Statement of Deficiencies	**(X1) Provider/Supplier/CLIA Identification Number**	**(X3) Date Survey Completed**
	10D2146458	11/19/2019
Name of Provider or Supplier	**Street Address, City, State**	
Women's Center Of Fort Lauderdale, Llc	2161 E Commercial Blvd Ste 400, Fort Lauderdale, 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	An announced CLIA recertification survey was conducted at Women's Center of Fort Lauderdale on 11/19/19. The laboratory is not in compliance with 42 CFR Part 493, Requirements for Laboratories. The following is a description of the standard level deficiencies:
D5417	TEST SYSTEMS, EQUIPMENT, INSTRUMENTS, REAGENT CFR(s): 493.1252(d) Reagents, solutions, culture media, control materials, calibration materials, and other supplies must not be used when they have exceeded their expiration date, have deteriorated, or are of substandard quality. This STANDARD is not met as evidenced by: Based on record review and interview with the Office Manager the laboratory used expired negative control for Rh typing for 5 days (08/14/18, 08/15/18, 08/16/18, 08/17/18, and 08/18/18) out of 4 months reviewed (09/19, 06/19, 08/18, and 05/18). Findings Included: Review of daily lab logs revealed on 08/14/18, 08/15/18, 08/16/18, 08/17/18, and 08/18/19 the negative control for Anti-D Rh testing expired on 08/12/18. There were 5 patients tested on 08/14/19, 7 patients on 08/15/18, 3 patients on 08/16/18, 2 patients on 08/17/18, and 13 patients on 08/18/18. Interview on 11/19/19 at 11:00 AM the Office Manager confirmed that the logs showed an expired negative control was used on the aforementioned dates.