

<b>Statement of Deficiencies</b>	<b>(X1) Provider/Supplier/CLIA Identification Number</b> 10D0998053	<b>(X3) Date Survey Completed</b> 09/13/2022
<b>Name of Provider or Supplier</b> Gastro-Intestinal Consultants Of	<b>Street Address, City, State</b> 2134 Vindale Rd, Tavares, FL	
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>D0000</b>	A recertification survey was conducted on September 13, 2022. Gastro-Intestinal Consultants of Central Florida clinical laboratory was not in compliance with 42 CFR 493, requirements for clinical laboratories.
<b>D5609</b>	<p><b>HISTOPATHOLOGY</b> CFR(s): 493.1273(e)(f)</p> <p>(e) The laboratory must use acceptable terminology of a recognized system of disease nomenclature in reporting results. (f) The laboratory must document all control procedures performed, as specified in this section.</p> <p>This STANDARD is not met as evidenced by: Based on record review and interview, the laboratory failed to document quality control information including the expiration dates, open dates and lot numbers for all reagents used in their Hematoxylin &amp; Eosin (H &amp; E), Alcian Blue/Periodic Acid Schiff's (AB/PAS), and Diff Quik from 12/21/2021 to 09/13/2022. Findings: Review of the laboratory's procedure for quality control records revealed there was no reagent log. On 09/13/2022 at 10:47 AM, the Histotechnologist stated she had never seen a reagent log in the laboratory.</p>