

**K091966 ANGIOSCULPT PTA SCORING BALLOON CATHETER,
MODELS 2076-6040, 2092-6040, 2105-6040**Jul 30, 2009
29 days to decisionK091966 · Product code: **PNO** · Cardiovascular
Source: <https://www.510kdatabase.net/k091966/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Percutaneous, Cutting/scoring (PNO)
Date received	Jul 1, 2009
Decision date	Jul 30, 2009
Days to decision	29 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Angioscore, Inc.
Location	Alameda, CA, US
Contact	KIMBERLEY KLINE
510(k) history	13 submissions · 13 cleared · 2005-2015

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k091966/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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