

**K761332 HI-SENSITIVITY MODE OPTION FOR S60-1A**Dec 30, 1976  
3 days to decisionK761332 · Product code: **JAK** · Radiology  
Source: <https://www.510kdatabase.net/k761332/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Tomography, Computed (JAK)
Date received	Dec 27, 1976
Decision date	Dec 30, 1976
Days to decision	3 days
Third-party review	No

**APPLICANT**

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Company	<b>Syntex Corp.</b>
Location	Mchenry, IL, US
510(k) history	10 submissions · 10 cleared · 1976-1992

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Device record: <https://www.510kdatabase.net/k761332/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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