

**K940073 EXTENSION TUBE, EXTENSION TUBE LENGTH 20, &  
30**Mar 23, 1994  
76 days to decisionK940073 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k940073/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Jan 6, 1994
Decision date	Mar 23, 1994
Days to decision	76 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Puritas Health Care, Inc.</b>
Location	Brookfield, CT, US
Contact	THOMAS P DIMAIO
510(k) history	24 submissions · 22 cleared · 1991-1994

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k940073/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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