

K083915 ANGELITE FAMILY OF INTENSE PULSED LIGHT SYSTEMS, MODELS ANGELITE-DDC AND ANGELITE-SDCJan 29, 2009
30 days to decisionK083915 · Product code: **GEX** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k083915/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Dec 30, 2008
Decision date	Jan 29, 2009
Days to decision	30 days
Third-party review	Yes
Summary / Statement	Statement

APPLICANT

Company	Advanced Technology Laser Co., Ltd.
Location	Shanghai, CN
Contact	Diana Hong
510(k) history	5 submissions · 5 cleared · 2009-2014

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k083915/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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