

**K090023 IF SERIES TRUE SINE INTERFERENTIAL
STIMULATOR, MODEL: WL-2206B AND WL-2106E**Jan 30, 2009
25 days to decisionK090023 · Product code: **LIH** · Neurology
Source: <https://www.510kdatabase.net/k090023/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Interferential Current Therapy (LIH)
Date received	Jan 5, 2009
Decision date	Jan 30, 2009
Days to decision	25 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Well-Life Healthcare Limited
Location	Kowloon, HK
Contact	JENNY HSIEH
510(k) history	22 submissions · 22 cleared · 2006-2024

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k090023/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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