



**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 Furosemide Tablets, USP, 80 mg.

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

On April 21, 2026, the United States Food and Drug Administration (FDA) issued a Class II recall of certain lot numbers of Furosemide Tablets, USP, 80 mg.

Leading Pharma, LLC issued this recall due to: CGMP Deviations; presence of N-nitroso-Furosemide (NNF) above the recommended intake limit.

**Product:** Furosemide Tablets, USP, 80 mg

**NDC:** 69315-0118-01

**Lot number:** H03125, H03225, H03325

**Expiration date:** 08/2027

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

**[wellpoint.com/nj/medicaid](https://wellpoint.com/nj/medicaid)**

1091139NJMESWLP 05/26

OMHC# 078-25-59

105-CNMCD26050WLPNJ 043026



**We're here to help**

If you have questions about the recall, please call Leading Pharma, LLC, the manufacturer, at 1-844-740-7500. You can contact the FDA toll-free at **888-INFO-FDA (888-463-6332)** or visit [fda.gov](http://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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<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 de la retirada del medicamento sobre su receta para comprimidos de furosemida, USP, 80 mg.

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

El 21 de abril de 2026, la Administración de Alimentos y Medicamentos (FDA) de los Estados Unidos emitió un retiro Clase II de determinados números de lote de comprimidos de furosemida, USP, 80 mg.

Leading Pharma, LLC emitió este retiro debido a: Desviaciones de las CGMP; presencia de N-nitroso-furosemida (NNF) por encima del límite de ingesta recomendado.

**Producto:** Comprimidos de furosemida, USP, 80 mg

**NDC:** 69315-0118-01

**Número de lote:** H03125, H03225, H03325

**Fecha de vencimiento:** 08/2027

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 médico puede hablar con usted sobre otras opciones que sean adecuadas para usted.

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