

K031008 AESCULAP NEURO PATTIES, NEUROSORB 4, 6, AND 75 (GREEN & WHITE)

May 9, 2003
39 days to decision

K031008 · Product code: **HBA** · Neurology
Source: <https://www.510kdatabase.net/k031008/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Neurosurgical Paddie (HBA)
Date received	Mar 31, 2003
Decision date	May 9, 2003
Days to decision	39 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	Aesculap
Location	Center Valley, PA, US
Contact	MATTHEW M HULL
510(k) history	10 submissions · 10 cleared · 2003-2010

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k031008/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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