

**K102532 HLS CANNULA NON-COATED, WITH BIOLINE
COATING AND WITH SOFTLINE COATING**Feb 25, 2011
175 days to decisionK102532 · Product code: **DWF** · Cardiovascular
Source: <https://www.510kdatabase.net/k102532/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Sep 3, 2010
Decision date	Feb 25, 2011
Days to decision	175 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Maquet Cardiopulmonary, AG
Location	Fairfield, IA, US
Contact	INGRID RICHTER
510(k) history	44 submissions · 44 cleared · 2005-2015

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k102532/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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