

**K800023 LUNG AEROSOL UNIT**Jan 1, 1985  
1821 days to decisionK800023 · Product code: **CAF** · General Hospital  
Source: <https://www.510kdatabase.net/k800023/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Nebulizer (direct Patient Interface) (CAF)
Date received	Jan 7, 1980
Decision date	Jan 1, 1985
Days to decision	1821 days
Third-party review	No

**APPLICANT**

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Company	<b>Cadema Medical Products, Inc.</b>
Location	Walker, MI, US
510(k) history	6 submissions · 6 cleared · 1985-1990

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k800023/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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