Polysomnography (PSG) is the gold standard method for diagnosing OSA. Level 1 PSG is performed in a sleep laboratory with a technician in attendance; it measures respiratory, cardiovascular, and neurologic parameters. However, increasing demand and limited capacity to perform level 1 PSG have resulted in protracted wait times in publicly funded health care systems [18]. Level 3 PSG is performed using portable monitors and may be done in the patient's home. Unlike level 1 PSG, it is not able to detect non-respiratory sleep disorders nor does it typically measure stages of sleep, duration of sleep, or arousals. Despite these limitations, current guidelines recommend that level 3 PSG can be used to diagnose OSA in patients with a high pretest probability of moderate to severe OSA and who do not have signif cant comorbid cardiopulmonary or neurologic conditions [19]. However, a major barrier to accessing level 3 PSG in our provincially-funded health care system (Alberta, Canada) is the need for the patient to bear the cost of these studies.

One potentially useful way to overcome these limitations is to use clinical predictors to identify patients with increased likelihood of OSA and perform testing in only these patients or prioritize these patients for faster objective testing—is approach could potentially streamline use of PSG, decrease wait-times and reduce costs. Screening questionnaires, such as the Epworth Sleepiness Scale or the Berlin questionnaire are not su—cient maccurate to be used as stand-alone tools [20]. Previous studies aimed at deriving clinical prediction rules to inform the need for PSG have shown inconsistent results [21-23]. erefore, no universally accepted and validated tool is currently being used in clinical practice.

e objective of this cross-sectional study was to identify clinical predictors of moderate-to-severe OSA in patients referred to a large Canadian bariatric care program — is program has a regional referral structure and is publicly funded; thus, the patient population is less highly selected than in previous studies [22,23]. We aimed to identify significant predictors of moderate-to-severe OSA and, to compare these with previous studies, and to generate a clinical prediction rule for this condition in a severely obese population. We postulated that this tool may be relevant to streamline referrals for PSG by identifying those that are most likely to have OSA (rather than sending all patients for PSG as is currently being suggested).

Methods

Approval to conduct this cross-sectional study was obtained from the University of Alberta Research Ethics Board (PRO0030092).

Participants and setting

Subjects were recruited from the Edmonton Weight Wise adult bariatric specialty clinic. is bariatric care clinic, established in 2005, serves as a catchment population of approximately 1.6 million residents and includes a central, reg Variable

89 and 91%. However, when this model was evaluated in a di erent sample of patients by Kolotkin et al., the sensitivity and specific itmwas found to be only 75% and 57%, respectively [22]. Instead, Kolotkin proposed a di erent 10-variable model consisting of neck circumference, systolic blood pressure, waist-hip ratio, waist, glucose,

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