FDA 510(k) Premarket Notification

K023092 BD K-4000 MICROKERATOME SYSTEM

CLEARED

Oct 18, 2002
30 days to decision

K023092 · Product code: **HMY** · Ophthalmic
Source: https://www.510kdatabase.net/k023092/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Keratome, Battery-powered (HMY)
Date received	Sep 18, 2002
Decision date	Oct 18, 2002
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Becton, Dickinson & CO**
Location	Franklin Lakes, NJ, US
Contact	EILEEN T SCHWEIGHARDT
510(k) history	190 submissions · 190 cleared · 2001-2016

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k023092/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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