FDA 510(k) Premarket Notification

K924081 LUMISONIC

CLEARED

Nov 9, 1992
88 days to decision

K924081 · Product code: **CAF** · Anesthesiology
Source: https://www.510kdatabase.net/k924081/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Nebulizer (direct Patient Interface) (CAF)
Date received	Aug 13, 1992
Decision date	Nov 9, 1992
Days to decision	88 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Lumiscope Co., Inc.**
Location	Edison, NJ, US
Contact	STEPHEN WHITTENBURG
510(k) history	13 submissions · 13 cleared · 1988-2003