FDA 510(k) Premarket Notification

K883233 EV MONOCANALICULAR STENT (TM)

CLEARED

Aug 22, 1988
21 days to decision

K883233 · Product code: **OKS** · Ophthalmic
Source: https://www.510kdatabase.net/k883233/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lacrimal Stents And Intubation Sets (OKS)
Date received	Aug 1, 1988
Decision date	Aug 22, 1988
Days to decision	21 days
Third-party review	No

APPLICANT

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