FDA 510(k) Premarket Notification

K924780 BURTON OPHTHALMIC EXAM UNIT - MODEL 2201

CLEARED

Apr 5, 1993
194 days to decision

K924780 · Product code: **HMF** · Ophthalmic
Source: https://www.510kdatabase.net/k924780/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stand, Instrument, Ac-powered, Ophthalmic (HMF)
Date received	Sep 23, 1992
Decision date	Apr 5, 1993
Days to decision	194 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

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