. Blood pressure 130/85 mm Hg, or 5. Fasting glucose 5.6 mmol/L.

**Hypertriglyceridemic waist:** Defined as a waist circumference of 90 cm with a triglyceride level of 2 mmol/L (men), or a waist circumference of 85 cm with a triglyceride level of 1.5 mmol/L (women). 10

## References

- Canadian Diabetes Association. Canadian Diabetes Association 2008 clinical practice guidelines. Toronto (ON): The Association; 2008. Available: www.diabetes.ca/for-professionals/resources/2008-cpg (accessed 2011 June 2).
- Hackam DG, Khan NA, Hemmelgarn BR, et al. The 2010 Canadian Hypertension Education Program recommendations for the management of hypertension: part 2 — therapy. Can J Cardiol 2010;26:249-58.
- 3. National Heart, Lung and, Blood Institute. Third report of the expert panel on detection, evaluation, and treatment of high blood cholesterol in adults (Adult Treatment Panel III). Bethesda (MD): The Institute; 2002. Available: www.nhlbi.nih.gov/guidelines/cholesterol/index.htm (accessed 2011 June 2).
- 4. Kim WR, Brown RS, Terrault NA, et al. Burden of liver disease in the United States: summary of a workshop. *Hepatology* 2002;36:227-42.
- 5. Clark JM, Brancati FL, Diehl AM. The prevalence and etiology of elevated aminotransferase levels in the United States. *Am J Gastroenterol* 2003;98:960-7.
- 6. Selvin E, Manzi J, Stevens L, et al. Calibration of serum creatinine in the National Health and Nutrition Examination Surveys (NHANES) 1988-1994, 1999-2004. *Am J Kidney Dis* 2007;50:918-26.
- 7. Levey A, Stevens L, Schmid C, et al. A new equation to estimate glomerular filtration rate. *Ann Intern Med* 2009;150:604-12.
- 8. Alley DE, Chang VW. The changing relationship of obesity and disability, 1988-2004. *JAMA* 2007;298:2020-7
- 9. Grundy SM, Hansen B, Smith SC, et al.; American Heart Association; National Heart, Lung, and Blood Institute; American Diabetes Association. Clinical management of metabolic syndrome: report of the American Heart Association/National Heart, Lung, and Blood Institute/American Diabetes Association conference on scientific issues related to management. *Circulation* 2004;109:551-6.
- 10. Arsenault BJ, Lemieux I, Després JP, et al. The hypertriglyceridemic-waist phenotype and the risk of coronary artery disease: results from the EPIC-Norfolk prospective population study. *CMAJ* 2010;182:1427-32

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