FDA 510(k) Premarket Notification

K231202 QuickRayPRO

CLEARED

Jun 22, 2023
56 days to decision

K231202 · Product code: **MUH** · Radiology
Source: https://www.510kdatabase.net/k231202/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Extraoral Source, Digital (MUH)
Date received	Apr 27, 2023
Decision date	Jun 22, 2023
Days to decision	56 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Denterprise International, Inc.**
Location	Ormond Beach, FL, US
Contact	Claude Berthoin
510(k) history	4 submissions · 4 cleared · 2012-2023