

<b>Statement of Deficiencies</b>	<b>(X1) Provider/Supplier/CLIA Identification Number</b>  46D2049037	<b>(X3) Date Survey Completed</b>  08/16/2018
<b>Name of Provider or Supplier</b>  Planned Parenthood Assn Of Utah-Metro Clinc	<b>Street Address, City, State</b>  160 S 1000 E Suite 120, Salt Lake City, UT	
For information on the provider's plan to correct this deficiency, please contact the provider or the state survey agency.		

<b>(X4) ID Prefix Tag</b>	<b>Summary Statement of Deficiencies</b>
<b>D3029</b>	<p><b>RETENTION REQUIREMENTS</b> CFR(s): 493.1105(a)(2)</p> <p>Test procedures. Retain a copy of each test procedure for at least 2 years after a procedure has been discontinued. Each test procedure must include the dates of initial use and discontinuance.</p> <p>This STANDARD is not met as evidenced by: Based on procedure manual review and interview with staff, the laboratory failed to document the date the director approved discontinuing potassium hydroxide (KOH) and wet preparation testing. Findings include: 1. The laboratory procedure manual review included the procedure for KOH and wet preparation testing. 2. In an interview with staff on 08/16/2018 at approximately 4:00 P.M. staff confirmed the lab had discontinued KOH and wet preparation testing in 2017 but had did not include the date the discontinuance date on the procedure.</p>