FDA 510(k) Premarket Notification

K012042 KINETIC CANNULA

CLEARED

Sep 20, 2001
83 days to decision

K012042 · Product code: **QPB** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k012042/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Suction, Lipoplasty For Removal (QPB)
Date received	Jun 29, 2001
Decision date	Sep 20, 2001
Days to decision	83 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Reliance Medical Corp.**
Location	Tucson, AZ, US
Contact	ROBERT W NICKS
510(k) history	3 submissions · 3 cleared · 1985-2001