

**K992999 DAISY PROTECTED SCALP VEIN SET**Nov 5, 1999  
59 days to decisionK992999 · Product code: **FPA** · General HospitalSource: <https://www.510kdatabase.net/k992999/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Sep 7, 1999
Decision date	Nov 5, 1999
Days to decision	59 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Diasol, Inc.</b>
Location	North Hollywood, CA, US
Contact	MONICA ABELES
510(k) history	10 submissions · 10 cleared · 1999-2014

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k992999/> 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 June 7, 2026