GLP-1 Receptor Agonists Decrease Medical Complications, Surgical Complications, and Readmission Rates Following Total Hip Arthroplasty

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INTRODUCTION:

Obesity is known to increase perioperative risk in total hip arthroplasty (THA). Glucagon-like peptide-1 receptor agonists (GLP1-RA) have become popularized due to their effectiveness as weight loss medications. The purpose of this study was to determine the effects of obese patients with osteoarthritis (OA) taking GLP1-RA.

METHODS:

All obese patients with OA undergoing primary THA from 2010 to 2022 were identified using an insurance claims database (n=310,678). Patients taking a GLP1-RA (n=11,883) were matched on a 1:1 basis to patients not taking the medication (n=11,883) using age, gender, body mass index (BMI), Elixhauser Comorbidity Index (ECI), and tobacco use. Outcomes included 90-day medical complications, 90-day readmission rates and 2-year surgical complications. RESULTS:

There were no differences in age, sex, BMI, tobacco use, and ECI between the two groups (p>0.05). Patients on GLP1-RA were found to have lower odds of developing ischemic stroke (0.15% versus 0.37%; OR 0.44; P < 0.05), deep vein thrombosis (0.33% versus 1%; OR 0.35; P < 0.05), pulmonary embolism (0.14% versus 0.50%; OR 0.3; P < 0.05), myocardial infarction (0.10% versus 0.29%; OR 0.4%; P < 0.05), pneumonia (0.45 versus 1.28%; OR 0.39; P < 0.05), acute kidney injury (0.81% versus 1.68%; OR 0.68; P < 0.05), and sepsis (0.14% versus 0.50; OR 0.36; P < 0.05). The odds of revision surgery was lower for GLP1 RA patients (3.03% versus 3.71%; OR 0.87; P < 0.05). When examining indications for revision surgery, patients taking a GLP1-RA had lower odds of prosthetic joint infection (0.30% vs. 0.76%, OR=0.42, p<0.05), periprosthetic fracture (0.12% vs. 0.30%, OR=0.54, p<0.05), and aseptic loosening (0.13% vs. 0.34%, OR=0.33, p<0.05).

DISCUSSION AND CONCLUSION:

Obese patients with OA taking GLP1-RAs had lower odds of 90-day medical complications, 90-day readmissions, and 2-year reoperations following THA when compared to a matched cohort not taking the medication.