

K232094 Orchid Safety Release ValveTMOct 3, 2023
82 days to decisionK232094 · Product code: **QOI** · General Hospital
Source: <https://www.510kdatabase.net/k232094/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intravenous Catheter Force-activated Separation Device. (QOI)
Date received	Jul 13, 2023
Decision date	Oct 3, 2023
Days to decision	82 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Linear Health Sciences, LLC
Location	Oklahoma City, OK, US
Contact	Daniel Clark
510(k) history	4 submissions · 4 cleared · 2022-2024

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k232094/> 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