FDA 510(k) Premarket Notification

K911721 KEELER PULSAIR NON CONTACT TONOMETER, MODIFICATION

CLEARED

Jun 25, 1991
69 days to decision

K911721 · Product code: **HKX** · Ophthalmic
Source: https://www.510kdatabase.net/k911721/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tonometer, Ac-powered (HKX)
Date received	Apr 17, 1991
Decision date	Jun 25, 1991
Days to decision	69 days
Third-party review	No

APPLICANT

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