FDA 510(k) Premarket Notification

K890999 MODIFIED QDR-1000 X-RAY BONE DENSITOMETER

CLEARED

Dec 21, 1989
297 days to decision

K890999 · Product code: **KGI** · Radiology
Source: https://www.510kdatabase.net/k890999/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Densitometer, Bone (KGI)
Date received	Feb 27, 1989
Decision date	Dec 21, 1989
Days to decision	297 days
Third-party review	No

APPLICANT

Company	**Hologic, Inc.**
Location	Waltham, MA, US
Contact	THOMAS L KELLY
Website	https://www.hologic.com/
510(k) history	115 submissions · 111 cleared · 1987-2025

Hologic, Inc. is a medical device company headquartered in Waltham, Massachusetts. The company specializes in women's health, diagnostics, and medical imaging technologies. Hologic has maintained a strong FDA 510(k) regulatory record since its founding in 1987. The company has received FDA 510(k) clearances from total submissions. Recent cleared devices span microbiology, radiology, and obstetrics & gynecology categories. The latest clearance in 2025 demonstrates continued active development and regulatory engagement. Hologic's cleared device portfolio includes molecular ...