FDA 510(k) Premarket Notification

CLEARED

K990672 LACRIMAL INTUBATION SET, DACRYOCYSTORHINOSTOMY (DCR) SET

May 18, 1999
77 days to decision

K990672 · Product code: **OKS** · Ophthalmic
Source: https://www.510kdatabase.net/k990672/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lacrimal Stents And Intubation Sets (OKS)
Date received	Mar 2, 1999
Decision date	May 18, 1999
Days to decision	77 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Hurricane Medical**
Location	Bradenton, FL, US
Contact	DAVID A CLAPP
510(k) history	3 submissions · 3 cleared · 1999-2000