

K792693 CRYPTOCOCCUS NEOFORMANS ANTISERUM, LATEXJan 21, 1980
25 days to decisionK792693 · 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

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Device record: <https://www.510kdatabase.net/k792693/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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