GLP1 Receptor Agonists Decrease Medical Complications, Surgical Complications, and Readmission Rates Following Total Knee Arthroplasty

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INTRODUCTION:

Obesity is associated with increased risk following total knee arthroplasty (TKA). Glucagon-like peptide-1 receptor agonists (GLP1-RA) have emerged as a promising therapy for obesity. The purpose of this study was to determine whether obese patients taking a GLP1-RA had different outcomes to patients not on the medication following TKA. METHODS:

All obese patients with OA undergoing primary TKA from 2010 to 2022 were identified using an insurance claims database (n=749,864). Patients taking a GLP1-RA (n=34,048) were matched on a 1:1 basis to patients not taking the medication (n=34,048) using age, gender, body mass index (BMI), Elixhauser Comorbidity Index (ECI), and tobacco use. All patients had a minimum of 2-year follow-up. The outcomes were 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 had lower odds of developing ischemic stroke (0.27% versus 0.62%; OR 0.58; P < 0.05), deep vein thrombosis (0.65% versus 1.58%; OR 0.47; P < 0.05), pulmonary embolism (0.29% versus 0.75%; OR 0.44; P < 0.05), myocardial infarction (0.14% versus 0.38%; OR 0.49; P < 0.05), pneumonia (0.74 versus 1.66%; OR 0.48; P < 0.05), acute kidney injury (1.08% versus 1.84%; OR 0.74; P < 0.05), and sepsis (0.29% versus 0.56; OR 0.67; P < 0.05). The odds of revision surgery was lower for patients on a GLP1-RA (3.11% versus 3.72%; OR 0.88; P < 0.05). Patients taking a GLP1-RA also had lower odds of prosthetic joint infection (0.33% vs. 0.98%, OR=0.39, p<0.05), periprosthetic fracture (0.05% vs. 0.09%, OR=0.44, p<0.05), and aseptic loosening (0.20% vs. 0.48%, OR=0.40, p<0.05).

DISCUSSION AND CONCLUSION:

Obese patients on GLP1-RA had lower odds of 90-day medical complications, 90-day readmissions, and 2-year reoperations following TKA compared to matched patients not taking the medication.