



# United States Pharmacopeia Quality Systems GMP Audited Certificate Dietary Supplement Manufacturing

**Name of Holder:** Premier Research Labs LLC

**Manufacturing Site Address:** 3500B Wadley Place, Austin, TX 78728

**Certificate Number:** QSGMP-PRL-01

**Valid:** October 25, 2017 – September 30, 2018

**Initial Verification Date:** October 25, 2017

Premier Research Labs LLC voluntarily applied for a Good Manufacturing Practice (GMP) audit of their facility under the USP Quality Systems GMP Audited Verification Program. During the verification process, USP conducted an on-site GMP audit of the manufacturing site referenced above. After auditing the facility and examining the information the manufacturer provided to USP during the GMP auditing process, USP finds that the dietary supplement manufacturer's quality system provides sufficient assurance that the site referenced above meets the applicable GMP audit requirements set forth in the 21 Code of Federal Regulations Part 111- *Current Good Manufacturing Practice in Manufacturing, Packaging, Labeling, or Holding Operations for Dietary Supplements*, and in USP General Chapter <2750> *Manufacturing Practices for Dietary Supplements*, in the current edition of the *United States Pharmacopeia–National Formulary*.

The assessment represents GMP audits conducted since May 2017, the latest of which was conducted from May 31-June 2, 2017. This certificate reflects the status of the manufacturing site at the time of the inspection previously noted. In order to maintain verification status, manufacturing, packaging, labeling and holding operations of dietary supplements must continue to take place under the same conditions under which the firm was audited.

USP is pleased to award this certificate to Premier Research Labs LLC.

A handwritten signature in black ink that reads "John B. Atwater".

John B. Atwater, Ph.D.  
Senior Director, USP Verification Programs

This certificate remains the property of USP. USP issues this GMP certificate for the period stated above or until any major change in the manufacturer's quality systems has taken place, as defined in the license agreement and/or the USP Quality Systems GMP Audited Verification Program Manual for Participants. Failure to comply with the provisions of the manual or the license agreement shall render this certificate void, and the right to use the USP Quality Systems GMP Audited Mark will be withdrawn. To check the validity of this certificate, call +1-301-816-8273 or visit [www.uspverified.org](http://www.uspverified.org).