FDA 510(k) Premarket Notification

K791351 CEMENT SPACERS

CLEARED

Aug 3, 1979
17 days to decision

K791351 · Product code: **KYB** · Ophthalmic
Source: https://www.510kdatabase.net/k791351/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lens, Guide, Intraocular (KYB)
Date received	Jul 17, 1979
Decision date	Aug 3, 1979
Days to decision	17 days
Third-party review	No

APPLICANT

Company	**Depuy, Inc.**
Location	Mchenry, IL, US
510(k) history	303 submissions · 239 cleared · 1976-2005