

**K980994 IV ADMINISTRATION SETS WITH NEEDLE ACCESS  
DEVICES**May 29, 1998  
73 days to decisionK980994 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k980994/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Mar 17, 1998
Decision date	May 29, 1998
Days to decision	73 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Venetec Intl., Inc.</b>
Location	Irvine, CA, US
Contact	ANITA PELLETIER
510(k) history	2 submissions · 2 cleared · 1998-1998

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k980994/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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