FDA 510(k) Premarket Notification

K071101 TRU RSV

CLEARED

Oct 18, 2007
182 days to decision

K071101 · Product code: **GOG** · Microbiology
Source: https://www.510kdatabase.net/k071101/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Antisera, Cf, Poliovirus 1-3 (GOG)
Date received	Apr 19, 2007
Decision date	Oct 18, 2007
Days to decision	182 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Meridian Bioscience, Inc.**
Location	Cincinnati, OH, US
Contact	SUSAN ROLIH
Website	https://www.meridianbioscience.com
510(k) history	38 submissions · 37 cleared · 2003-2025

Meridian Bioscience, Inc. is a diagnostic and life science solutions company with a manufacturing facility in Cincinnati, US. The company develops integrated diagnostic products and molecular reagents for clinical and research applications. Meridian has received FDA 510(k) clearances from total submissions since 2003. The company specializes exclusively in Microbiology devices, including molecular detection assays, pathogen identification systems, and diagnostic analyzers. The latest clearance in 2025 reflects continued regulatory activity and product innovation. Recent c...