FDA 510(k) Premarket Notification

K780544 DISC-PAK ROTATOR

CLEARED

Apr 24, 1978
21 days to decision

K780544 · 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