

K202252 ASAHI Corsair Armet 18 Microcatheter (Model: CSAR060-18N), ASAHI Corsair Armet 18 Microcatheter (Model: CSAR090-18N), ASAHI Corsair Armet 18 Microcatheter (Model: CSAR110-18N), ASAHI Corsair Armet 18 Microcatheter (Model: CSAR135-18N),Sep 8, 2020
29 days to decisionK202252 · Product code: **DQY** · Cardiovascular
Source: <https://www.510kdatabase.net/k202252/>**SUBMISSION DETAILS**

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|-----------------------|------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Special |
| Device classification | Catheter, Percutaneous (DQY) |
| Date received | Aug 10, 2020 |
| Decision date | Sep 8, 2020 |
| Days to decision | 29 days |
| Third-party review | No |
| Combination product | No |
| PCCP authorized | No |
| Summary / Statement | Summary |

APPLICANT

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

REGULATORY CONSULTANT

| | |
|-----------------|-------------------------------|
| Consulting firm | Asahi Intecc USA, Inc. |
| Contact | Cynthia Valenzuela |

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)

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