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CLIA Laboratory Details

Please note that all of the laboratories found using the CMS CLIA Laboratory Demographic Information Tool on the CMS website are certified by the United States Government Department of Health and Human Services under the Clinical Laboratory Improvement Amendments of 1988 (CLIA), 42 U.S.C. §263a to perform laboratory testing as of the Data Source Date listed. Please note the most up to date information may not appear on the linked certificate.

CLIA Identification Number: 05D0592241

Facility Name: EUROFINS ASCEND CLINICAL, LLC
Lab Director: DR. RUSSELL L. KERSCHMANN
Address: 435 OAKMEAD PARKWAY

SUNNYVALE, CA 94085-4709

Phone Number:800 800-5655Certificate Type:ComplianceCertificate Effective Date:09/14/2022Certificate Expiration Date:02/28/2025Facility Type:Independent

CENTERS FOR MEDICARE & MEDICAID SERVICES CLINICAL LABORATORY IMPROVEMENT AMENDMENTS

CERTIFICATE OF COMPLIANCE

LABORATORY NAME AND ADDRESS

ASCEND CLINICAL, LLC 435 OAKMEAD PARKWAY SUNNYVALE, CA, 94085-1101 CLIA ID NUMBER

05D0592241

EFFECTIVE DATE

09/14/2020

LABORATORY DIRECTOR

RUSSELL KERSCHMANN

EXPIRATION DATE

05/31/2023

Pursuant to Section 353 of the Public Health Services Act (42 U.S.C. 263a) as revised by the Clinical Laboratory Improvement Amendments (CLIA), the above named laboratory located at the address shown hereon (and other approved locations) may accept human specimens for the purposes of performing laboratory examinations or procedures.

This certificate shall be valid until the expiration date above, but is subject to revocation, suspension, limitation, or other sanctions for violation of the Act or the regulations promulgated thereunder.



HEMATOLOGY (400)

Gregg Brandush, Director

Division of Clinical Laboratory Improvement & Quality

Quality & Safety Oversight Group Center for Clinical Standards and Quality

If you currently hold a Certificate of Compliance or Certificate of Accreditation, below is a list of the laboratory specialties/subspecialties you are certified to perform and their effective date:

LAB CERTIFICATION (CODE)	EFFECTIVE DATE	LAB CERTIFICATION (CODE)	EFFECTIVE DATE
MICROBIOLOGY - BACTERIOLOGY (110)	09/14/1994		
MICROBIOLOGY - MYCOLOGY (120)	04/01/2021		
MICROBIOLOGY - VIROLOGY (140)	01/04/2021		
DIAGNOSTIC IMMUNOLOGY - GENERAL IMMUNOLOGY (220)	09/14/1994		
CHEMISTRY - ROUTINE CHEMISTRY (310)	09/14/1994		
CHEMISTRY - URINALYSIS (320)	09/07/2010		
CHEMISTRY - ENDOCRINOLOGY (330)	09/14/1994		
CHEMISTRY - TOXICOLOGY (340)	09/14/1994		

09/14/1994



State of California—Health and Human Services Agency California Department of Public Health



March 24, 2023

Russell Kerschmann, Laboratory Director ASCEND CLINICAL, LLC 435 OAKMEAD PARKWAY SUNNYVALE, CA 94085

RE: VERIFICATION OF CERTIFICATION

CLIA Number: 05D 0592241

Dear Dr. Kerschmann,

The entity listed at the above address is currently certified under the Clinical Laboratory Improvement Amendments of 1988 (CLIA) program with a Certificate of Compliance and continues to meet all the appropriate regulatory requirements until a survey is completed.

This letter is proof, until such time that all other appropriate documents are issued, that the above stated entity continues to participate in the CLIA program.

If you have any questions regarding this letter, please call Donna McCallum at (213) 620-6570.

Sincerely,

Donna McCallum Section Chief, CLIA

Department of Public Health Laboratory Field Services



State of California-Health and Human Services Agency

California Department of Public Health



TOMÁS J. ARAGÓN, M.D., Dr.P.H. Director and State Public Health Officer

> December 19, 2024 Russell Kerschmann, M.D., Director Eurofins Ascend Clinical, LLC 435 Oakmead Parkway Sunnyvale, CA 94085-4709

CLIA #05D0592241

State I.D. # CLF- 00004141

RE: CERTIFICATION OF COMPLIANCE

Dear Director/Owner(s):

In order for a laboratory to perform testing under the Clinical Laboratory Improvement Amendments of 1988 (CLIA), Public Law 100-578, it must comply with all CLIA requirements. These requirements are found in section 353 of the Public Health Service Act (42 U.S.C. § 263a) and 42 Code of Federal Regulations, Part 493 (42 C.F.R. § 493). Federal regulations require onsite surveys to determine whether or not a laboratory is in compliance with the applicable regulations. Compliance with these regulations is a condition of certification for the CLIA program.

California Department of Public Health, Laboratory Field Services conducted a Recertification survey of your laboratory on December 10, 2024. The results of the survey showed that all CLIA Condition-level requirements were met during the time of the onsite survey. We are recommending to CMS that your laboratory be recertified in the CLIA program.

Please note that the routine survey takes an overview of the laboratory through random sampling. By its nature, the routine survey may not find every instance of non-compliance that may have occurred in the laboratory. It remains the responsibility of the laboratory and its director to ensure that the laboratory is at all times following all CLIA requirements, to identify any problems in the laboratory and take corrective action specific to the problems, and to institute appropriate quality assessment measures to ensure that the deficient practices do not recur.

If you have questions regarding this letter, please contact me at (213) 620-2013.

Sincerely, Jaleh Samani, Examiner I Laboratory Field Services Enclosure: CMS-2567

