Statement of Deficiencies	**(X1) Provider/Supplier/CLIA Identification Number**	**(X3) Date Survey Completed**
	29D2106915	04/21/2025
Name of Provider or Supplier	**Street Address, City, State**	
Center For Wellness And Pain Care Of Las Vegas	6930 S Cimarron Rd Ste 260, Las Vegas, NV	

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	This Statement of Deficiencies was created as a result of an on-site CLIA recertification survey conducted at your facility on April 21, 2025. The findings and conclusions of any investigation by the Division of Public and Behavioral Health shall not be construed as prohibiting any criminal or civil investigations, actions or other claims for relief that may be available to any party under applicable federal, state, or local laws.
D2007	TESTING OF PROFICIENCY TESTING SAMPLES CFR(s): 493.801(b)(1) (b)(1) The samples must be examined or tested with the laboratory's regular patient workload by personnel who routinely perform the testing in the laboratory, using the laboratory's routine methods. This STANDARD is not met as evidenced by: Based on a review of the 2024 American Proficiency Institute (API) Proficiency Testing (PT) records, a review of the director approved policy and procedure for proficiency testing, a completed CMS-209 form listing the technical personnel of the laboratory, and an interview with the laboratory supervisor, the laboratory failed to ensure that the proficiency testing specimens were rotated among all testing personnel. Findings include: 1. A review of the 2024 API Miscellaneous Chemistry PT records revealed that testing personnel number two listed on the CMS-209 form failed to participate in the performance of testing for PT Test Event one and Test event two for the urine drug screens. 2. The director approved policy entitled 'Proficiency Testing' stated, "Samples should be rotated among the testing employees actively working in the laboratory at the time of the survey. If applicable, samples will be divided among multiple testing employees." 3. The findings were confirmed during an interview with the laboratory supervisor conducted on April 21, 2025 at approximately 10:00 AM. The laboratory performs approximately 300 chemistry tests annually.

D3031	RETENTION REQUIREMENTS CFR(s): 493.1105(a)(3) Analytic systems records. Retain quality control and patient test records (including instrument printouts, if applicable) and records documenting all analytic systems activities specified in 493.1252 through 493.1289 for at least 2 years. In addition, retain the following: This STANDARD is not met as evidenced by: Based on a random patient audit of six patients tested between the dates September 7, 2023 and February 17, 2025, and an interview with the laboratory supervisor, the laboratory failed to ensure that the calibration records for the urine drug screen tests were retained for two years. Findings include: 1. A random patient audit of six patients tested between the dates of September 7, 2023 and February 17, 2025 revealed that there were no records of the urine drug screen calibrations for the following dates of patients tested: September 7, 2023, November 21, 2023 and January 9, 2024. 2. The findings were confirmed during an interview with the laboratory supervisor conducted on April 21, 2025 at approximately 11:30 AM. The laboratory performs approximately 300 chemistry tests annually.