

**K023830 AIROCID T102**Feb 4, 2003  
78 days to decisionK023830 · Product code: **FRA** · General Hospital  
Source: <https://www.510kdatabase.net/k023830/>**SUBMISSION DETAILS**

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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**

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Company	<b>Kes Science &amp; Technology, Inc.</b>
Location	Kennesaw, GA, US
Contact	JOHN HAYMAN, JR
510(k) history	1 submissions · 1 cleared · 2003-2003

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k023830/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated July 2, 2026