



<<Date>>

<<First Name>> <<Last Name>>

<< Address line 1>>

<< Address line 2>>

<<City>> <<ST>> << Zip Code>>

### **IMPORTANT RECALL INFORMATION**

Dear <<First Name>> <<Last Name>>,

We are writing to update you about Quinapril 20 mg and 40 mg tablets and tell you what you'll need to do next.

On December 21, 2022, the United States Food and Drug Administration (FDA) published a notice regarding a recall of Quinapril 20 mg and 40 mg tablets manufactured by Lupin Pharmaceuticals, Inc. **This recall was issued because of the presence of a nitrosamine impurity greater than the US Food and Drug Administration (FDA) Acceptable Daily Intake (ADI) level. This could be harmful to your health.**

Our records show you may have recently filled a prescription for Quinapril 20 mg and/or 40 mg tablets from your retail pharmacy.

#### **This recall affects:**

**Product:** Quinapril 20 mg Tablet, 90 count  
**NDC:** 68180-0558-09  
**Lot Number/Expiration Date:** G102929 exp. 04/2023

**Product:** Quinapril 40 mg Tablet, 90 count  
**NDC:** 68180-0554-09  
**Lot Number/Expiration Date:** G100533 exp. 12/2022  
G100534 exp. 12/2022  
G203071 exp. 03/2024

To see if your product was recalled, check the lot number. The lot number is on the manufacturer's label on the bottle.

- If your drug has a different lot number, it's not affected by this recall.
- If your drug is from one of the affected lot numbers or filled in a pharmacy bottle, the FDA is advising that patients taking these products are advised to continue taking their medication and contact their medical provider for advice regarding an alternative treatment. **Call your doctor right away if you may be using the affected product or if you're unsure.** Your doctor knows your medical history and can suggest the best treatment option for you.
- For assistance with returning affected product, please call the number on your benefit ID card.

This document contains references to brand-name prescription drugs that are trademarks or registered trademarks of pharmaceutical manufacturers not affiliated with CVS Caremark. Your privacy is important to us. Our employees are trained regarding the appropriate way to handle your private health information. This information is not a substitute for medical advice or treatment. Talk to your doctor or health care provider about this information and any health-related questions you have. CVS Caremark assumes no liability whatsoever for the information provided or for any diagnosis or treatment made as a result of this information.

For more information, call Lupin Medical Information toll-free at 1-800-399-2561. You may also call the U.S. Food and Drug Administration at 1-888-INFO-FDA (1-888-463-6332). Or visit [fda.gov](http://fda.gov).

- This letter is offered in other languages. Call the number on your benefit ID card. TTY users should call 1-800-863-5488.
- Esta información está disponible en otros idiomas. Por favor llame a nuestro departamento de servicios al cliente al número gratuito que aparece en su tarjeta de identificación. Los usuarios de equipo teleescritor (TTY) deben llamar al 1-800-863-5488.

With your safety in mind,

CVS Caremark® Medical Affairs