

**K792096 2 FR KELMAN CHAMBER MAINTAINER**Nov 16, 1979  
29 days to decisionK792096 · Product code: **FMI** · General HospitalSource: <https://www.510kdatabase.net/k792096/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Oct 18, 1979
Decision date	Nov 16, 1979
Days to decision	29 days
Third-party review	No

**APPLICANT**

---

Company	<b>Heyer Schulte Corp.</b>
Location	Mchenry, IL, US
510(k) history	9 submissions · 9 cleared · 1976-1979

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k792096/> 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 2, 2026