BD – Recall of BD insulin syringes

- BD announced a consumer-level recall of one lot of BD Ultra-Fine™ needles ½ mL 12.7 mm 30G, catalog # 328466, because some polybags in the lot are incorrectly labeled as BD Ultra-Fine needle ½ mL 8 mm 31G, catalog # 328468.
    - The shelf carton and case carton are correctly labeled as BD Insulin Syringes with BD Ultra-Fine needle ½ mL 12.7 mm 30G.
    - Additional information to identify the recalled product can be found here.
- The recalled lot was distributed from March 3, 2017 to present.

Product Description	NDC #	Catalog #	Lot #
BD Insulin Syringes with the BD Ultra-Fine™ needle ½ mL 12.7 mm 30G	08290-8466-01; 08290-3284-66	328466	6291768
BD Insulin Syringes with the BD Ultra-Fine needle ½ mL 8 mm 31G	08290-8468-01	328468	

- The polybags of BD Ultra-Fine needles that are incorrectly labeled as ½ mL 8 mm 31G, contain syringes that are ½ mL 12.7 mm 30G. This may present a health hazard to patients using the product affected by this recall. Using a 12.7 mm needle for injection when an 8 mm was intended could result in an increased risk of an inadvertent intramuscular injection, which may lead to unanticipated hypoglycemia.
- Anyone with an existing inventory of the recalled product should stop use and distribution, and quarantine the product immediately. Patients should contact their healthcare provider if they have experienced any problems that may be related to using the recalled product.
- BD will assist patients with the return of the recalled product and how to obtain a replacement at no charge.
- For questions regarding this recall, contact BD at **1-888-345-5346**.