FDA 510(k) Premarket Notification

K041020 CADSTREAM VERSION 3.1

CLEARED

Jul 7, 2004
78 days to decision

K041020 · 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