

K182967 Visor SystemNov 23, 2018
28 days to decisionK182967 · Product code: **QAF** · Neurology
Source: <https://www.510kdatabase.net/k182967/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Impedance Plethysmograph, Cranial (QAF)
Date received	Oct 26, 2018
Decision date	Nov 23, 2018
Days to decision	28 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Cerebrotech Medical Systems, Inc.
Location	Pleasanton, CA, US
Contact	Dawnel Scott
510(k) history	1 submissions · 1 cleared · 2018-2018

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k182967/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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