

K781870 IMPLANT, H.P. CONDYLARFeb 9, 1979
98 days to decisionK781870 · Product code: **KWD** · Orthopedic
Source: <https://www.510kdatabase.net/k781870/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Toe, Hemi-, Phalangeal (KWD)
Date received	Nov 3, 1978
Decision date	Feb 9, 1979
Days to decision	98 days
Third-party review	No

APPLICANT

Company	Dow Corning Corp. Healthcare Industries Materials
Location	Mchenry, IL, US
510(k) history	31 submissions · 31 cleared · 1976-1985

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k781870/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 22, 2026