Impact and management of comorbidities in people with obesity: a multinational survey

Swarna Khare¹, Josefine Redig², Arash Tahbaz¹, Victoria Higgins³, Andrea Leith³, Tamara Mensah³, Esther Artime⁴

¹Eli Lilly and Company, Bracknell, UK, ²Eli Lilly and Company, Stockholm, Sweden, ³Adelphi Real World, Bollington, UK, ⁴Eli Lilly and Company, Alcobendas, Spain



OBJECTIVE

■ To describe the presence of comorbidities in people/patients with obesity (PwO), and associated concomitant treatments, weight management and hospitalisation in PwO with ≥1 comorbidities (PwOC)

CONCLUSION

- Results indicate high presence of obesity-related comorbidities in PwO. Presence of comorbidities was a common physician-reported reason for initiation of weight management and treatment-related discussions
- Preventing the occurrence of, or controlling comorbidities in, PwOC could help reduce overall impact on PwO outcomes, highlighting the need for effective weight management and earlier intervention

International Society for Pharmacoeconomics and Outcomes Research – 26th Annual European Congress; Copenhagen, Denmark; 12-15 November 2023

Analysis funded by Eli Lilly and Company

BACKGROUND

- Obesity is a chronic progressive disease that implicates several organ systems¹ and is associated with various comorbidities, such as type 2 diabetes mellitus (T2DM) and cardiovascular conditions.² Obesity is also linked with mental health issues, such as anxiety and depression¹
- There is a need to better understand the role of comorbidities in PwO and obesity management in real-world settings

STUDY DESIGN

- Data were drawn from the Adelphi Real World Obesity Disease Specific Programme (DSP)™, a real-world, cross-sectional survey of physicians and the PwO they manage, conducted in Brazil, Canada, China, Japan, Kingdom of Saudi Arabia (KSA) and United Arab Emirates (UAE) between April and December 2022
 - The DSP methodology has been published and validated previously³⁻⁵
- Physicians responsible for managing PwO were recruited with specialties varying by country:
 - Brazil: primary care physicians (PCP), diabetologist/endocrinologists and cardiologists; Canada: PCPs and diabetologist/endocrinologists; China: Internists and diabetologist/endocrinologists; Japan: Internists, diabetologist/endocrinologists and cardiologists, and KSA/UAE: PCPs, diabetologist/endocrinologists and obstetricians/gynaecologists
- Physicians completed an attitudinal survey reporting reasons for initiating a discussion with PwO about their weight. Physicians also reported demographics, comorbidities, clinical characteristics and management for up to eight of their consecutive PwO meeting the inclusion criteria
- PwO inclusion criteria: at time of data collection must be aged ≥18 years old, not involved in a clinical trial, on a weight management programme and/or have a BMI of ≥30 in Brazil, Canada, KSA and UAE, ≥28 in China and ≥25 Japan and recruited via one of two groups:
 - No anti-obesity medication (AOM) PwO: not on an AOM at time of data collection
 - AOM PwO: on an AOM at time of data collection
- Analysis criteria: qualifying PwO that had ≥1 comorbidities at time of data collection (PwOC)
- All respondents provided informed consent and ethics exemption was obtained from Pearl IRB. All analyses were descriptive

RESULTS

- A total of 431 physicians provided data on 2839 PwO (Table 1). Overall, mean ± standard deviation (SD) number of comorbidities was 2.1 ± 2.0
- Among these PwO, 83% (n=2351) had ≥1 comorbidity (PwOC).

Table 1: Sample size and AOM status among PwO*

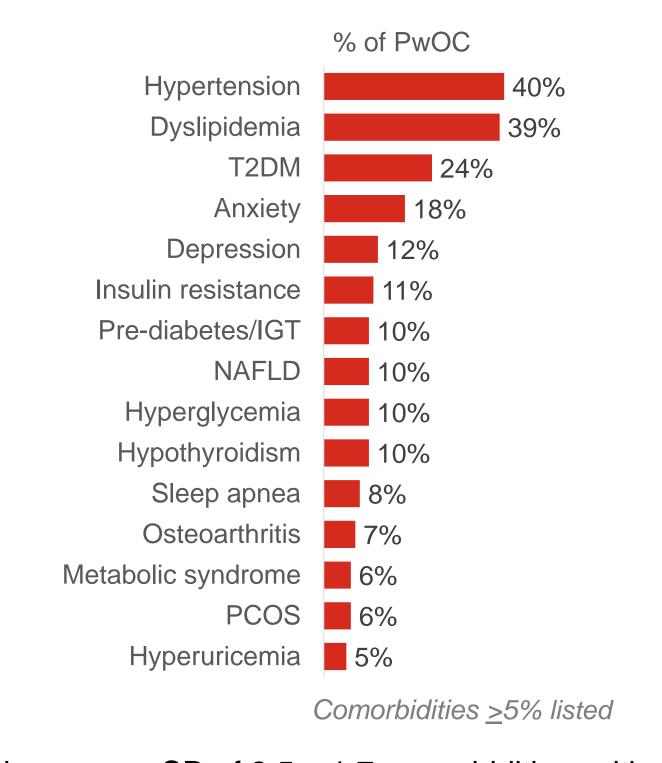
	n	AOM, n (%)
Total	2839	1121 (39)
Brazil	895	480 (54)
Canada	199	156 (78)
China	801	-
Japan	543	285 (52)
KSA	200	100 (50)
UAE	201	100 (50)

*Target quotas by country were established to ensure enough number of PwO in each group

Table 2: Demographics and clinical characteristics among PwOC

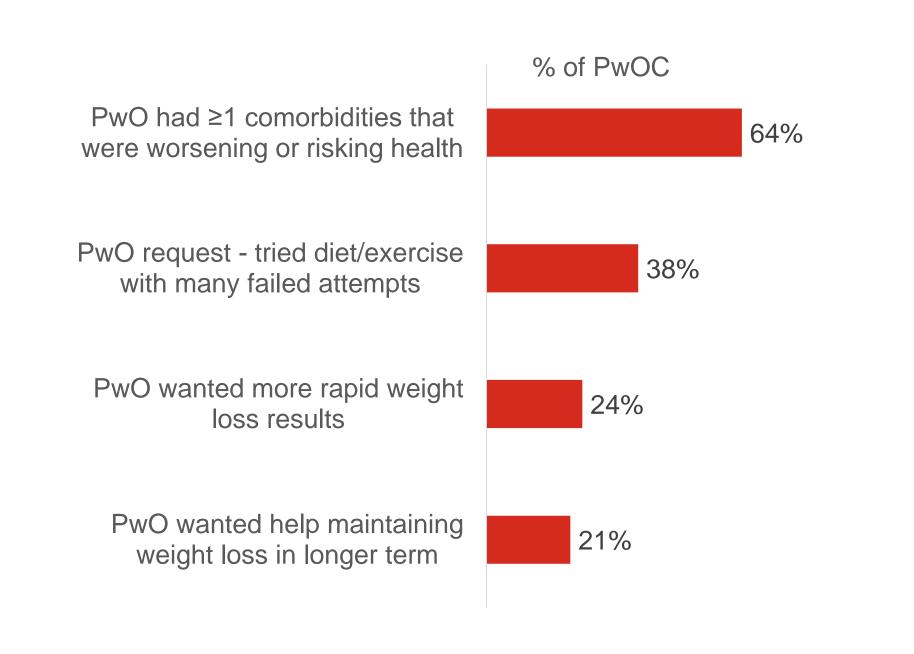
Mean ± SD or n (%)	Total (n=2351)		
Age (years)	44.3 ± 13.5		
Female*	1283 (55)		
BMI (kg/m²) at diagnosis	35.2 ± 7.9		
BMI (kg/m²) at time of data collection	33.2 ± 7.2		
Employed (part-/full-time)	1694 (72)		
Receiving AOM at time of data collection	950 (40)		
Concomitant medications	2.3 ± 2.0		
Top three concomitant medications			
Statin	837 (36)		
Metformin	624 (27)		
Angiotensin-II receptor inhibitor	596 (25)		
Hospitalised in the last 12 months in relation to obesity	72 (3)		
Hospitalised in the last 12 months in relation to any obesity-related comorbidities *Three intersex PwO in Brazil	113 (5)		

Figure 1: Most prevalent comorbidities among PwOC were hypertension, dyslipidemia and T2DM



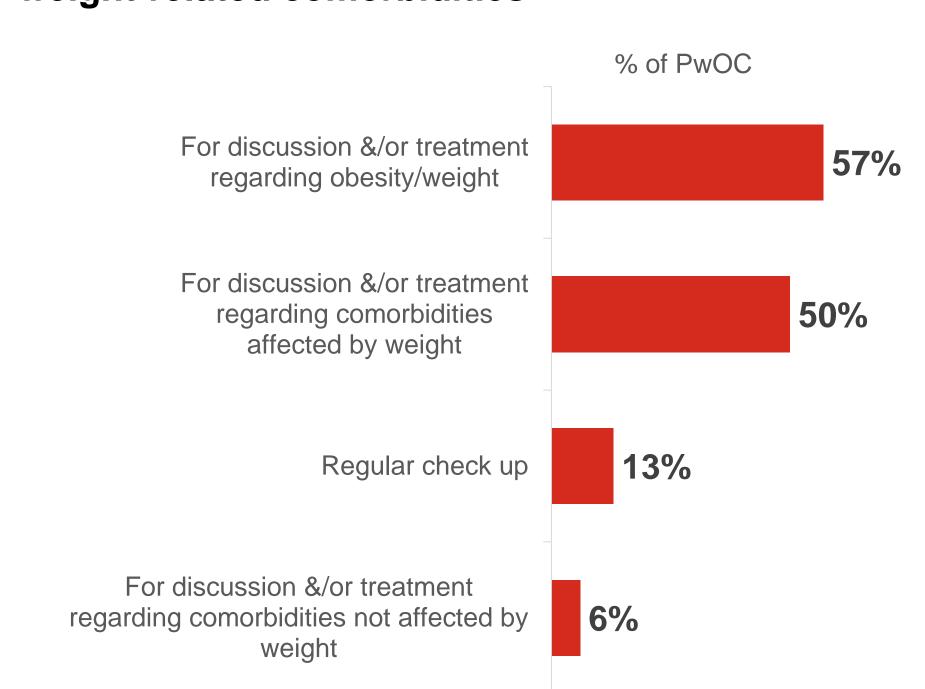
- PwOC had a mean ± SD of 2.5 ± 1.7 comorbidities, with 40% having three or more comorbidities
- Cardiovascular conditions were most prevalent, followed by T2DM and psychological conditions

Figure 3: Most common reason why PwOC started their current weight loss attempt was due to the presence of comorbidities that were worsening or risking their health



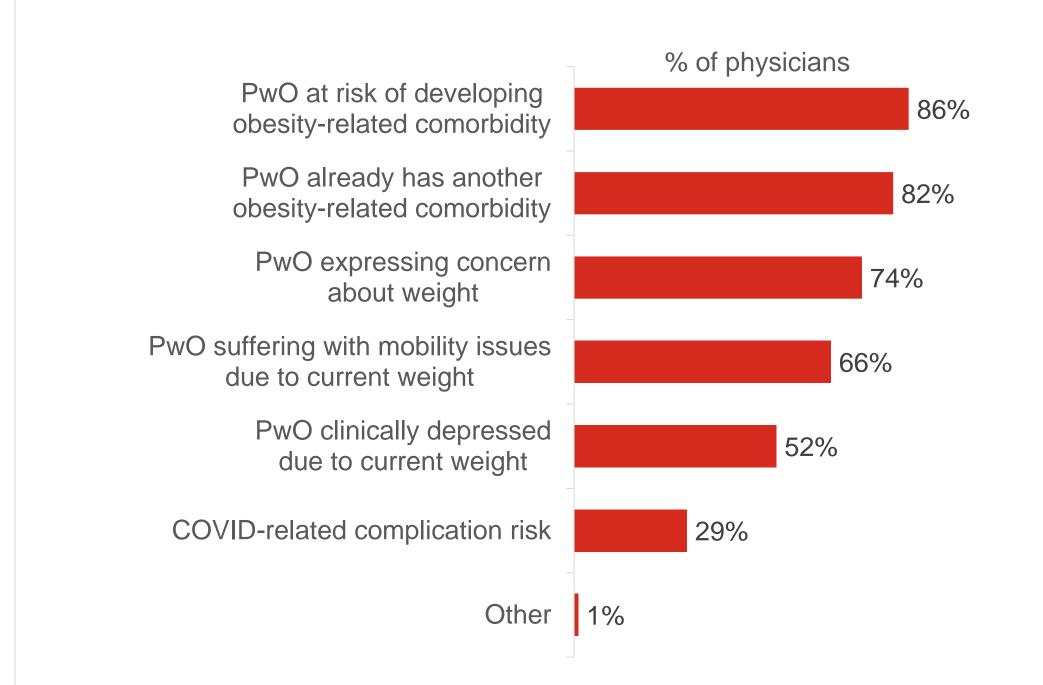
The most common current comorbidities physicians were trying to manage in PwOC were hypertension (47%) and dyslipidemia (39%)

Figure 2: Most PwOC first consulted their physician for discussion/treatment regarding their obesity/weight or for weight-related comorbidities



Among PwOC who consulted for discussion &/or treatment regarding comorbidities affected by weight, the most common comorbidities that triggered these discussions were hypertension (47%), T2DM (41%) and high LDL levels (27%)

Figure 4: Most common reasons why physicians initiated a weight discussion with PwO were the risk of or presence of obesity-related comorbidities



Other important reasons were PwO own concerns about their weight

LIMITATIONS

- Physician participation could have been influenced by willingness to complete the survey
- Recall bias, a common limitation of surveys, may have affected responses to the questionnaires. However, physicians had access to patient records during the survey, minimising recall bias
- The cross-sectional design of this study prevents any conclusions about causal relationships, however identification of significant associations is possible
- No p values were calculated. Any differences noted in reported results are based on summaries of descriptive data, rather than reflecting formal hypothesis testing

REFERENCES:

- 1. Uranga RM et al. (2019) Frontiers in Neuroscience. 24;13
- 2. Khaodhiar L et al. (1999) Clinical Cornerstone. 2(3):17–31 3. Anderson P et al. (2008) Curr Med Res Opin. 24(11):3063-72 4. Babineaux SM et al. (2016) BMJ Open. 6:e010352
- 5. Higgins V et al. (2016) Diabetes Metab Syndr Obes 9:371–380

ABBREVIATIONS:

AOM, Anti-obesity medication; BMI, Body mass index; DSP, Disease specific programme; IGT, impaired glucose tolerance; KSA, Kingdom of Saudi Arabia; LDL, Low density lipoprotein; NAFLD, Non-alcoholic fatty liver disease; NASH, Non-alcoholic steatohepatitis; PCOS, Polycystic ovary syndrome; **PCP**, Primary care physician; **PwO**, People with obesity; **PwOC**, People with obesity with ≥1 comorbidity; **SD**, Standard deviation; **T2DM**, Type 2 diabetes mellitus; **UAE**, United **Arab Emirates**

DISCLOSURES:

VH, AL and TM are employees of Adelphi Real World. SK, JR, AT and EA are employees and shareholders of Eli Lilly and Company. The DSP is a wholly owned Adelphi product. Eli Lilly and Company was one of multiple subscribers to the DSP