

**K120573 MEDART 620**Jun 13, 2012  
107 days to decisionK120573 · Product code: **GEX** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k120573/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Feb 27, 2012
Decision date	Jun 13, 2012
Days to decision	107 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

Company	<b>Medart A/S</b>
Location	Hvidovre, DK
Contact	DARREN THOMAS
510(k) history	3 submissions · 3 cleared · 2009-2012

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k120573/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 14, 2026