

Laboratory Detection of *Trichomonas vaginalis* by Nucleic Acid Amplification

To: Clients of UR Medicine Labs

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According to data from the CDC, trichomoniasis is a highly prevalent sexually transmitted infection affecting women and men. With respect to available laboratory test methods for the detection of *Trichomonas vaginalis* in clinical specimens, Nucleic Acid Amplification Testing (NAAT) is significantly more sensitive than traditional detection methods, i.e., microscopy (50%-60% sensitive) and culture (75% to 95% sensitive).

Beginning December 21st, we will provide testing for *T. vaginalis* by NAAT on the BD Viper platform (Becton Dickinson ProbeTec™) which we currently use for detection of *Chlamydia trachomatis* and *Neisseria gonorrhoeae*. Using this method, a single provider-collected swab of the endocervix or a single self-collected swab of the vagina can be used to detect nucleic acid from one or all three organisms; although urine is an FDA-cleared specimen type it is the least preferred specimen due to potential cross-contamination of specimens by the “leaky” urine container. Until urines from males are in-house validated for testing by NAAT, culture for *T. vaginalis* in male urines is the preferred method. The NAAT for *T. vaginalis* will be performed Monday thru Friday (same as for Chlamydia/GC). Until individual EMRs are updated to include this test, clients should request “Trichomonas NAAT”.

Refer to the UR Medicine Clinical Labs Test Index for availability of specimen collection/transport devices and details of specimen collection and transport to the laboratory: www.testmenu.com/rochester. For additional information contact Dr. Hardy by any of the methods listed above.