

K780544 DISC-PAK ROTATORApr 24, 1978
21 days to decisionK780544 · Product code: **GLE** · Hematology
Source: <https://www.510kdatabase.net/k780544/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Mixer, Blood Tube (GLE)
Date received	Apr 3, 1978
Decision date	Apr 24, 1978
Days to decision	21 days
Third-party review	No

APPLICANT

Company	Techhnilab Instruments, Inc.
Location	Mchenry, IL, US
510(k) history	6 submissions · 6 cleared · 1977-1981

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Device record: <https://www.510kdatabase.net/k780544/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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