

FDA RECALLS TESTS MANUFACTURED BY UNIVERSAL MEDITECH, INC.

The FDA (Food and Drug Administration) recently released a statement warning consumers and patients to no longer use specific tests manufactured by Universal Meditech, Inc. including their One Step Pregnancy Test, HealthWiser UriTest UTI Test Strips and PrestiBio Ovulation Strips. Below is the full list of tests included in this recall and what you should do if you have or recently used any of these products.

RECALLED TESTS

- One Step Pregnancy Test
- DiagnosUS One Step Ovulation Test
- HealthyWiser UriTest 10 Parameter Reagent Test Strips for Urinalysis
- HealthyWiser UriTest UTI Test Strips
- HealthyWiser KetoFast Ketone Test Strips
- HealthyWiser pH-Aware pH Test Strips
- To Life hCG Pregnancy Urine Test
- Am I Pregnant Pregnancy Midstream Test
- DeTec hCG Pregnancy Urine Test
- PrestiBio Pregnancy Strips
- PrestiBio Rapid Detection Pregnancy Test Midstream
- PrestiBio Ovulation Strips
- PrestiBio Urinalysis Test Strip 10 Parameters
- PrestiBio Ketone Test Strips
- PrestiBio Breast Milk Alcohol Test Strips



This recall occurred after Universal Meditech, Inc. told the FDA it will no longer provide support for its tests. Since the FDA cannot confirm the performance of UMI's tests, their safety and effectiveness cannot be ensured.

WHAT TO DO IF YOU HAVE THESE TESTS:

Destroy and discard the test kits. **Currently, no refunds or replacements are being offered in relation to this recall**

If you've recently used these tests, consider discussing the accuracy of the results with your healthcare provider. If you still desire results from similar tests, purchase FDA approved testing kits.



For more information on recalled tests visit:

www.fda.gov/safety/recalls-market-withdrawals-safety-alerts