Clonazepam - Endo

On July 5, 2024. Endo announced a voluntary recall of one lot of clonazepam 0.25 mg orally disintegrating tablets (ODT) because they were packaged in cartons labeled with the wrong strength, clonazepam 0.125 mg ODT.

On July 16, 2024, Endo announced a voluntary recall of one lot of clonazepam 0.125 mg ODT. This is an expansion of the original recall.

Per Endo, inadvertently taking clonazepam 0.25 mg rather than the intended 0.125 mg dose may lead to drowsiness, confusion, diminished reflexes and other adverse effects. To date, Endo has not received any reports of adverse effects related to this recall.

Use the table below to check the NDC number on your prescription. Call your pharmacist if you need help.

Clonazepam Orally Disintegrating Tablets Recalled by Endo

Product Description	NDC#	Lot# (Expiration Date)
Clonazepam orally disintegrating tablets, 0.125 mg	49884-306-02	550147301 (Aug 2026)
Clonazepam orally disintegrating tablets, 0.25 mg	49884-307-02	550147301 (Aug 2026)

If you have additional questions:

- Contact your doctor
- Contact Inmar (appointed company for Endo) at **1-877-890-0765** (9:00 a.m. – 9:00 p.m. EST, Monday through Friday).

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