

K820468 PROFILE GENERAL SET ASSAYED SERUM CONTRMar 25, 1982
31 days to decisionK820468 · Product code: **DIF** · Toxicology
Source: <https://www.510kdatabase.net/k820468/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drug Mixture Control Materials (DIF)
Date received	Feb 22, 1982
Decision date	Mar 25, 1982
Days to decision	31 days
Third-party review	No

APPLICANT

Company	General Diagnostics
Location	Mchenry, IL, US
510(k) history	39 submissions · 39 cleared · 1976-1988

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Device record: <https://www.510kdatabase.net/k820468/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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