

## **Metformin ER - Viona**

Viona announced a consumer level recall of twenty-three lots of metformin 750 mg tablets due to the detection of N-nitrosodimethylamine (NDMA) levels above the acceptable daily intake limit in one lot.

Product	Metformin ER
Manufacturer	Viona
Recall identification date	01/07/2022
Affected NDCs	72578-036-01

Consumers should contact their physician or healthcare provider if they have experienced any problems that may be related to taking or using this product.