

Estimated Healthcare Resource Utilisation and Costs Associated With Semaglutide 2.4mg Life Expectancy Gains in a Real-World Population With CVD and Overweight or Obesity in the United Kingdom

Hageman S.H.J.¹, Westerink J²,da Rocha Fernandes J.D.³,Arnaut T³, Holloway S⁴, Lee P⁴, Pearson-Stuttard J^{4,5}

Presented by: Silvia Capucci³

Correspondence to: jonathan.pearson-stuttard@lcp.uk.com



Aim

- To estimate the impact of semaglutide 2.4mg treatment on healthcare resource utilisation (HCRU) and costs attributable to life expectancy and ASCVD event-free life-year gains in a UK real-world population with established ASCVD.

Introduction

- In the SELECT cardiovascular (CV) outcomes trial (ClinicalTrials.gov NCT03574597), semaglutide 2.4mg significantly reduced the incidence of major adverse cardiovascular events (MACE: non-fatal myocardial infarction, non-fatal stroke, CV mortality) versus placebo in patients aged ≥45 years with established ASCVD and overweight or obesity, without type 2 diabetes (T2D).¹
- While the cost-effectiveness and life expectancy benefits of semaglutide 2.4mg have been previously demonstrated,^{2,3} the potential HCRU reductions and cost savings associated with life expectancy gains have yet to be quantified.

Methods

- This observational cohort study used data from Discover, a real-world database of linked primary and secondary electronic health records covering 2.8 million people residing in North-West London.⁴
- Individuals aged ≥45 years, with a BMI ≥27 kg/m², established CVD (previous MI, stroke, or PAD) and no history of T2D were included.
- First recurrent CV events and associated healthcare resource utilisation (HCRU) and HCRU costs were assessed between January 2015 – December 2019.
- Approach:**
 - Individuals were partitioned into five health states (CV death, non-CV death, non-fatal myocardial infarction (MI), non-fatal stroke, no event) using extended actuarial life tables.
 - Outcomes were estimated in 5-year age bands over a 5-year horizon.
 - Age-specific observed HCRU events and HCRU costs per person per year were applied to time spent in each state.
 - Age-specific effect estimates from SELECT for 3-point major adverse cardiovascular events (MACE: non-fatal MI, non-fatal stroke, CV mortality), non-CV mortality, and composite 3-point MACE (incl. non-CV mortality) endpoints were applied to aggregated observed outcomes.
 - Reductions in HCRU events and HCRU costs were estimated for semaglutide treatment versus no treatment.
- Primary analysis:** focused on individuals aged 55-74 years, to demonstrate the impact of treatment on HCRU events and HCRU costs among age groups with statistically significant reductions in CV events and mortality observed in SELECT.
- Sensitivity analysis: repeated analyses in the full eligible cohort, applying overall SELECT trial effect estimates for CV event and mortality reductions, to quantify treatment impacts on HCRU and HCRU costs.

Results

- In total, 16,310 SELECT-like individuals aged 45 and over were included (**Table 1**).
 - The percentage of individuals in each 5-year age band from 45 to 70 years is nearly equally distributed, ranging from 12.6% to 13.9%. The proportion of individuals then progressively decreases in the older age bands.
 - The majority (63.8%) of the study population were men.
 - Compared with the SELECT trial participants, this cohort has a smaller proportion of men (63.8% vs. 72%), and greater ethnic diversity (White: 60.6% vs. 84%; Asian: 21.8% vs. 8%; Black: 8.1% vs. 4%).¹
- Main findings suggest that the CVD risk reduction observed with semaglutide 2.4mg treatment leads to 20-25% reduction in HCRU events in those aged 55-74
- Patients observed 3,581 per 100,000 patient years fewer HCRU events associated with composite 3-point MACE, with a reduction in HCRU costs of over £1.9M per 100,000 patient years (**Figure 1**).
- In addition, semaglutide 2.4mg treatment led to a reduction in HCRU events and HCRU costs following a CV event across all age bands, with the greatest reduction in HCRU events (22,252 per 100,000 patient years) and HCRU costs (~£7.4M per 100,000 patient years) observed in the 70 – 74 age band (**Figure 2**).
- When including all age groups, findings were broadly consistent with the main findings from (**Figure 3**).

Table 1: Baseline characteristics of the study population

Characteristics	Overall (N=16,310)
Sex – Male, n (%)	10,406 (63.8)
Ethnicity, n(%)	
White	9,892 (60.6)
Asian or Asian British	3,561 (21.8)
Other	1,454 (8.9)
Black or Black British	1,319 (8.1)
Unknown	84 (0.5)
Age group, n (%)	
45-49 years	2,264 (13.9)
50-54 years	2,130 (13.1)
55-59 years	2,055 (12.6)
60-64 years	2,232 (13.7)
65-69 years	2,139 (13.1)
70-74 years	1,861 (11.4)
75+ years	3,629 (22.3)
CVD type at baseline, n(%)	
MI only	9,010 (55.2)
Stroke only	6,693 (41.0)
PAD only	567 (3.5)
2+ CVD	40 (0.2)
Number of ORCs, n (%)	
2 or more ORCs	14,071 (86.3)
3 or more ORCs	9,608 (58.9)

CVD, cardiovascular disease; MI, myocardial infarction; ORC, obesity-related complication; PAD, peripheral artery disease; HCRU, healthcare resource utilisation; RCT, randomized clinical trial

Figure 1. 3-point MACE and mortality associated HCRU events and costs with and without semaglutide 2.4mg

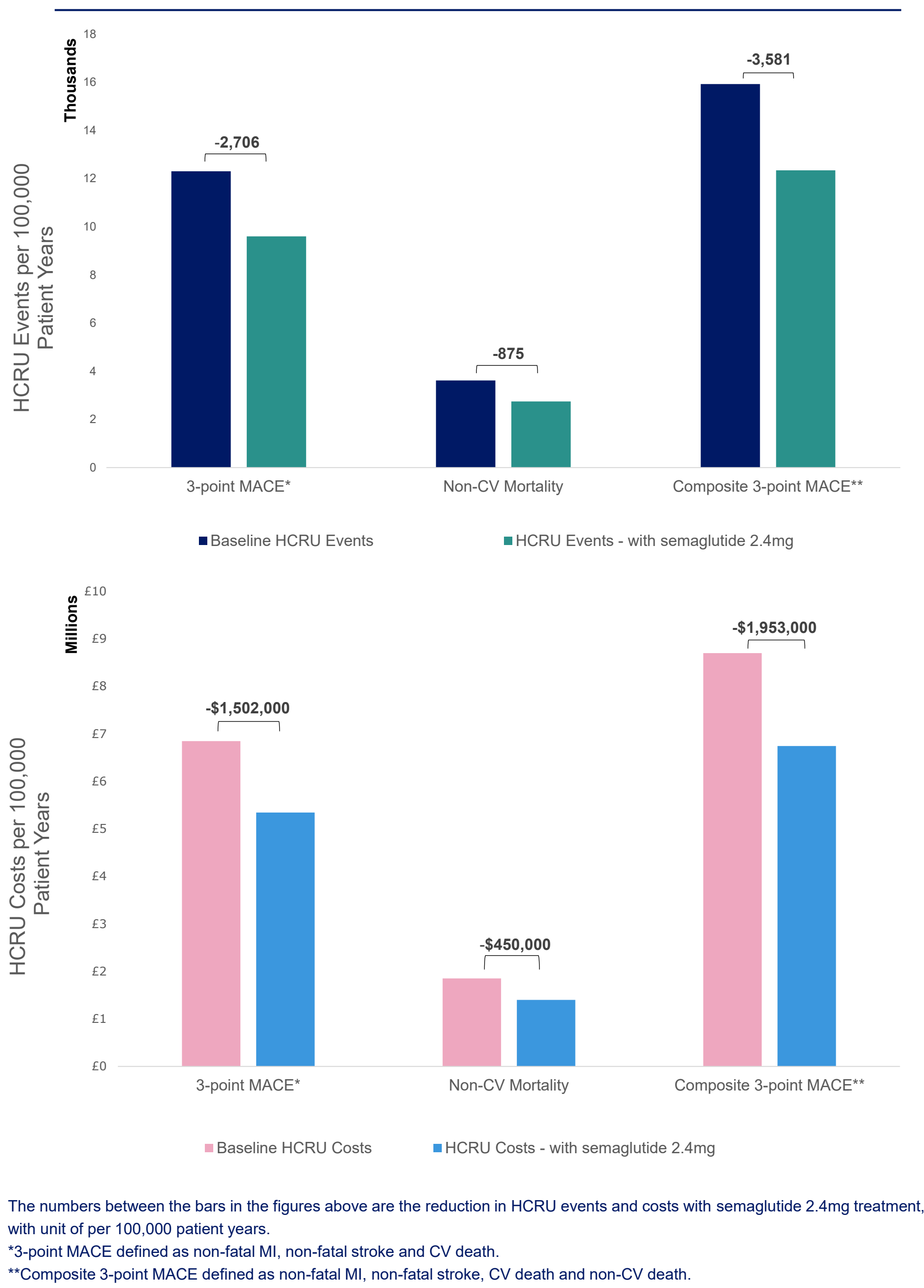


Figure 2. HCRU events and costs following a CV event with and without semaglutide 2.4mg

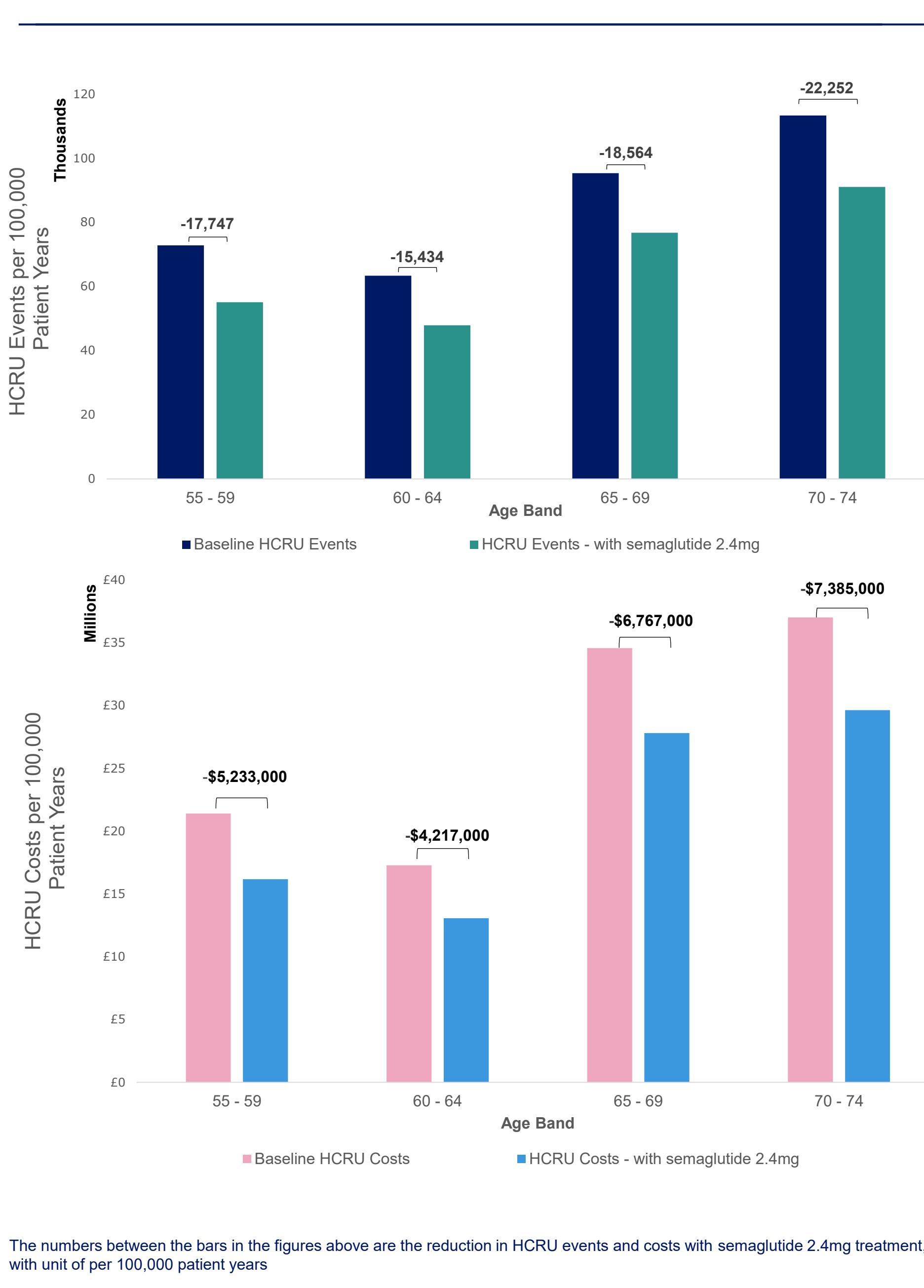


Figure 3. Results from sensitivity analyses including all eligible individuals ≥45 years of age



Strengths and limitations

- Strengths:**
 - This study leverages data from high quality sources such as SELECT and Discover
 - Captures a more diverse, real-world population than typical RCTs.
- Limitations:**
 - Outcomes focus exclusively on HCRU and cost reductions associated with life expectancy and CV event-free life expectancy gains from semaglutide 2.4mg treatment
 - Further research is needed to assess whether these modelling findings are observed in routine clinical practice.

Disclosures:

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Conclusion

- Use of semaglutide 2.4mg treatment was associated with reductions in HCRU events and healthcare costs in people living with overweight or obesity and ASCVD in a North-West London real-world population.
- These cost savings highlight the healthcare system benefits of addressing ASCVD in people living with overweight or obesity and established ASCVD.

References:

¹Department of Vascular Medicine, University Medical Center Utrecht, Utrecht, Netherlands; ²Department of Internal Medicine, Isala, Zwolle, Netherlands; ³Novo Nordisk A/S, Søborg, Denmark; ⁴Health Analytics, Lane Clark & Peacock LLP, London, UK; ⁵Department of Epidemiology and Biostatistics, School of Public Health, Imperial College London, London, UK

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