FDA 510(k) Premarket Notification

K023830 AIROCIDE TI02

CLEARED

Feb 4, 2003
78 days to decision

K023830 · Product code: **FRA** · General Hospital
Source: https://www.510kdatabase.net/k023830/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Purifier, Air, Ultraviolet, Medical (FRA)
Date received	Nov 18, 2002
Decision date	Feb 4, 2003
Days to decision	78 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Kes Science & Technology, Inc.**
Location	Kennesaw, GA, US
Contact	JOHN HAYMAN, JR
510(k) history	1 submissions · 1 cleared · 2003-2003