FDA 510(k) Premarket Notification

K884888 MODIFIED NEW JERSEY ACETABULAR COMP. & BONE SCREWS

CLEARED

Dec 29, 1988
37 days to decision

K884888 · Product code: **JDI** · Orthopedic
Source: https://www.510kdatabase.net/k884888/

SUBMISSION DETAILS

Decision	Substantially Equivalent - SN
Submission type	Traditional
Device classification	Prosthesis, Hip, Semi-constrained, Metal/polymer, Cemented (JDI)
Date received	Nov 22, 1988
Decision date	Dec 29, 1988
Days to decision	37 days
Third-party review	No

APPLICANT

Company	**Protek, Inc.**
Location	Indianapolis, IN, US
Contact	KENNETH EPLING
510(k) history	25 submissions · 20 cleared · 1985-1989