FDA 510(k) Premarket Notification

CLEARED

K961524 STERILAB CANNULAS, STERILAB CYSTOTMES, STERILAB NEEDLES AND STERILAB LENS MANIPULATORS

May 16, 1996
24 days to decision

K961524 · Product code: **HNY** · Ophthalmic
Source: https://www.510kdatabase.net/k961524/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cystotome (HNY)
Date received	Apr 22, 1996
Decision date	May 16, 1996
Days to decision	24 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Sterilab, Inc.**
Location	Carlsbad, CA, US
Contact	JONATHAN WOODWARD
510(k) history	1 submissions · 1 cleared · 1996-1996