VOLUNTARY RECALL NOTICE:

ELIQUIS 5MG TABLETS

Partial Lot Recall

June 20, 2017

Dear Care1st Providers & Office Staff:

Please be aware that Bristol-Myers Squibb voluntarily recalled one lot of Eliquis 5mg tablets due to a customer complaint that one bottle of Eliquis 5mg was found to contain Eliquis 2.5mg tablets.

At this time there have not been any adverse events reported. However, if a patient takes a lower dose of Eliquis for a prolonged period of time, it could result in an increased probability of stroke, blood clot, or death.

The recall impacts Eliquis 5mg tablets, 60-count bottles. The affected product was distributed to wholesalers and retail pharmacies in February 2017.

The following lot is being recalled in the United States:

- Eliquis 5mg, NDC 00003-0894-21, lot HN0063 (Expiration date: September 2019)

Patients with the affected lot should contact their physician and call Bristol-Myers Squibb at 1-800-332-2056.

For additional information regarding the recall, please refer to the FDA Enforcement Report found at:
https://www.fda.gov/Safety/Recalls/ucm563002.htm?source=govdelivery&utm_medium=email&utm_source=govdelivery"

If you have questions please use the information below to contact us.

Thank you!

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