

K072316 PRECIOUS PLUS GUIDE CATHETERSep 17, 2007
31 days to decisionK072316 · Product code: **DQY** · CardiovascularSource: <https://www.510kdatabase.net/k072316/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Percutaneous (DQY)
Date received	Aug 17, 2007
Decision date	Sep 17, 2007
Days to decision	31 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Asahi Intecc Co., Ltd.
Location	Seto-Shi, JP
Contact	YOSHI TERAJ
Website	https://www.asahi-intecc.com
510(k) history	84 submissions · 84 cleared · 2003-2026

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k072316/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 26, 2026