MORE THAN JUST TEST RESULTS.

cGMP Certification Statement

Facility Registration

FOCUS Scientific Services, Inc. dba FOCUS Laboratories is a contract testing laboratory operating in two separate facilities, each registered as a Drug Establishment with a business operation listed as Analysis, and therefore not required to maintain a Drug Master File, nor do we operate under GDFA.

FDA Establishment Identifier	DUNS	Business Operations	Address	Registration Expiration
3006576330	078488452	ANALYSIS;	894 Marcon Blvd Ste 150 Allentown, Pennsylvania (PA) 18109-9603 United States (USA)	12/31/2025
3017875888	117786631	ANALYSIS;	2660 Alt 19 Ste B Palm Harbor, Florida (FL) 34683-2629 United States (USA)	12/31/2025

Debarment Notification

FOCUS Scientific Services, Inc. dba FOCUS Laboratories has not been debarred by the FDA nor is currently involved in any debarment proceeding with the FDA. Determined by a signed and dated certification statement, no person employed by FOCUS Laboratories has currently or in the past five years been convicted of any crime described in Sections 306 (a) or (b) of the Generic Drug Enforcement Act of 1992. FOCUS Laboratories has not, and will not, use the services of any person debarred under Section 306 of the Generic Drug Enforcement Act of 1992.

Current Good Manufacturing Practices

FOCUS Scientific Services, Inc. dba FOCUS Laboratories has implemented current Good Manufacturing Practices (cGMPs) as stated in the code of Federal Regulations. These regulations apply to the facility or controls to be used for the “Manufacture, Processing, Packing or Holding of Drugs”. Although our facility does not perform these functions and are therefore not subject to these practices, we do operate within the following sections of the regulations: 21 CFR Parts 210 and 211, Subpart B, Subpart C, Subpart D, Subpart E, Subpart I, and Subpart J.

Anthony Grilli
Chief Executive Officer
FOCUS Laboratories
Phone: 610-866-7272