FDA 510(k) Premarket Notification

K897021 MARTIN-HENSLEY SEGMENT LENS

CLEARED

Jun 22, 1990
185 days to decision

K897021 · Product code: **HPX** · Ophthalmic
Source: https://www.510kdatabase.net/k897021/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lens, Contact (polymethylmethacrylate) (HPX)
Date received	Dec 19, 1989
Decision date	Jun 22, 1990
Days to decision	185 days
Third-party review	No

APPLICANT

Company	**Bruce W. Martin O.D. and Edward Hensley**
Location	Gladwin, MI, US
Contact	MARTIN, OD
510(k) history	1 submissions · 1 cleared · 1990-1990