

**K120944 KENDALL SCD 700 SEQUENTIAL COMPRESSION CONTROLLER**May 31, 2012  
63 days to decisionK120944 · Product code: **JOW** · Cardiovascular  
Source: <https://www.510kdatabase.net/k120944/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Sleeve, Limb, Compressible (JOW)
Date received	Mar 29, 2012
Decision date	May 31, 2012
Days to decision	63 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Covidien</b>
Location	North Haven, CT, US
Contact	JENNIFER SULLIVAN
510(k) history	130 submissions · 126 cleared · 2008-2024

Covidien is an Irish-registered global healthcare products company headquartered in North Haven, Connecticut. Now part of Medtronic following a 2015 acquisition, the brand continues to operate as a major medical device manufacturer. Covidien maintains a strong FDA 510(k) regulatory record with cleared devices from total submissions spanning 2008 to 2024. The company specializes in General & Plastic Surgery devices, with a dominant focus on surgical staplers, sutures, and wound closure systems. Recent clearances include advanced stapling technologies, endotracheal tubes, a...

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k120944/> 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](http://accessdata.fda.gov). - Generated June 14, 2026