Bryant Ranch Repack – Recall of spironolactone

- On March 9, 2021, the FDA announced a voluntary, patient-level recall of four lots of Bryant Ranch Repack spironolactone tablets because some prepackaged bottles labeled spironolactone 50 mg may contain spironolactone 25 mg tablets and prepackaged bottles of spironolactone 25 mg may contain spironolactone 50 mg tablets. This recall was originally announced by Bryant Ranch Repack on January 29, 2021.

- As of 3/9/2021 Bryant Ranch Repack has not received any reports of adverse events related to this recall.

- The recalled lots are listed below:

Product Description	NDC#	Lot# (Expiration Date)
Spironolactone 25 mg tablets	63629106401	148969 (7/31/2022)
	63629106402	148791 (7/31/2022)
	63629106403	148991 (7/31/2022)
Spironolactone 50 mg tablets	63629106701	CT003 (5/2022)

- Spironolactone is indicated as a diuretic in the treatment of high blood pressure, heart failure, hypokalemia, and edema.

- A patient who consumes spironolactone 25 mg instead of the prescribed spironolactone 50 mg may experience an elevation in blood pressure or edema if taking the product chronically.

- It is possible that patients could experience a decrease in potassium if taking half of the expected dose which could lead to hypokalemia, a condition associated with cardiac arrhythmias.

- Furthermore, patients who consume spironolactone 50 mg instead of the prescribed spironolactone 25 mg could experience an increase in potassium which could be life-threatening. Patients with renal insufficiency or those taking concomitant renin-angiotensin-aldosterone system inhibitors would be at increased risk.

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

- Anyone with an existing inventory of the recalled product should stop use, distribution and quarantine the product immediately.

- Contact Bryant Ranch Repack at **1-877-885-0882** for more information about this recall.