

**K802213 WHITE KNIGHT BOVIE CORD/SUCTION TUBE**Oct 10, 1980  
28 days to decisionK802213 · Product code: **KKX** · General Hospital  
Source: <https://www.510kdatabase.net/k802213/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drape, Surgical (KKX)
Date received	Sep 12, 1980
Decision date	Oct 10, 1980
Days to decision	28 days
Third-party review	No

**APPLICANT**

---

Company	<b>G.D. Searle and Co.</b>
Location	Mchenry, IL, US
510(k) history	56 submissions · 56 cleared · 1976-1982

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k802213/> 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 May 22, 2026