

K171862 GALAXY G3 Mini Microcoil Delivery SystemSep 25, 2017
95 days to decisionK171862 · Product code: **HCG** · Neurology
Source: <https://www.510kdatabase.net/k171862/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Neurovascular Embolization (HCG)
Date received	Jun 22, 2017
Decision date	Sep 25, 2017
Days to decision	95 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Codman & Shurtleff, Inc.
Location	Mchenry, IL, US
Contact	Yoon Hee Beatty
510(k) history	152 submissions · 151 cleared · 1976-2020

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k171862/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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