

K840012 INDIRECT FLUORESCENT ANTIBODY MYCOFeb 27, 1984
54 days to decisionK840012 · Product code: **GRZ** · Microbiology
Source: <https://www.510kdatabase.net/k840012/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Antisera, Fluorescent, All, Mycoplasma Spp. (GRZ)
Date received	Jan 4, 1984
Decision date	Feb 27, 1984
Days to decision	54 days
Third-party review	No

APPLICANT

Company	Zeus Technologies
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1983-1984

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Device record: <https://www.510kdatabase.net/k840012/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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