

K932070 DURACON STABILIZER FEMORAL COMPONENT AND INSERTMar 22, 1995
693 days to decisionK932070 · Product code: **JWH** · Orthopedic
Source: <https://www.510kdatabase.net/k932070/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent - SN
Submission type	Traditional
Device classification	Prosthesis, Knee, Patellofemorotibial, Semi-constrained, Cemented, Polymer/metal/polymer (JWH)
Date received	Apr 28, 1993
Decision date	Mar 22, 1995
Days to decision	693 days
Third-party review	No

APPLICANT

Company	Howmedica Corp.
Location	Mchenry, IL, US
Contact	MARGARET F CROWE
510(k) history	373 submissions · 325 cleared · 1976-1998

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k932070/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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