

**K944817 NEUTRALECT ADULT STANDARD, RETURN (REM),  
PAEDIATRIC STANDARD, RETURN (REM)**Nov 9, 1994  
41 days to decisionK944817 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k944817/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Sep 29, 1994
Decision date	Nov 9, 1994
Days to decision	41 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Msb , Ltd.</b>
Location	Marlborough, Wiltshire, GB
Contact	MARTIN BEAUMONT
510(k) history	7 submissions · 7 cleared · 1994-1998

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k944817/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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