FDA 510(k) Premarket Notification

K043223 DEPUY CAS KNEE INSTRUMENTATION

CLEARED

Mar 4, 2005
102 days to decision

K043223 · Product code: **HAW** · Neurology
Source: https://www.510kdatabase.net/k043223/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Neurological Stereotaxic Instrument (HAW)
Date received	Nov 22, 2004
Decision date	Mar 4, 2005
Days to decision	102 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**DePuy Orthopaedics, Inc.**
Location	Warsaw, IN, US
Contact	RANDA FRANKLIN
510(k) history	206 submissions · 204 cleared · 1998-2023

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k043223/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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