

K960852 HALL MODULAR ACETABULAR REAMER SYSTEMMar 20, 1996
19 days to decisionK960852 · Product code: **HWE** · Orthopedic
Source: <https://www.510kdatabase.net/k960852/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Surgical, Orthopedic, Ac-powered Motor And Accessory/attachment (HWE)
Date received	Mar 1, 1996
Decision date	Mar 20, 1996
Days to decision	19 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Linvatec Corp.
Location	Research Triangle Pa, NC, US
Contact	CAROL A WEIDEMAN,PH.D.
510(k) history	93 submissions · 87 cleared · 1992-2009

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k960852/> 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 14, 2026