

## VIDEO ROUNDTABLE

### ABSTRACT

# Safe and Appropriate Use of GLP-1 RAs in Treating Adult Patients With Type 2 Diabetes and Macrovascular Disease

**James LaSalle, DO; Lucia M. Novak, CRNP; Lawrence Blonde, MD, MACE, FACP**

10.12788/jfp.0442

**M**acrovascular complications, particularly cardiovascular disease (CVD), are the greatest contributors to the morbidity, mortality, and cost of diabetes mellitus. Atherosclerotic cardiovascular

disease (ASCVD) is the most important macrovascular complication, and type 2 diabetes (T2D) and its associated hyperglycemia are major risk factors for ASCVD. Some antihyperglycemic therapies for T2D, including some sodium-glucose cotransporter-2 inhibitors, thiazolidinediones, and glucagon-like peptide-1 receptor agonists (GLP-1 RAs) have demonstrated benefit in CVD and chronic kidney disease (CKD).

GLP-1, a potent incretin hormone, enhances insulin and inhibits glucagon secretion in a glucose-dependent manner; it also inhibits gastric emptying and gastric acid secretion and increases satiety. GLP-1 RAs mimic endogenous GLP-1 but have a longer half-life. In addition to lowering glucose levels, some GLP-1 RAs have demonstrated the ability to reduce the risk of CVD events. GLP-1 RAs have also been shown to decrease the progression of CKD and to help manage obesity and nonalcoholic steatohepatitis. Adverse events related to GLP-1 RAs include gastrointestinal issues such as nausea, vomiting, diarrhea, and abdominal pain and discomfort.

The roundtable videos in this supplement, developed for primary care clinicians, aim to review the mechanisms of action, clinical trial data, and real-world evidence for the use of GLP-1 RAs in the safe and effective care of individuals with T2D and macrovascular disease. ●

**James LaSalle, DO<sup>1</sup>; Lucia M. Novak, CRNP<sup>2</sup>; Lawrence Blonde, MD, MACE, FACP<sup>3</sup>**

<sup>1</sup>Family Medicine Specialist, The Excelsior Springs Clinic, Excelsior Springs, MO

<sup>2</sup>Nurse Practitioner, Diabetes Consulting Services, North Bethesda, MD

<sup>3</sup>Director, Ochsner Diabetes Clinical Research Unit, Frank Riddick Diabetes Institute, Department of Endocrinology, Ochsner Health, New Orleans, LA

### DISCLOSURES

**James LaSalle, DO**, has disclosed the following relevant financial relationships: Consultant and member of the speakers bureau for Novo Nordisk Inc.

**Lucia M. Novak, CRNP**, has disclosed the following relevant financial relationships: Consultant, speaker, and/or advisor for Abbott Diabetes Care, Novo Nordisk Inc., Provention Bio, Xeris

**Lawrence Blonde, MD, MACE, FACP**, has disclosed the following relevant financial relationships: Consultant for Corcept Therapeutics, Gilead, Janssen Pharmaceuticals Inc, Lyndra Therapeutics, Merck, Novo Nordisk Inc., Salix Pharmaceuticals, Sanofi, and member of the speakers bureau for Sanofi

### ACKNOWLEDGEMENT

The authors acknowledge professional medical writing support from Amy Ross, PhD, PRECISIONscientia, Yardley, PA, which was supported financially by Novo Nordisk Inc., Plainsboro, NJ.

### FINANCIAL SUPPORT

This digital publication was funded by Novo Nordisk Inc., Plainsboro, NJ. Authors received no remuneration for their development of this digital publication.

### VIDEO

The video roundtable associated with this abstract can be found online at <https://www.mdedge.com/JFP/macrovascular-disease>

The video roundtable was peer reviewed by *The Journal of Family Practice*.