

**K071189 CORDIS AVIATOR PLUS PTA BALLOON DILATATION  
CATHETER**May 16, 2007  
16 days to decisionK071189 · Product code: **LIT** · Cardiovascular  
Source: <https://www.510kdatabase.net/k071189/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Angioplasty, Peripheral, Transluminal (LIT)
Date received	Apr 30, 2007
Decision date	May 16, 2007
Days to decision	16 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Cordis Europa, N.V.</b>
Location	Warren, NJ, US
Contact	HARM HOVINGA
510(k) history	7 submissions · 5 cleared · 2001-2007

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k071189/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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