

K884888 MODIFIED NEW JERSEY ACETABULAR COMP. & BONE SCREWSDec 29, 1988
37 days to decisionK884888 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k884888/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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