Sun – Recall of Fyremadel® (ganirelix acetate) injection

- On April 18, 2023, Sun announced a consumer-level recall of one lot of Fyremadel (ganirelix acetate) injection because glass particulate was observed in one syringe.
    - Other Fyremadel injection products are available for patients to use.

Product Description	NDC	Lot number
Fyremadel (ganirelix acetate) injection, 250 mcg/0.5 mL	55566-1010-1	HAD1190A

- Fyremadel injection is indicated for the inhibition of premature luteinizing hormone surges in women undergoing controlled ovarian hyperstimulation.
- To date, Sun has not received any reports of adverse events with this lot number related to this recall.
- Contact Sun by phone at 1-800-406-7984 or by email at **drug.safetyUSA@sunpharma.com** for questions regarding this recall.