

**K123370 EZ GLIDE AORTIC CANNULA, AORTIC PERFUSION CANNULA WITH DURAFLO COATING, DISPERSION AORTIC PERFUSION CANNULA**Mar 13, 2013  
132 days to decisionK123370 · Product code: **DWF** · Cardiovascular  
Source: <https://www.510kdatabase.net/k123370/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Nov 1, 2012
Decision date	Mar 13, 2013
Days to decision	132 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Edwards Lifesciences, LLC</b>
Location	Irvine, CA, US
Contact	KAREN JONES
Website	<a href="https://www.edwards.com">https://www.edwards.com</a>
510(k) history	135 submissions · 129 cleared · 1979-2026

Edwards Lifesciences, LLC is a global structural heart innovation company headquartered in Irvine, California. The company specializes in advanced medical devices for cardiovascular disease management. Edwards Lifesciences has established a strong FDA 510(k) regulatory track record with FDA 510(k) cleared devices from total submissions since 1979. The company's portfolio is dominated by Cardiovascular devices, which represent 88% of all submissions. The latest clearance was received in 2026, demonstrating continued active development and regulatory engagement. Recent clea...