

**K944320 VENTED/NON-VENTED GEMINI ADMINISTRATION SET**Feb 14, 1995  
169 days to decisionK944320 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k944320/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Aug 29, 1994
Decision date	Feb 14, 1995
Days to decision	169 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Imed Corp.</b>
Location	Mchenry, IL, US
Contact	AHMAD SAJADI
510(k) history	43 submissions · 43 cleared · 1977-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k944320/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 28, 2026