Drug Recall

DATE OF RECALL: January 05, 2019

DRUG NAME: Ceftriaxone

RECALLING FIRM: Lupin Pharmaceuticals Inc

REASON FOR RECALL: This recall was issued because the products have been found to contain visual grey particulate matter in reconstituted vials.

FDA LINK: https://www.fda.gov/Safety/Recalls/ucm629298.htm

OTHER DETAILS: See FDA link.

RECALLED PRODUCT:

Product Name	NDC	Lot	Expiration
Ceftriaxone for Injection USP, 250mg	68180-611-10 (10 pack carton) 68180-611-01 (single-vial)	C600182 C600136 C600142 C700147 C700207	09/2019 08/2019 08/2019 05/2020 09/2020
Ceftriaxone for Injection USP, 500mg	68180-622-01 (single-vial) 68180-622-10 (10 pack carton)	C600218 C600219 C600126 C600127 C600137 C600143 C600173 C700146 C700208 C700209	09/2019 09/2019 08/2019 08/2019 08/2019 08/2019 08/2019 05/2020 09/2020 09/2020
Ceftriaxone for Injection USP, 1g	68180-633-10 (10 pack carton) 68180-633-01 (single-vial)	C600106 C600108 C600110 C600174 C600179 C600180 C600181 C700110 C700111 C700131 C700132 C700138	05/2019 05/2019 05/2019 09/2019 09/2019 09/2019 09/2019 03/2020 03/2020 05/2020 05/2020 05/2020

Product Name	NDC	Lot	Expiration
		C700143 C600128 C600130 C600138 C700108 C700109 C700112 C700129 C700130 C700142 C700145 C700113	05/2020 08/2019 08/2019 08/2019 03/2020 03/2020 03/2020 05/2020 05/2020 05/2020 05/2020 03/2020
Ceftriaxone for Injection USP, 2g	68180-644-01 (single-vial) 68180-644-10 (10 pack carton)	C600109 C600129 C600135	05/2019 08/2019 08/2019