

K230680 Anthogyr Surgical CassettesJul 11, 2023
120 days to decisionK230680 · Product code: **KCT** · General Hospital
Source: <https://www.510kdatabase.net/k230680/>**SUBMISSION DETAILS**

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|-----------------------|---------------------------------------------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Sterilization Wrap Containers, Trays, Cassettes & Other Accessories (KCT) |
| Date received | Mar 13, 2023 |
| Decision date | Jul 11, 2023 |
| Days to decision | 120 days |
| Third-party review | No |
| Combination product | No |
| PCCP authorized | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|-----------------------------------------|
| Company | Anthogyr |
| Location | Sallanches, FR |
| Contact | Aude Goulet |
| 510(k) history | 10 submissions · 10 cleared · 2004-2024 |

REGULATORY CONSULTANT

| | |
|-----------------|---------------------------|
| Consulting firm | Straumann USA, LLC |
| Contact | Jennifer M. Jackson |

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k230680/> 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 May 26, 2026