

**K012886 CBYON SYSTEM WITH IMAGE ENHANCED  
FLUOROSCOPY MODULE**Oct 9, 2001  
42 days to decisionK012886 · Product code: **HAW** · Neurology  
Source: <https://www.510kdatabase.net/k012886/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Neurological Stereotaxic Instrument (HAW)
Date received	Aug 28, 2001
Decision date	Oct 9, 2001
Days to decision	42 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Cbyon, Inc.</b>
Location	Palo Alto, CA, US
Contact	KRISHNA SUDHAKARAN
510(k) history	2 submissions · 2 cleared · 2000-2001

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k012886/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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