

explain the weight gain observed with basal insulin therapy?

Before blindly raising the basal insulin dose based on morning fasting glucose, perhaps we should evaluate the effect of late-night eating. Also, with once-nightly basal insulin, we should test plasma glucose during the expected physiologic glucose nadir between 2 and 6 AM (3).

DISCLOSURE

The author has been a speaker, consultant, and received research grants from Sanofi-Aventis, Eli Lilly, and NovoNordisk.

Allen B. King, MD
1260 S. Main Street
Salinas, CA 93901
E-mail: aking@diabetescarecenter.com

REFERENCES

1. **Strange P.** Treat-to-target insulin titration algorithms when initiating long or intermediate acting insulin in type 2 diabetes. *J Diabetes Sci Technol.* 2007;1:540-548.
2. **King AB, Clark D, Wolfe GS.** The number of basal rates required to achieve near-normal basal glucose control in pump-treated type 2 diabetes. *Diab Technol Therap.* 2012;14:900-903.
3. **Riddle MC, Rosenstock J, Gerich J, Insulin Glargine 4002 Study Investigators.** The treat-to-target trial: randomized addition of glargine or human NPH insulin to oral therapy of type 2 diabetic patients. *Diabetes Care.* 2003;26:3080-3086.

AACE COMPREHENSIVE DIABETES MANAGEMENT ALGORITHM 2013 ENDOCRINE PRACTICE

To the Editor:

The new Comprehensive Diabetes Management Algorithm 2013, which was recently released by the American Association of Clinical Endocrinologists (AACE), modifies previous recommendations made in the 2012 clinical guidelines published by the American Diabetes Association and the European Association for the Study of Diabetes (ADA/EASD). What makes the 2013 Comprehensive Diabetes Management Algorithm innovative is that it visually facilitates comprehension of management and emphasizes the need for individualized therapy, including specific guidelines for the management of obesity, prediabetes, and comorbidities. However, regarding the oral antidiabetic drug schemes, we believe that it is important to point out the increased risk reported on March 14 of this year by the U.S. Food and Drug Administration

(FDA) in relation to pancreatitis and pancreatic neoplasias associated with incretin mimetic therapy.

Although it could well be argued that this information is recent, there are previous publications by the same group of researchers that link this risk with this class of drugs (1). In addition, this association has been described for other drugs, such as thiazolidinediones (2), with metformin use acting as a protective factor (3).

Because of the aforementioned points, we feel that it is necessary to add a column to the Table of the profile of antidiabetic medications, which describes the adverse effects, to mention the relationship between pancreatic tumors and the presence of pancreatitis in an effort to enhance the safety and effectiveness of these medical treatment options.

Finally, some of the drugs reported in the algorithm that represent an alternative to dual and triple therapy that has been recommended in recent years are not available in many countries including our own, such as colesevelam, quick-release bromocriptine, pramlintide, and sodium glucose transporter-2 inhibitors.

DISCLOSURE

The authors have no multiplicity of interest to disclose.

ACKNOWLEDGMENT

The author would like to thank Dr. Sergio Lozano.

Héctor Eloy Tamez-Pérez, MD, Dr. Med¹
Calle Dr. Eduardo Aguirre Pequeño s/n
Colonia Mitras Centro
Monterrey, Nuevo León, 64460, Mexico
E-mail: hectoreloytp@gmail.com

Stephanie Lissette Proskauer-Peña, MD¹
Calle Dr. Eduardo Aguirre Pequeño s/n
Colonia Mitras Centro
Monterrey, Nuevo León, 64460, Mexico
E-mail: mdsproskauer@gmail.com

Mayra Ivonne Hernández-Coria, MD²
Calle Dr. Eduardo Aguirre Pequeño s/n
Colonia Mitras Centro
Monterrey, Nuevo León, 64460, Mexico
E-mail: mayrahc@hotmail.com

¹Subdirección de Investigación

²Servicios Médicos Universidad Autónoma de Nuevo León
Facultad de Medicina y Hospital Universitario "Dr. José Eleuterio González," Universidad Autónoma de Nuevo León, Mexico

REFERENCES

1. **Butler AE, Campbell- Thompson M, Gurlo T, Dawson DW, Atkinson M, Butler PC.** Marked expansion of exocrine and endocrine pancreas with incretin therapy in humans with increased exocrine pancreas dysplasia and the potential for glucagon-producing neuroendocrine tumors. *Diabetes.* 2013;62:2595-2604.