

K041020 CADSTREAM VERSION 3.1Jul 7, 2004
78 days to decisionK041020 · Product code: **LNH** · Radiology
Source: <https://www.510kdatabase.net/k041020/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Nuclear Magnetic Resonance Imaging (LNH)
Date received	Apr 20, 2004
Decision date	Jul 7, 2004
Days to decision	78 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Confirma, Inc.
Location	Kirkland, WA, US
Contact	PATRICIA A MILBANK
510(k) history	9 submissions · 9 cleared · 2002-2009

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k041020/> 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 17, 2026