FDA 510(k) Premarket Notification

K831580 PARAFFIN INFILTRATION MEDIUM

CLEARED

Jun 17, 1983
31 days to decision

K831580 · Product code: **KEO** · Pathology
Source: https://www.510kdatabase.net/k831580/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Formulations, Paraffin, All (KEO)
Date received	May 17, 1983
Decision date	Jun 17, 1983
Days to decision	31 days
Third-party review	No

APPLICANT

Company	**Surgipath Medical Industries, Inc.**
Location	Mchenry, IL, US
510(k) history	30 submissions · 30 cleared · 1977-1988