

**K895352 KENDALL INFANT ECMO VENOVENOUS CATHETER-DUAL LUMEN**

Nov 29, 1989  
89 days to decision

K895352 · Product code: **DWF** · Cardiovascular  
Source: <https://www.510kdatabase.net/k895352/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Sep 1, 1989
Decision date	Nov 29, 1989
Days to decision	89 days
Third-party review	No

**APPLICANT**

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Company	<b>Kendall Healthcare Products Co. Div.Of Tyco Health</b>
Location	Mansfield, MA, US
Contact	JANE ARNOLD
510(k) history	66 submissions · 50 cleared · 1989-1997

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k895352/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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