FDA 510(k) Premarket Notification

CLEARED

K213824 Sonopet iQ Ultrasonic Aspirator System

Feb 3, 2022
57 days to decision

K213824 · Product code: **LFL** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k213824/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Instrument, Ultrasonic Surgical (LFL)
Date received	Dec 8, 2021
Decision date	Feb 3, 2022
Days to decision	57 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Stryker Corporation**
Location	Malwah, NJ, US
Contact	Thomas Flannelly
Website	http://www.stryker.com/
510(k) history	81 submissions · 81 cleared · 2010-2023

Stryker Corporation is an American multinational medical technology company headquartered in Portage, Michigan. The company develops and markets surgical equipment, neurotechnology, orthopedic implants, and patient safety systems used globally across medical specialties. Stryker has received FDA 510(k) clearances from total submissions between 2010 and 2023. The company's cleared devices span orthopedic surgery, neurosurgery, general and plastic surgery, and ear, nose, and throat specialties. This regulatory record reflects the company's broad portfolio across surgical an...