



**IMPORTANT ANAGRELIDE CAPSULES, USP 0.5 MG
RECALL NOTICE**

Dear Member,

Your health is important to us. On **May 23, 2022** the U.S. Food and Drug Administration (FDA) announced a drug recall notice. It is for **one** lot of Teva's **Anagrelide Capsules, USP 0.5 mg**.¹ The FDA issued the recall because the product may have a safety concern.

Products that may be affected by this recall:

| Product | NDC | Lot # | Exp. Date |
|-----------------------------------|--------------|--------------|------------------|
| Anagrelide capsules, USP 0.5mg | 0172-5241-60 | GD01090 | 05/2022 |

Additional information is available at: https://www.fda.gov/safety/recalls-market-withdrawals-safety-alerts/teva-issues-voluntary-nationwide-recall-one-lot-anagrelide-capsules-usp-05-mg-due-dissolution-test?utm_medium=email&utm_source=govdelivery

Please call your pharmacy to find out if your drug could be part of this recall. The pharmacy may replace the drug. We are writing this letter to give you information. This should not replace your doctor's advice. Only your doctor can decide what drugs are right for you.

Please disregard this letter if you are not currently taking **Anagrelide Capsules, USP 0.5 mg** or if this safety warning does not apply to you. Our records may not be complete, or your doctor may have discontinued this medication.

Sincerely,

Doctors HealthCare Plans, Inc.

ATENCIÓN: Si habla español, tiene a su disposición servicios gratuitos de asistencia lingüística. Llame al 786-460-3427 o 833-342-7463(TTY: 711).

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