FDA 510(k) Premarket Notification

K920571 TRIAL LENS SET

CLEARED

May 7, 1992
90 days to decision

K920571 · Product code: **HPC** · Ophthalmic
Source: https://www.510kdatabase.net/k920571/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Lens, Trial, Ophthalmic (HPC)
Date received	Feb 7, 1992
Decision date	May 7, 1992
Days to decision	90 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**R.H. Burton Co.**
Location	Grove City, OH, US
Contact	TAMMY DERN
510(k) history	19 submissions · 19 cleared · 1992-1993