**Important Safety Notice**

The FDA has required this safety update as part of the TANZEUM® REMS (Risk Evaluation and Mitigation Strategy) to inform healthcare providers about the following serious risks of TANZEUM (albiglutide):

- **Potential Risk of Medullary Thyroid Carcinoma (MTC)**
  - Carcinogenicity of albiglutide could not be assessed in rodents, but other glucagon-like peptide-1 (GLP-1) receptor agonists have caused thyroid C-cell tumors in rodents at clinically relevant exposures. It is unknown whether TANZEUM causes thyroid C-cell tumors, including MTC in humans.
  - Cases of MTC in patients treated with liraglutide, another GLP-1 receptor agonist, have been reported in the postmarketing period. The data in these reports are insufficient to establish or exclude a causal relationship between MTC and GLP-1 receptor agonist use in humans.

- **Acute Pancreatitis**
  - In clinical trials, there were more cases of acute pancreatitis among patients treated with TANZEUM than among patients treated with comparators.

**TANZEUM is not recommended as first-line therapy for patients inadequately controlled on diet and exercise.**

A non-promotional factsheet, reviewed by the FDA, with more detailed safety information about these risks is enclosed.

**Indication:** TANZEUM is a GLP-1 receptor agonist indicated as an adjunct to diet and exercise to improve glycemic control in adults with type 2 diabetes mellitus.
Please visit www.TANZEUMREMS.com for more information.

This letter does not contain the complete safety profile for TANZEUM. Please see the enclosed Prescribing Information and Medication Guide.

**Reporting Adverse Events**
You are encouraged to report negative side effects of prescription drugs to GlaxoSmithKline (the Sponsor) at 1-888-825-5249 and/or the FDA www.fda.gov/medwatch, or call 1-800-FDA-1088.

If you have any questions about the information contained in this letter or the use of TANZEUM, you may contact:

- US Medical Information Department at 1-877-356-8368

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