

**K911721 KEELER PULSAIR NON CONTACT TONOMETER,
MODIFICATION**Jun 25, 1991
69 days to decisionK911721 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k911721/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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