FDA 510(k) Premarket Notification

K882871 CASF DRILL GUIDE DG 170

CLEARED

Sep 23, 1988
74 days to decision

K882871 · Product code: **LYQ** · Orthopedic
Source: https://www.510kdatabase.net/k882871/

SUBMISSION DETAILS

Decision	Substantially Equivalent - SN
Submission type	Traditional
Device classification	Accessories, Fixation, Spinal Intervertebral Body (LYQ)
Date received	Jul 11, 1988
Decision date	Sep 23, 1988
Days to decision	74 days
Third-party review	No

APPLICANT

Company	**Terray Manufacturing, Inc.**
Location	Ontario K7s 1l8, CA
Contact	JUDY WILSON
510(k) history	9 submissions · 8 cleared · 1987-1992