

**K073520 ANGIOTECH CVC**Apr 11, 2008  
119 days to decisionK073520 · Product code: **FOZ** · General Hospital  
Source: <https://www.510kdatabase.net/k073520/>**SUBMISSION DETAILS**

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|                       |  |
|-----------------------|--|
| Decision              | Substantially Equivalent (Cleared)                                       |
| Submission type       | Traditional  |
| Device classification | Catheter, Intravascular, Therapeutic, Short-term Less Than 30 Days (FOZ) |
| Date received         | Dec 14, 2007   |
| Decision date         | Apr 11, 2008   |
| Days to decision      | 119 days   |
| Third-party review    | No   |
| Summary / Statement   | Summary  |

**APPLICANT**

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|----------------|--|
| Company        | <b>Angiotech Pharmaceuticals, Inc.</b> |
| Location       | Herndon, VA, US                        |
| Contact        | TRUDY D ESTRIDGE                       |
| 510(k) history | 1 submissions · 1 cleared · 2008-2008  |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k073520/> 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 25, 2026