

K791351 CEMENT SPACERSAug 3, 1979
17 days to decisionK791351 · 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

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Device record: <https://www.510kdatabase.net/k791351/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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