FDA 510(k) Premarket Notification

K032121 MODIFICATION TO E-SCAN XQ

CLEARED

Aug 13, 2003
35 days to decision

K032121 · Product code: **LNH** · Radiology
Source: https://www.510kdatabase.net/k032121/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	System, Nuclear Magnetic Resonance Imaging (LNH)
Date received	Jul 9, 2003
Decision date	Aug 13, 2003
Days to decision	35 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Esaote, S.P.A.**
Location	Genoa, IT
Contact	COLLEEN J DENSMORE
Website	https://www.esaote.com
510(k) history	67 submissions · 67 cleared · 2003-2026

Esaote, S.P.A. is a medical diagnostic imaging company based in Genoa, Italy. The company specializes in ultrasound, MRI, and healthcare IT solutions for clinical settings. Esaote has received FDA 510(k) clearances from total submissions since 2003. The company's regulatory portfolio is dominated by Radiology devices, representing 100% of its FDA submissions. Recent cleared devices include the MyLab ultrasound systems and Magnifico Open imaging platforms. The company remains actively engaged in FDA regulatory submissions, with the latest clearance in 2026. Esaote's produc...