FDA 510(k) Premarket Notification

K032260 ARAMIS II DERMATOLOGICAL LASER

CLEARED

Nov 20, 2003
121 days to decision

K032260 · Product code: **GEX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k032260/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Jul 22, 2003
Decision date	Nov 20, 2003
Days to decision	121 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Quantel Medical**
Location	Cournon D'Auvergne-Cedex, FR
Contact	ROGER W BARNES
Website	https://www.quantelmedical.com
510(k) history	30 submissions · 30 cleared · 2000-2026