FDA 510(k) Premarket Notification

K792693 CRYPTOCOCCUS NEOFORMANS ANTISERUM, LATEX

CLEARED

Jan 21, 1980
25 days to decision

K792693 · Product code: **GMD** · Microbiology
Source: https://www.510kdatabase.net/k792693/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Antisera, Latex Agglutination, Cryptococcus Neoformans (GMD)
Date received	Dec 27, 1979
Decision date	Jan 21, 1980
Days to decision	25 days
Third-party review	No

APPLICANT

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