

**K930926 DURHAM**Oct 12, 1993  
231 days to decisionK930926 · Product code: **FPA** · General HospitalSource: <https://www.510kdatabase.net/k930926/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Feb 23, 1993
Decision date	Oct 12, 1993
Days to decision	231 days
Third-party review	No
Summary / Statement	Statement
Other names	Y CHECK SET-LUER LOCK W/INJECTION SITE

**APPLICANT**

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Company	<b>Intl. Medical Consultants</b>
Location	Washington Crossing, PA, US
Contact	JOHN ROMANO
510(k) history	6 submissions · 6 cleared · 1993-1993

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k930926/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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