FDA 510(k) Premarket Notification

K881151 MICROSCRAPE

CLEARED

Apr 6, 1988
20 days to decision

K881151 · Product code: **LXK** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k881151/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Scrapper, Skin Specimen (LXK)
Date received	Mar 17, 1988
Decision date	Apr 6, 1988
Days to decision	20 days
Third-party review	No

APPLICANT

Company	**Ortho Pharmaceutical Corp.**
Location	Walker, MI, US
Contact	RUSSELL J HUME
510(k) history	9 submissions · 9 cleared · 1979-1996