FDA 510(k) Premarket Notification

K894720 UNIVERSAL EYESHIELD

CLEARED

Sep 14, 1989
51 days to decision

K894720 · Product code: **HOY** · Ophthalmic
Source: https://www.510kdatabase.net/k894720/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Shield, Eye, Ophthalmic (including Sunlamp Protective Eyewear And Post-mydriatic Eyewear) (HOY)
Date received	Jul 25, 1989
Decision date	Sep 14, 1989
Days to decision	51 days
Third-party review	No

APPLICANT

Company	**Trident Medical Products, Inc.**
Location	Fort Worth, TX, US
Contact	DAN E BRUHL
510(k) history	3 submissions · 3 cleared · 1989-1989