Utilization of glucagon-like peptide-1 receptor agonist at the time of total hip arthroplasty for patients with morbid obesity

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INTRODUCTION:

Morbid obesity negatively affects outcomes after total hip arthroplasty (THA). The optimal strategy for weight loss prior to THA has not been identified. Recently, glucagon-like peptide-1 receptor agonists (GLP-1 RA) have been used for their ability to promote pharmacologic weight loss in the medical management of obesity. The goal of this study was to evaluate the effect of perioperative use in GLP-1 RA in patients with morbid obesity undergoing primary THA on postoperative outcomes.

METHODS:

Using an administrative claims database, patients with morbid obesity (BMI \geq 40.0 kg/m²) undergoing primary THA were identified. Patients with morbid obesity and GLP-1 RA use for 3 months before and after surgery were matched to patients with morbid obesity without GLP-1 RA use (controls) and to comparison group of patients with severe obesity (35.0-39.9 kg/m²) in a 1:4:4 ratio based on patient age, gender, diagnosis of type II diabetes mellitus (TIIDM), and Charlson Comorbidity Index (CCI). Univariable tests were performed to compare overall group differences in 90-day and 2-year postoperative outcomes between matched cohorts, followed by post hoc pairwise testing and p-value adjustment for multiple comparisons.

RESULTS:

Patients with morbid obesity on GLP-1 RA had a significantly lower rate of 90-day periprosthetic joint infection (PJI) (1.6% vs. 3.2%; P=0.034), readmission (6.9% vs 9.7%; P=0.043), any medical complication (10.5% vs 14.1%; P=0.028), and postoperative hematoma formation (0.0% vs. 1.3%, P=0.001) compared to controls. Patients with morbid obesity on GLP-1 RA demonstrated lower rates of hematoma formation (0.0% vs 1.0%; P=0.003) compared to patients with severe obesity (BMI=35.0-39.9kg/m²). There were no differences in other medical complications or 2-year surgical complications. DISCUSSION AND CONCLUSION:

Perioperative use of GLP-1 RA in patients with morbid obesity reduced the risk of acute PJI and 90-day hospital readmission. The risk is reduced to a level comparable to obese patients with BMI <40.0 kg/m². These medications may be a viable strategy in the optimization of this challenging patient population.

Table 2. No-Day Pestoperative Outcomes					
56-Day Outcomes	Severe Obesity (BMI=35- 39.9) (n=3,084)	Morbid Obesity (BMI>→80) GLP-1 RA use (x=771)	Morbid Obesity (BMI>=40) GLP-1 RA non-une (n=3,084)	P-value	Posthoc Pairwise comparisons (adjusted P-value)
Readmissions	273 (8.9%)	53 (6.9%)	300 (9.7%)	0.043	Groups 1 vs 2 (P=0.247); Groups 1 vs 3 (P=0.762); Groups 2 vs 3 (P=0.043)
Any Medical Composition	416 (13.5%)	81 (10.5%)	435 (14.1%)	0.632	Groups 1 vs 2 (P=0.079); Groups 1 vs 3 (P=1.099); Groups 2 vs 3 (P=0.028)
Acute Kidney Injury (AKI)	124 (4.0%)	25 (3.2%)	134 (4.3%)	0.375	
Cardiac Arrest	<11	<11	<11	0.829	
Deep Vein Thrombosis (DVT)	26 (0.8%)	41	36 (1.2%)	0.272	
Wound Defineence	25 (1.8%)	18 (2.5%)	10 (2.3%)	0.345	
Hematoma	32 (1.0%)	0 (0.0%)	39 (1.3%)	0.008	Groups 1 vs 2 (P=0.003); Group 1 vs 3 (P=1.000); Groups 2 vs 3 (P=0.001)
Nerve Injury	0 (0.0%)	0 (0.0%)	0 (0.0%)		
Preumonia (PNA)	52 (1.7%)	<11	50 (1.6%)	0.286	
Painonary Embotism (PE)	×11	511	811	1.000	
Transfesion	65 (2.1%)	<11	65 (2.1%)	0.078	
Urinary Tract Infection (UTD	139 (4.5%)	28 (3.6%)	152 (4.9%)	0.291	
90-Day Surgical Complications	Severe Obesity (BMI=35- 39,9) (n=3,084)	Murbid Obesity (BMI>=40) GLP-1 RA use (n=771)	Morbid Obesity (BMI>=40) GLP-1 RA non-use (n=3,084)	P-value	Postho: Pairwise comparisons (adjusted P-value)
Compenent Revision	55 (1.8%)	<11	59 (1.9%)	0.092	
Prosthetic Joint Infection	68 (2.2%)	12 (1.6%)	100 (3.2%)	0.006	Groups 1 vs 2 (P=0.967); Group 1 vs 3 (P=0.046); Groups 2 vs 3 (P=0.034)
Periprosthetic Fracture	18 (0.6%)	<11	28 (9.1%)	0.315	
Dislocation	55 (1.8%)	12 (1.6%)	53 (1.7%)	0.909	
Aseptic Leonening	<11	0	<11	0.062	
Instability	<11	0	<11	0.457	
Ostrolysb	N11	0	511	0.883	
Wear	0 (0.0%)	0	<11	0.535	