





# Obesity and kidney function decline in 60-80 years old stateof-the-art drug-treated post-myocardial infarction patients

Kevin Esmeijer<sup>1,2</sup>, Johanna M. Geleijnse<sup>3</sup>, Ph.D., Erik J. Giltay<sup>4</sup>, M.D., Ph.D., Theo Stijnen<sup>5</sup>, Ph.D., Friedo W. Dekker<sup>1</sup>, Ph.D., Johan W. de Fijter<sup>2</sup>, M.D., Ph.D, Daan Kromhout<sup>3,6</sup>, M.P.H., Ph.D., Ellen K. Hoogeveen, M.D., Ph.D.<sup>1,2,7</sup>

1. Dept of Epidemiology, LUMC, Leiden, 2. Dept of Nephrology, LUMC, Leiden, 3. Division of Human Nutrition, Wageningen University, Wageningen, 4. Dept of Psychiatry, LUMC, Leiden, 5. Dept of Medical Statistics and Bioinformatics, LUMC, Leiden, 6. Dept of Medical Sciences, University Medical Center Groningen, 7. Dept of Nephrology, Jeroen Bosch Hospital, Den Bosch, all in the Netherlands

The prevalence of overweight and obesity has increased to epidemic proportions and is globally ranking in the top 5 risk factors for death. Obesity is associated with accelerated decline of kidney function in the general population, kidney transplant and dialysis patients.<sup>2</sup> Obesity may promote kidney damage through both hemodynamic and hormonal effects. Accelerated kidney function decline is associated with increased morbidity and mortality. Little is known about the effect of obesity on kidney function decline in stable post-myocardial infarction (MI) patients, prompting us to investigate this relation.

This prospective cohort study is a follow-up of the Alpha Omega Trial; 2,410 post MI-patients with optimal pharmacotherapeutical treatment, aged 60-80 years (79% men, 18% diabetes), were followed for 40 months. We assessed serum cystatin C (cysC) at baseline and after 40 months.<sup>3,4</sup> We estimated eGFR<sub>cvsC</sub> with the CKD-EPI equation. We used body mass index (BMI) and waist circumference (WC) measured at baseline as an index of adiposity and defined obesity as a BMI ≥30 kg/m<sup>2</sup>. We assessed the effect of BMI and WC on eGFR<sub>cvsC</sub> decline by multivariate linear regression. Results were stratified for sex.

# Objective

To study the relation of overweight and obesity with the rate of kidney function decline in older state-of-the-art drug treated post-myocardial infarction (MI) patients.

### Results

At baseline patients had a mean (SD) BMI of 27.7 (3.6) kg/m<sup>2</sup>, 23% were obese. The baseline mean (SD) eGFR $_{\text{cysC}}$  was 81.5 (19.6) ml/min/1.73m<sup>2</sup>. Annual mean (95%-CI) eGFR $_{\text{cysC}}$  decline was -1.33 ml/min/1.73m<sup>2</sup>.

We found that both BMI (Figure 1) and WC (not shown) were associated with higher kidney function decline (Table 2). After adjustment for age, sex, smoking, alcohol-use and education level, we found:

- for every 5 kg/m $^2$  increment of BMI an additional annual eGFR $_{cvsC}$  decline of -0.28 ml/min/1.73m $^2$  (21%).
- for every 10 cm increment of WC an additional annual eGFR<sub>cvsC</sub> decline of -0.21 ml/min/1.73m $^2$  (16%).
- obese compared to normal weight patients had an extra 23% yearly eGFR  $_{\mbox{\scriptsize cysC}}$  decline of -0.30 ml/min/1.73m<sup>2</sup>.

We found no evidence for effect modification between sex and BMI or WC with regard to kidney function decline.

Table 2: Effect of baseline body mass index and waist circumference on yearly cystatin C based kidney function decline in post-myocardial infarction patients, for men and women separately.

	Annual eGFR <sub>cysC</sub> decline	Annual eGFR <sub>cysC</sub> decline	Annual eGFR <sub>cysC</sub> decline
	mean (95%-CI), total	mean (95%-CI), men	mean (95%-CI), women
	n = 2,410	n = 1,914	n = 496
Mean annual eGFR <sub>cysC</sub> decline	-1.33 (-1.46 to -1.21)	-1.45 (-1.59 to -1.31)	-0.89 (-1.21 to -0.58)
Per 5 kg/m <sup>2</sup> increment of BMI			
Crude	-0.20 (-0.38 to -0.02)	-0.27 (-0.48 to -0.06)	-0.15 (-0.49 to 0.19)
Model 1	-0.29 (-0.46 to -0.11)	-0.36 (-0.57 to -0.15)	-0.16 (-0.50 to 0.18)
Model 2	-0.28 (-0.46 to -0.10)	-0.35 (-0.56 to -0.14)	-0.21 (-0.56 to 0.14)
Per 10 cm increment of WC			
Crude	-0.24 (-0.37 to -0.11)	-0.20 (-0.35 to -0.05)	-0.19 (-0.47 to 0.08)
Model 1	-0.21 (-0.34 to -0.08)	-0.22 (-0.37 to -0.06)	-0.20 (-0.47 to 0.08)
Model 2	-0.21 (-0.34 to -0.08)	-0.21 (-0.37 to -0.06)	-0.22 (-0.50 to 0.05)

Model 1: adjusted for four randomized treatment groups (3 dummy variables), age, and sex. Model 2: model 1, additionally adjusted for lifestyle variables; smoking (yes/no), alcohol use (yes/no), level of education.

## Conclusion

We found that both body-mass index and waist circumference had a progressive relation with decline in kidney function in stable post-MI patients.

References

- 1. WHO: Global Health Risks: Mortality and burden of disease attributable
- to selected major risks. WHO Press: Chapter 2: results, 2009 2. Grubbs V et al: Body mass index and early kidney function decline in
  - young adults: a longitudinal analysis of the CARDIA (Coronary Artery Risk Development in Young Adults) study. Am J Kidney Dis 63: 590-597, 2014 Geleijnse JM et al: Alpha Omega Trial G: Effect of low doses of n-3 fatty acids on cardiovascular diseases in 4,837 post-myocardial infarction patients: design and baseline characteristics of the Alpha Omega Trial.
- Hoogeveen EK et al: Effect of omega-3 fatty acids on kidney function after myocardial infarction: the Alpha Omega Trial. Clin J Am Soc Nephrol 9: 1676-1683, 2014

Table 1: baseline characteristics.

- Labre 11 Saconino characteriotico.	
	n = 2,410
Age, years	68.9±5.4
Men, no. (%)	1914 (79.4)
Smoking, no. (%)	384 (15.9)
BMI, $kg/m^2$	
Men	27.5±3.3
Women	$28.4 \pm 4.6$
Waist circumference, cm	
Men	$102.5 \pm 9.1$
Women	96.4±11.6
Diabetes, no. (%)	444 (18.4)
Anti-hypertensive drugs, no. (%)	2097 (87.0)
Glucose lowering drugs, no. (%)	315 (13.1)
Lipid modifying drugs, no. (%)	2076 (86.1)
Anti-thrombotic agents, no. (%)	2353 (97.6)
Baseline eGFR <sub>cysC</sub> , ml/min/1.73m <sup>2</sup>	81.5 (19.6)

Displayed as Mean ± SD or percentage

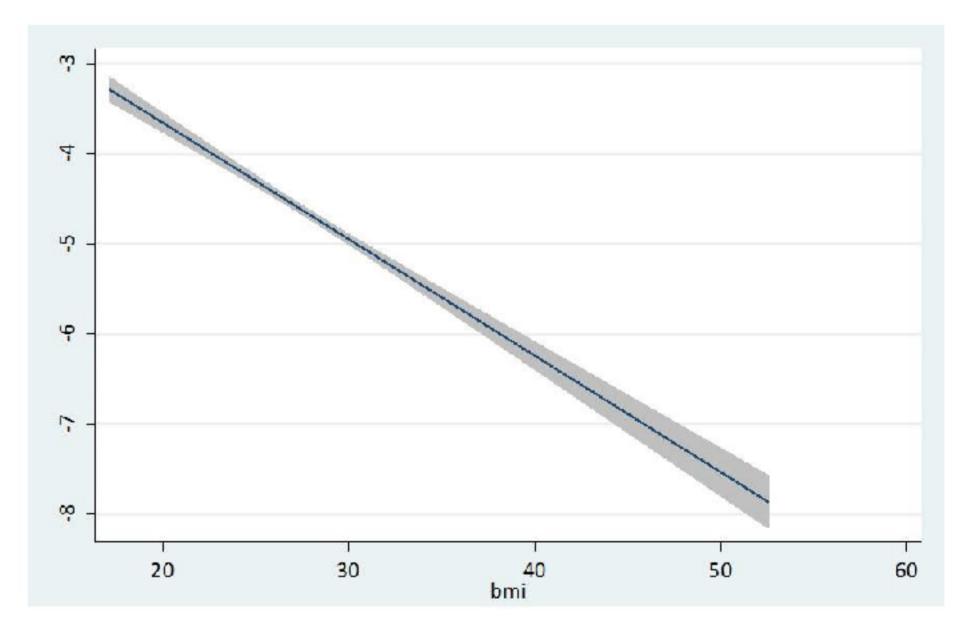


Figure 1: linear relation with 95% confidence interval between BMI (X-axis) and eGFR<sub>cvsC</sub> decline in 40 months (Y-axis).



Corresponding author: k.esmeijer@umail.leidenuniv.nl



Leven gaat voor

Am Heart J 159: 539-546 e532, 2010











