

K811630 KINEMATIC ROTATING HINGE TOTAL KNEE PR.Jul 1, 1981
22 days to decisionK811630 · Product code: **LGE** · Orthopedic
Source: <https://www.510kdatabase.net/k811630/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Knee, Femorotibial, Semi-constrained, Cemented, Trunnion-bearing (LGE)
Date received	Jun 9, 1981
Decision date	Jul 1, 1981
Days to decision	22 days
Third-party review	No

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k811630/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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