Magno-Humphries Laboratories – Recall of Senna Laxative

- On January 22, 2018, the FDA announced a consumer level recall of one lot of Magno-Humphries Laboratories' Basic Drugs Brand of senna laxative 8.6 mg tablets due to a customer complaint that their bottle labeled as senna laxative actually contained Basic Drugs Brand of naproxen sodium 220 mg tablets.

- Basic Drugs Brand senna laxative tablets were distributed nationwide in the U.S. to secondary distributors, retail pharmacies and via the internet.

Product Description	NDC #	Lot # (Expiration Date)
Basic Drugs Brand of senna laxative tablets, 8.6 mg, 100-count bottle	00761-0790-02	352300 (1/2019)

- Senna is used to relieve occasional constipation and naproxen is used as a pain reliever/fever reducer and is a nonsteroidal anti-inflammatory drug (NSAID).

- Unintentional consumption of naproxen sodium potentially could result in fatal adverse events in patients with underlying illnesses, including known allergy to the hidden ingredients, cardiac, gastrointestinal, hepatic, and renal conditions as well as patients who recently undergone cardiac bypass graft surgery.

- Patients may inadvertently overdose by taking another NSAID concurrently, thus increasing the risk for NSAID associated adverse events, which include but are not limited to, myocardial infarction, stroke, congestive heart failure, renal toxicity, bleeding, ulceration, or perforation of the stomach or the intestines. The populations most at risk are children, pregnant women, nursing mothers, and surgical patients.

- To date, Magno-Humphries Laboratories has not received any reports of adverse events related to this recall.

- Magno-Humphries Laboratories has notified its distributor by e-mail and is arranging for the return of all recalled products. Anyone that has the recalled Basic Drugs Brand senna laxative should stop use and return the product to Magno-Humphries Laboratories or their distributor.

- Contact Magno-Humphries Laboratories at **1-503-684-5464** or **1-800-935-6737** for more information regarding this recall.