

K014300 SAFETOUCHMar 1, 2002
60 days to decisionK014300 · Product code: **NHM** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k014300/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Sheet, Recording, Breast Examination (NHM)
Date received	Dec 31, 2001
Decision date	Mar 1, 2002
Days to decision	60 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	James D Rusin MD, MBA
Location	Anoka, MN, US
Contact	JAMES D RUSIN
510(k) history	1 submissions · 1 cleared · 2002-2002

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k014300/> 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 June 29, 2026