

K191689 EleGARD Patient Positioning SystemFeb 13, 2020
233 days to decisionK191689 · Product code: **FOA** · General Hospital
Source: <https://www.510kdatabase.net/k191689/>**SUBMISSION DETAILS**

| | |
|-----------------------|------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Board, Cardiopulmonary (FOA) |
| Date received | Jun 25, 2019 |
| Decision date | Feb 13, 2020 |
| Days to decision | 233 days |
| Third-party review | No |
| Combination product | No |
| PCCP authorized | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---|
| Company | Minnesota Resuscitation Solutions, D.B.D Advancedcpr Solutio |
| Location | Edina, MN, US |
| Contact | Philip Tetzlaff |
| 510(k) history | 1 submissions · 1 cleared · 2020-2020 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k191689/> 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 29, 2026