Shire – Recall of Kalbitor® (ecallantide)

- On April 12, 2017, the FDA announced a Class II recall of one lot of Kalbitor (ecallantide) injection due to the presence of glass particulate matter.

Product Description	NDC #	Lot # (Expiration date)
Kalbitor (ecallantide) 10 mg/mL single-use vials, packaged as 3 vials per carton	47783-101-01	A1500009 (1/31/2019)

- Kalbitor is indicated for the treatment of acute attacks of hereditary angioedema in patients ≥ 12 years of age.
- Healthcare providers, distributors, wholesalers, and patients should immediately check inventory, quarantine, discontinue distribution, and return all recalled product.
- For questions regarding this recall, contact Shire at **1-800-828-2088**.