FDA 510(k) Premarket Notification

K841624 PROBE

CLEARED

Aug 27, 1984
130 days to decision

K841624 · Product code: **HRX** · Orthopedic
Source: https://www.510kdatabase.net/k841624/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Arthroscope (HRX)
Date received	Apr 19, 1984
Decision date	Aug 27, 1984
Days to decision	130 days
Third-party review	No

APPLICANT

Company	**Plastafil, Inc.**
Location	Walker, MI, US
510(k) history	13 submissions · 13 cleared · 1984-1984