

K992120 DYNAWELL OR DYNAWELLAug 13, 1999
51 days to decisionK992120 · Product code: **LNH** · Radiology
Source: <https://www.510kdatabase.net/k992120/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Nuclear Magnetic Resonance Imaging (LNH)
Date received	Jun 23, 1999
Decision date	Aug 13, 1999
Days to decision	51 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Ekeroth Quality AB
Location	Stockholm, SE
Contact	NILS EKEROTH
510(k) history	1 submissions · 1 cleared · 1999-1999

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k992120/> 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 19, 2026