FDA 510(k) Premarket Notification

K810888 BOUNDARY TUBING HOLDERS

CLEARED

Apr 23, 1981
21 days to decision

K810888 · Product code: **FZX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k810888/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Guide, Surgical, Instrument (FZX)
Date received	Apr 2, 1981
Decision date	Apr 23, 1981
Days to decision	21 days
Third-party review	No

APPLICANT

Company	**The Buckeye Cellulose Corp.**
Location	Mchenry, IL, US
510(k) history	18 submissions · 18 cleared · 1980-1983