Mylan – Recall of alprazolam

- On October 23, 2019, Mylan announced a consumer-level recall of one batch of alprazolam 0.5 mg tablets due to the remote potential for foreign material on the tablets.
- The recalled lot was distributed between July 2019 and August 2019.

Product Description	NDC#	Lot #
Alprazolam 0.5 mg tablets, 500 count bottle	0378-4003-05	8082708

- Alprazolam tablets are indicated for the management of anxiety disorder, the short-term relief of symptoms of anxiety, and for the treatment of panic disorder, with or without agoraphobia.
- Clinical impact from the foreign material, if present, is expected to be rare, but the remote risk of infection to a patient cannot be ruled out.
- Mylan has not received any reports of adverse events, to date, for the recalled lot.
- Patients should contact their physician or healthcare provider if they have experienced any problems that may be related to using the recalled alprazolam.
- Anyone with an existing inventory of the recalled product should quarantine and discontinue distribution of the product immediately.
- Contact Stericycle (appointed company for Mylan) at **1-888-843-0255** for further information regarding this recall.