Statement of Deficiencies	**(X1) Provider/Supplier/CLIA Identification Number**	**(X3) Date Survey Completed**
	10D0274847	10/31/2019
Name of Provider or Supplier	**Street Address, City, State**	
Quest Diagnostics Clinical Laboratories Inc	2423 Sand Lake Road Suite 110b, Orlando, FL	

For information on the provider's plan to correct this deficiency, please contact the provider or the state survey agency.

(X4) ID Prefix Tag	Summary Statement of Deficiencies
D0000	A recertification survey was conducted on October 31, 2019. Quest Diagnostics Clinical Laboratories Inc. was found not in compliance with 42 CFR 493, requirements for clinical laboratories.
D2010	TESTING OF PROFICIENCY TESTING SAMPLES CFR(s): 493.801(b)(2) The laboratory must test samples the same number of times that it routinely tests patient samples. This STANDARD is not met as evidenced by: Based on record review and interview, the laboratory failed to run proficiency testing (PT) samples the same number of times, as it routinely tests patient samples for 2019 1st event for the specialty of Urinalysis. Findings: Review of the College of American Pathologists (CAP) PT records showed that for the 1st event of 2019 there were two sets of handwritten urinalysis cell identification results. The attestation for the 1st event for 2019 was signed by Testing Personnel B and the first set of handwritten results had Testing Personnel B's name written on it . The second set of result had Testing Personnel A's name on it and was dated 5/14/19. The CAP submission date for PT results for the 2019 1st event was 5/28/19. During an interview on 10/30/19 at 12:15 PM, the Testing Personnel A stated that he performed the cell identification to use for their comparison study but was unaware he needed to wait until after the submission date.