

Drug Recall

DATE OF RECALL: November 21, 2023

DRUG NAME: Vitrakvi® (larotrectinib) Oral Solution 20 mg/mL

RECALLING FIRM: Bayer Healthcare Pharmaceuticals Inc.

REASON FOR RECALL: This recall was issued due to microbial contamination identified as *Penicillium brevicompactum* observed during routine ongoing stability testing.

FDA LINK: <https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/bayer-issues-voluntary-recall-nationwide-vitrakvir-larotrectinib-oral-solution-20-mgml-due-presence>

OTHER DETAILS: N/A

RECALLED PRODUCT:

Product Description	NDC Number	Lot Number	Expiration Date
Vitrakvi® (larotrectinib) Oral Solution 20 mg/mL	50419-0392-01	2114228	2/29/2024