

**K821500 STERILE CATH CARE KIT**Jun 23, 1982  
35 days to decisionK821500 · Product code: **KDD** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k821500/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Kit, Surgical Instrument, Disposable (KDD)
Date received	May 19, 1982
Decision date	Jun 23, 1982
Days to decision	35 days
Third-party review	No

**APPLICANT**

---

Company	<b>Pro-Pak</b>
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1982-1982

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k821500/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated July 9, 2026