FDA 510(k) Premarket Notification

K933369 CUDA PRODUCTS CORP. LIGHTSOURCES

CLEARED

Jan 3, 1994
178 days to decision

K933369 · Product code: **FCW** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k933369/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light Source, Fiberoptic, Routine (FCW)
Date received	Jul 9, 1993
Decision date	Jan 3, 1994
Days to decision	178 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Cuda Products Co.**
Location	Jacksonville, FL, US
Contact	CYNTHIA ARCUSA
510(k) history	17 submissions · 17 cleared · 1988-1998