

**K020784 JOSTRA SINGLE STAGE VENOUS RETURN
CATHETERS**Jan 9, 2003
304 days to decisionK020784 · Product code: **DWF** · Cardiovascular
Source: <https://www.510kdatabase.net/k020784/>**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 | Mar 11, 2002 |
| Decision date | Jan 9, 2003 |
| Days to decision | 304 days |
| Third-party review | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|-----------------------------------------|
| Company | Jostra AG |
| Location | Newark, DE, US |
| Contact | KATHLEEN JOHNSON |
| 510(k) history | 13 submissions · 13 cleared · 2000-2004 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k020784/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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