## Bayer – Update to Recall of Kogenate® FS antihemophilic factor (recombinant)

- On July 19, 2019, Bayer announced a consumer-level recall of two lots of Kogenate® FS antihemophilic factor (recombinant) 2000 IU vials because some of these vials contained Jivi® antihemophilic factor (recombinant) PEGylated-aucl 3000 IU.

    — Further information about this recall is found in a previous RxNews (from July 19).

- Although an initial claims assessment did not indicate that any members were affected, a second review of the data did identify that some members were impacted. Therefore, Clinical Services will conduct mailings to members that may be affected by the Kogenate FS recall, and to providers who prescribed for Kogenate FS recently.