FDA 510(k) Premarket Notification

K831685 SF REFEEDING MEDIA

CLEARED

Jul 6, 1983
43 days to decision

K831685 · Product code: **KIT** · Pathology
Source: https://www.510kdatabase.net/k831685/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Media And Components, Synthetic Cell And Tissue Culture (KIT)
Date received	May 24, 1983
Decision date	Jul 6, 1983
Days to decision	43 days
Third-party review	No

APPLICANT

Company	**Viromed Laboratories, Inc.**
Location	Mchenry, IL, US
510(k) history	25 submissions · 25 cleared · 1983-1996

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k831685/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 14, 2026