FDA 510(k) Premarket Notification

K151103 ASAHI Corsair Microcatheter

CLEARED

Jun 23, 2015
60 days to decision

K151103 · Product code: **DQY** · Cardiovascular
Source: https://www.510kdatabase.net/k151103/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Apr 24, 2015
Decision date	Jun 23, 2015
Days to decision	60 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Asahi Intecc Co., Ltd.**
Location	Seto-Shi, JP
Contact	YOSHIHIKO FUKUI
Website	https://www.asahi-intecc.com
510(k) history	83 submissions · 83 cleared · 2003-2026

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k151103/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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