

K100382 SALUTUNNEL NERVE PROTECTOR, MODEL ED-20329, ED-20330, ED-20331

Aug 5, 2010
170 days to decision

K100382 · Product code: **JXI** · Neurology
Source: <https://www.510kdatabase.net/k100382/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cuff, Nerve (JXI)
Date received	Feb 16, 2010
Decision date	Aug 5, 2010
Days to decision	170 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Salumedica, LLC
Location	Atlanta, GA, US
Contact	BOB BRADEN
510(k) history	2 submissions · 2 cleared · 2000-2010

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k100382/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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