FDA 510(k) Premarket Notification

K834026 RDW PARAMETER FOR THE TECHNICON H6000

CLEARED

Jan 30, 1984
69 days to decision

K834026 · Product code: **GKL** · Hematology
Source: https://www.510kdatabase.net/k834026/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Counter, Cell, Automated (particle Counter) (GKL)
Date received	Nov 22, 1983
Decision date	Jan 30, 1984
Days to decision	69 days
Third-party review	No

APPLICANT

Company	**Technicon Instruments Corp.**
Location	Mchenry, IL, US
510(k) history	157 submissions · 156 cleared · 1976-1991