



Wellpoint New Jersey

P. O. Box 62509

Virginia Beach, VA 24466-2509

<Date>

<Parent or Guardian of>

<Member Name>

<Member Address>

<Member City, State ZIP>

Dear Member or Parent/Guardian of <Member Name>:

We have important drug recall information about your prescription for Tramadol Hydrochloride Tablets 50 mg.

Our records show that you may have recently filled a prescription for this drug.

On March 30, 2026, the United States Food and Drug Administration (FDA) issued a Class II recall of two lot numbers of Tramadol Hydrochloride Tablets 50 mg.

Amneal Pharmaceuticals, LLC issued this recall due to Failed Impurities/Degradation Specifications: N-nitroso-desmethyl-tramadol impurity (NDSRI) was found out of the specification at 24-month stability interval during Long term stability (25°C/60%RH).

<p>Product: Tramadol Hydrochloride Tablets 50 mg NDC: 60219-2348-05 Lot number/Expiration date: AM230987 exp. 05/2026, AR232387 exp. 12/2026</p>

Talk to your doctor before you stop taking any medication. Stopping a drug without a replacement could have health risks. Your doctor can talk to you about other options that are right for you.

What you need to do next:

Although the distributor doesn't need any action on your part, if you have questions or concerns about this recall, contact the distributor.

wellpoint.com/nj/medicaid

1090596NJMESWLP 04/26

OMHC# 078-25-59

105-CNMCD26034WLPNJ 041426



We're here to help

If you have questions about the recall, please call Amneal Pharmaceuticals, LLC, the distributor, at 1-833-582-0812, Monday through Friday, 9:00 am to 5:00 pm EST. You can contact the FDA toll-free at **888-INFO-FDA (888-463-6332)** or visit fda.gov for more information.

You can also call your doctor or pharmacist if you have questions about the drug or about this letter. If you have questions about your prescription drug benefits, please call Pharmacy Member Services at the phone number on your member ID card (TTY users can call **711**).

Sincerely,

Your Wellpoint service team

Enclosures: Language assistance sheet
Nondiscrimination notice

This information is provided for educational purposes only and should not be considered medical advice. Talk to your physician for specific treatment options and recommendations best suited for you and before beginning any lifestyle program.



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P. O. Box 62509

Virginia Beach, VA 24466-2509

<Date>

<Padre o tutor de>

<Member Name>

<Member Address>

<Member City, State ZIP>

Estimado/a miembro o padre/tutor de <Member Name>:

Tenemos información importante del retiro de medicamento sobre su receta para clorhidrato de tramadol en comprimidos de 50 mg.

Nuestros registros muestran que usted tal vez ha surtido recientemente una receta de este medicamento.

El 30 de marzo de 2026, la Administración de Alimentos y Medicamentos (FDA) de los Estados Unidos emitió un retiro Clase II de dos números de lote de clorhidrato de tramadol en comprimidos de 50 mg.

Anneal Pharmaceuticals, LLC emitió este retiro debido al incumplimiento de las especificaciones de impurezas/degradación: se detectó la impureza N-nitroso-desmetiltramadol (NDSRI) fuera de los límites especificados en el intervalo de estabilidad de 24 meses durante estudios de estabilidad a largo plazo (25 °C/60% HR).

<p>Producto: Clorhidrato de tramadol en comprimidos de 50 mg NDC: 60219-2348-05 Número de lote/Fecha de vencimiento: AM230987 vto. 5/2026, AR232387 vto. 12/2026</p>

Hable con su médico antes de dejar de tomar cualquier medicamento. Dejar de tomar un medicamento sin un reemplazo podría tener riesgos para la salud. Su

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