FDA 510(k) Premarket Notification

K992120 DYNAWELL OR DYNAWELL

CLEARED

Aug 13, 1999
51 days to decision

K992120 · 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