

**K172843 TwitchView System**Mar 29, 2018  
191 days to decisionK172843 · Product code: **KOI** · Anesthesiology  
Source: <https://www.510kdatabase.net/k172843/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Peripheral, Electric (KOI)
Date received	Sep 19, 2017
Decision date	Mar 29, 2018
Days to decision	191 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Blink Device Corporation</b>
Location	Seattle, WA, US
Contact	Justin Hulvershorn
510(k) history	1 submissions · 1 cleared · 2018-2018

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k172843/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 24, 2026