

**K955157 PROSPECT GIARDIA/CRYPTOSPORIDIUM  
MICROPLATE ASSAY**Feb 9, 1996  
88 days to decisionK955157 · Product code: **MHI** · Microbiology  
Source: <https://www.510kdatabase.net/k955157/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Giardia Spp. (MHI)
Date received	Nov 13, 1995
Decision date	Feb 9, 1996
Days to decision	88 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

Company	<b>Alexon Biomedical, Inc.</b>
Location	Glen Ellyn, IL, US
Contact	SUSAN TURNER
510(k) history	5 submissions · 5 cleared · 1988-1997

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k955157/>. 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 June 29, 2026