K010170 TRIDENT POROUS TITANIUM ACETABULAR COMPONENT

CLEARED

Apr 18, 2001
90 days to decision

K010170 · Product code: **LPH** · Orthopedic
Source: https://www.510kdatabase.net/k010170/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Hip, Semi-constrained, Metal/polymer, Porous Uncemented (LPH)
Date received	Jan 18, 2001
Decision date	Apr 18, 2001
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Howmedica Osteonics Corp.**
Location	Allendale, NJ, US
Contact	MARGARET F CROWE
510(k) history	288 submissions · 288 cleared · 1999-2020

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k010170/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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