

**K240657 ArtiSential Laparoscopic Instruments-Electrodes (AMHD01-LH)**

Jun 6, 2024  
90 days to decision

K240657 · Product code: **GEI** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k240657/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Mar 8, 2024
Decision date	Jun 6, 2024
Days to decision	90 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary
Other names	ArtiSential Laparoscopic Instruments-Electrodes (AMHD01S-LH); ArtiSential Laparoscopic Instruments-Electrodes (AMHD01L-LH); ArtiSential Laparoscopic Instruments-Electrodes (5AMHD01-LH); ArtiSential Laparoscopic Instruments-Electrodes (5AMHD01S-LH); ArtiSential Laparoscopic Instruments-Electrodes (AMHD01-FH); ArtiSential Laparoscopic Instruments-Electrodes (AMHD01S-FH); ArtiSential Laparoscopic Instruments-Electrodes (AMHD01L-

**APPLICANT**

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Company	<b>Livsmed, Inc.</b>
Location	Seongnam-Si, KR
Contact	Hyeonbin Chung
510(k) history	12 submissions · 12 cleared · 2020-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k240657/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://accessdata.fda.gov)).

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