

K955111 MAXI-TORQUE PLUS ANGIOGRAPHIC CATHETERJan 31, 1996
84 days to decisionK955111 · Product code: **DQO** · Cardiovascular
Source: <https://www.510kdatabase.net/k955111/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Nov 8, 1995
Decision date	Jan 31, 1996
Days to decision	84 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Adam Spence Corp.
Location	Wall, NJ, US
Contact	AL WEISENBORN
510(k) history	3 submissions · 3 cleared · 1995-1999

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k955111/> Data sourced from FDA 510(k) public records (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. - Generated July 4, 2026