

**K810588 SYNCHRONIZER #208**Apr 10, 1981  
37 days to decisionK810588 · Product code: **IXO** · Radiology  
Source: <https://www.510kdatabase.net/k810588/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Synchronizer, Ecg / Respirator, Radiographic (IXO)
Date received	Mar 4, 1981
Decision date	Apr 10, 1981
Days to decision	37 days
Third-party review	No

**APPLICANT**

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Company	<b>Brattle Instrument Corp.</b>
Location	Mchenry, IL, US
510(k) history	7 submissions · 7 cleared · 1977-1981

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

Device record: <https://www.510kdatabase.net/k810588/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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