

K912408 MERIFLUOR-CRYPTOSPORIDIUM/GIARDIAJul 24, 1991
54 days to decisionK912408 · Product code: **MHI** · Microbiology
Source: <https://www.510kdatabase.net/k912408/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Giardia Spp. (MHI)
Date received	May 31, 1991
Decision date	Jul 24, 1991
Days to decision	54 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Meridian Diagnostics, Inc.
Location	Walker, MI, US
Contact	ALLEN D NICKOL
510(k) history	92 submissions · 92 cleared · 1980-1999

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k912408/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 14, 2026