FDA 510(k) Premarket Notification

K933564 SKYTRON EXAM LIGHT

CLEARED

Feb 10, 1994
203 days to decision

K933564 · Product code: **KZF** · General Hospital
Source: https://www.510kdatabase.net/k933564/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Medical Examination, Ac Powered (KZF)
Date received	Jul 22, 1993
Decision date	Feb 10, 1994
Days to decision	203 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Skytron, Div. the Kmw Group, Inc.**
Location	Mchenry, IL, US
Contact	DAVID M MEHNEY
510(k) history	19 submissions · 19 cleared · 1981-2007