

K002702 NEUROTOLOGY DRAPE, MODEL TBDOct 26, 2000
57 days to decisionK002702 · Product code: **KKX** · General Hospital
Source: <https://www.510kdatabase.net/k002702/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drape, Surgical (KKX)
Date received	Aug 30, 2000
Decision date	Oct 26, 2000
Days to decision	57 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Gyrus Ent LLC
Location	Bartlett, TN, US
Contact	GREG SREDIN
510(k) history	28 submissions · 28 cleared · 1997-2004

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k002702/> 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 June 30, 2026