FDA 510(k) Premarket Notification

K934880 MTI PHOTOSCREENER

CLEARED

Apr 6, 1994
176 days to decision

K934880 · Product code: **MMF** · Ophthalmic
Source: https://www.510kdatabase.net/k934880/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Photorefractor (MMF)
Date received	Oct 12, 1993
Decision date	Apr 6, 1994
Days to decision	176 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Medical Technology and Innovations, Inc.**
Location	Grand Prairie, TX, US
Contact	JEREMY P FEAKINS
510(k) history	3 submissions · 3 cleared · 1986-2002