FDA 510(k) Premarket Notification

K010534 CROSS-CHECKS DUAL, MODEL CI 125

CLEARED

Aug 6, 2001
164 days to decision

K010534 · Product code: **JOJ** · General Hospital
Source: https://www.510kdatabase.net/k010534/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Indicator, Physical/chemical Sterilization Process (JOJ)
Date received	Feb 23, 2001
Decision date	Aug 6, 2001
Days to decision	164 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Steritec Products, Inc.**
Location	Castle Rock, CO, US
Contact	LINDA NELSON
510(k) history	12 submissions · 12 cleared · 2000-2009