

**From:** [CDRH Registration and Listing](#)  
**To:** [Melinda Leishear](#)  
**Subject:** Registration Number 1121996: Successful 2025 Medical Device Establishment Registration  
**Date:** Friday, November 22, 2024 8:25:27 AM  
**Attachments:** [SignatureBlockLogo.png](#)

---

Dear Melinda Leishear:

This e-mail provides confirmation that the annual registration for the following medical device establishment has been successfully completed for 2025:

Registration Number: 1121996  
Owner Operator Number: 1121996  
AMERICAN I.V. PRODUCTS, INC.  
7485 SHIPLEY AVE.  
HARMANS, MD 21077  
UNITED STATES

If you do not see a registration number assigned to the establishment and your establishment previously had one, please send an email to [reglist@cdrh.fda.gov](mailto:reglist@cdrh.fda.gov) and include the registration number you believe is assigned to your establishment. We will review and determine if a duplicate registration has been created for your establishment.

Your registration is valid until December 31, 2025. Registration for 2026 will be conducted between October 1 and December 31, 2025.

Please note that registering your device facility and listing your devices does not, in any way, constitute FDA approval of your facility or your devices.

Should you have any questions, please send an e-mail to the CDRH Registration and Listing Helpdesk at [reglist@cdrh.fda.gov](mailto:reglist@cdrh.fda.gov).

CDRH Registration and Listing Helpdesk  
Imports & Registration and Listing Team  
Division 2 Establishment Support  
Office of Regulatory Programs  
Office of Product Evaluation and Quality  
Center for Devices and Radiological Health  
U.S. Food and Drug Administration

Tel: 301-796-7400, Option 1  
Email: [reglist@cdrh.fda.gov](mailto:reglist@cdrh.fda.gov)