From: CDRH Registration and Listing <reglist@CDRH.FDA.GOV>
Sent: Saturday, December 28, 2024 8:02 PM
To: USSTP Pentair Quality <PentairQuality@Pentair.com>
Subject: Registration Number 2132517: Successful 2025 Medical Device Establishment Registration

Dear Roger Hanzalik:

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

Registration Number: 2132517
Owner Operator Number: 9009435
Pentair Filtration Solutions
1350 Hammond Rd
Saint Paul, MN 55110
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 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.

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