FDA 510(k) Premarket Notification

K030353 MONOCANALICULAR STENTS, LACRIMAL INTUBATION SETS

CLEARED

Apr 25, 2003
81 days to decision

K030353 · Product code: **OKS** · Ophthalmic
Source: https://www.510kdatabase.net/k030353/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lacrimal Stents And Intubation Sets (OKS)
Date received	Feb 3, 2003
Decision date	Apr 25, 2003
Days to decision	81 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Eagle Vision, Inc.**
Location	Bethesda, MD, US
Contact	BILLY HANNAFORD
510(k) history	7 submissions · 7 cleared · 1988-2006