Trividia Health – Recall of Insulin Syringes

- On September 7, 2018, Trividia Health announced a voluntary, patient-level recall of several lots of TRUEplus® and store brand insulin syringes because certain lots contain a defect in which a small crack in the top end of the barrel near the needle creates the inability to aspirate insulin into the syringe barrel from the insulin vial. Inability to draw insulin into the syringe deems the syringe unusable.

- There are 26 affected lots that were manufactured for Trividia Health from April 30, 2018 thru August 21, 2018, of which 3 lots were distributed by Trividia Health for sale in the U.S. The other 23 affected lots have been contained and not released in the market.

- The recalled lots are listed below:

Product Description	NDC #	Lot #
TRUEplus 0.3cc, 29G insulin syringe	56151-1711-01	NP18196
Leader 0.3cc, 31G insulin syringe	56151-1731-01	NP18123; NP18130

- 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.

    — To date, Trividia Health has not received any reports of patient injuries related to this voluntary recall.

- Patients should contact their physician or healthcare provider if they have experienced any problems that may be related to using the recalled product.

- For more information regarding this recall contact Trividia Health at **1-800-588-1685**.