



February 28, 2011

Dear AmeriHealth Mercy Participating Provider,

As you may be aware, Qualitest Pharmaceuticals issued a voluntary recall of Hydrocodone/Acetaminophen 10mg/500mg and Phenobarbital 32.4mg on February 5, 2011 due to a bottle of hydrocodone being inadvertently labeled as phenobarbital.

Any patients who have received the affected product(s) should stop using them and contact Qualitest at 1-800-444-4011. In addition Qualitest is notifying all customers who may have received affected product and arranging for the return of any affected product.

Please note these tablets have the following appearance:

- Hydrocodone Bitartrate and Acetaminophen Tablets are large (approximately 16.5 mm in length), pink, capsule-shaped tablets, debossed (3600) on one side, and debossed (V) on the reverse side
- Phenobarbital Tablets are small (approximately 6.4 mm in diameter), white, round, biconvex, scored tablets, debossed (5012) and (V) on one side and plain on the reverse side.

The specific lot numbers affected are:

- Hydrocodone Bitartrate and Acetaminophen Tablets, USP 10mg / 500mg, NDC 0603-3888-20, 60 count, Lot Numbers T150G10B, T120J10E and T023M10A
- Phenobarbital Tablets, USP 32.4 mg, NDC 0603-5166-32, 1000 count, Lot Numbers T150G10B, T120J10E and T023M10A

Thank you for your attention to this matter.

Sincerely,

A handwritten signature in black ink that reads "Eric J. Berman, DO, MS".

Eric J. Berman, DO, MS
Chief Medical Officer
Pennsylvania Managed Care