

**K823536 INTRALIG**Dec 28, 1982  
28 days to decisionK823536 · Product code: **EJI** · DentalSource: <https://www.510kdatabase.net/k823536/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Cartridge (EJI)
Date received	Nov 30, 1982
Decision date	Dec 28, 1982
Days to decision	28 days
Third-party review	No

**APPLICANT**

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Company	<b>Miltex Instrument Co.</b>
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1981-1992

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

Device record: <https://www.510kdatabase.net/k823536/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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