FDA 510(k) Premarket Notification

K942180 KEELER MULTILASE 3000

CLEARED

Nov 22, 1994
202 days to decision

K942180 · Product code: **HQF** · Ophthalmic
Source: https://www.510kdatabase.net/k942180/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laser, Ophthalmic (HQF)
Date received	May 4, 1994
Decision date	Nov 22, 1994
Days to decision	202 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Keeler Instruments, Inc.**
Location	Mchenry, IL, US
Contact	EUGENE R VAN ARSDALE
510(k) history	60 submissions · 60 cleared · 1981-2019