

K011708 MODIFICATION TO WARTNER WART REMOVAL SYSTEM

Feb 20, 2002
261 days to decision

K011708 · Product code: **GEH** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k011708/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Cryosurgical, Accessories (GEH)
Date received	Jun 4, 2001
Decision date	Feb 20, 2002
Days to decision	261 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Wartner Medical Products
Location	Brampton, Ontario, CA
Contact	NANCY LUM-WILSON
510(k) history	3 submissions · 3 cleared · 2000-2003

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k011708/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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