

K811955 CALIBRATED BREATHING SYSTEM COLLECTIONAug 4, 1981
34 days to decisionK811955 · Product code: **KDQ** · General Hospital
Source: <https://www.510kdatabase.net/k811955/>**SUBMISSION DETAILS**

| | |
|-----------------------|------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Bottle, Collection, Vacuum (KDQ) |
| Date received | Jul 1, 1981 |
| Decision date | Aug 4, 1981 |
| Days to decision | 34 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---------------------------------------|
| Company | Anchor Brush Co., Inc. |
| Location | Mchenry, IL, US |
| 510(k) history | 1 submissions · 1 cleared · 1981-1981 |

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k811955/> 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 July 10, 2026