Compounded Injectables – Trimix Formulas F-9, T-105, SB-4, Sermorelin, Sincalide, Hydroxocobalamin, and NAD – Recall Alert

Date of Notice: March 10, 2022

Brief Description of Recall Alert

Olympia Pharmacy is voluntarily recalling eleven (11) specific lots of Trimix Formulas F-9, T-105, SB-4, Sermorelin, Sincalide, Hydroxocobalamin, and NAD, compounded injectables. These compounded products were found to be "out-of-specification".

Administration of subpotent (below expected strength) hydroxocobalamin in infants, pregnant/ breastfeeding women, and elderly populations are at risk for vitamin B12 deficiency, and there is a reasonable probability they could experience adverse events including muscle weakness, neurological peripheral neuropathic numbness or pain, vision loss, and psychiatric disorders (e.g., depression, memory loss).

Affected Products

Drug Name & Strength	NDC	Lot	Expiration Date
NAD, 500 mg vial	N/A	C41008	03/08/2022
NAD, 500 mg vial	N/A	D24005	04/05/2022
Sincalide, 5 mcg vial	N/A	D24001	04/01/2022
Trimix Formula F9, 10 mL vial	N/A	D41C19	04/19/2022
Sermorelin Acetate, 9 mg vial	N/A	D44026	04/26/2022
Sermorelin Acetate, 9 mg vial	N/A	F42104	06/04/2022
Trimix T-105, 5 mL vial	N/A	E41F10	05/10/2022
Trimix T-105, 10 mL vial	N/A	E41G10	05/10/2022
Trimix SB-4, 5 mL vial	N/A	E41C18	05/18/2022
Trimix SB-4, 10 mL vial	N/A	E41D18	05/18/2022
Hydroxocobalamin, 1 mg/mL, 30 mL vial	N/A	E47025	05/21/2022

Prescriber Information

Olympia Pharmacy is notifying its customers by mail and is arranging for return and replacement of all recalled compounded products. Patients and health clinics that have any of the listed compounded products which are being recalled should stop using them and return the recalled product(s) to Olympia Pharmacy.

For questions regarding this recall, contact the recall coordinator by phone at 407-250-4000 or e-mail clientservices@olympiapharmacy.com Monday through Friday from 9 am to 6 pm Eastern Standard Time.

Adverse reactions or quality problems experienced with the use of this product may be reported to the FDA's MedWatch Adverse Event Reporting program either online, by regular mail or by fax.

- Complete and submit the report Online
- Regular Mail or Fax: Download form or call 1- 800-332-1088 to request a reporting form, then complete and return to the address on the pre-addressed form, or submit by fax to 1-800-FDA-0178

Member Information

Members should contact their doctor or healthcare provider if they have experienced any problems that may be related to taking or using this drug product.

Members with medical-related questions, who wish to report an adverse event or quality issues about the products being recalled, should contact Olympia Pharmacy by phone at 407-250-4000 or e-mail clientservices@olympiapharmacy.com Monday through Friday from 9 am to 6 pm Eastern Standard Time.

RxAdvance Response

Members should continue taking their drug until a doctor or pharmacist provides replacement drug (if needed) or a different treatment option is given. RxAdvance is in the process of contacting members and prescribers to advise them of this recall.