FDA 510(k) Premarket Notification

K880240 INSTANT EOSIN-ALCOHOLIC

CLEARED

Mar 3, 1988
43 days to decision

K880240 · Product code: **HYB** · Pathology
Source: https://www.510kdatabase.net/k880240/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Eosin Y (HYB)
Date received	Jan 20, 1988
Decision date	Mar 3, 1988
Days to decision	43 days
Third-party review	No

APPLICANT

Company	**Shandon, Inc.**
Location	Mchenry, IL, US
Contact	CROOKHAM, BA
510(k) history	5 submissions · 5 cleared · 1978-1989