

Device Return

- Devices are returned to the Sponsor for several reasons which include but are not limited to:
 - Product expiration
 - o Device malfunction
 - Shipping/ordering error
 - Inventory return/exchange

1. What Do I Do if I Need to Return a Device/Devices?

- All returned devices must have an RGA (Returned Goods Authorization) number. Please ask your **Field Clinical Specialist** to create one on case day or reach out to your **Site Manager.**
 - Note: product may be returned for many reasons (device deficiency, expiration, site transfer, etc.).

2. Where Do I Record the RGA Number?

- In the appropriate column of the Device Accountability Application (screenshot below)
- AND somewhere visible on the packaging of the return devices

Unique ID Site Id	Product Code Description Order#	Query/ Monitor Status	Lot# Expy Date	Subject ID	Disposition Date	Deficient	Used	Disposed	Returned	RGA	Transferred	Other	Search	× C
TEST2	TESTPRODUCT - TESTPRODUCT - Used for development 0-240125074143		TST-101 5/15/2024		12/31/2001				~	RGA				*
TEST2	TESTPRODUCT - TESTPRODUCT - Used for development 0-240117155550		TST-101 5/15/2024		12/31/2001									
TEST2	TESTPRODUCT - TESTPRODUCT - Used for development 0-240103112713		TST-101 5/15/2024		12/31/2001									
TEST2	TESTPRODUCT - TESTPRODUCT - Used for development 0-231121085818		TST-101 5/15/2024		12/31/2001									
TEST2	TESTPRODUCT - TESTPRODUCT - Used for development 0-231120074825		TST-101 5/15/2024		12/31/2001									
TECTO	TESTPRODUCT - TESTPRODUCT - Used for development 0-230809184819	~	TST-101 5/15/2024		12/31/2001									
TECTO	TESTPRODUCT - TESTPRODUCT - Used for development 0-230809184819	~	TST-101 5/15/2024		12/31/2001									
TEST2	TESTPRODUCT2 - TESTPRODUCT2 - Used for development		TST-101 5/15/2023		12/31/2001									•
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3. How do I get a return shipping label?

- Use return labels included with original product shipment OR
- Reach out to the Site Manager or Field Clinical Specialist
 - Site Manager or FCS will generate the return shipping label with FedEx and send it to the site contact via email
- Print the shipping label and place it on the original device packaging
 - Write RGA number somewhere visible on the device packaging

4. What Do I Do if There was a Device Deficiency?

- Check off the appropriate box on the Device Accountability Log
 - \circ $\;$ Note: not all product with a device deficiency needs to be returned.



The Shape of Stroke Prevention

• Record the device deficiency in EDC

Unique ID Site Id	Product Code Description Order#	Query/ Monitor Status	Lot# Expy Date	Subject ID	Disposition Date	Deficient	Used	Disposed	Returned	RGA	Transferred	Other	Search	XQ
#3833 TEST2	TESTPRODUCT - TESTPRODUCT - Used for development 0-240125074143		TST-101 5/15/2024		12/31/2001				v	RGA				*
#3761 TEST2	TESTPRODUCT - TESTPRODUCT - Used for development 0-240117155550		TST-101 5/15/2024		12/31/2001									
#3687 TEST2	TESTPRODUCT - TESTPRODUCT - Used for development 0-240103112713		TST-101 5/15/2024		12/31/2001									
#3538 TEST2	TESTPRODUCT - TESTPRODUCT - Used for development O-231121085818		TST-101 5/15/2024		12/31/2001									
#3530 TEST2	TESTPRODUCT - TESTPRODUCT - Used for development O-231120074825		TST-101 5/15/2024		12/31/2001									
#2873 TEST2	TESTPRODUCT - TESTPRODUCT - Used for development O-230809184819	~	TST-101 5/15/2024		12/31/2001									
#2874 TEST2	TESTPRODUCT - TESTPRODUCT - Used for development O-230809184819	~	TST-101 5/15/2024		12/31/2001									
#2936 TEST2	TESTPRODUCT2 - TESTPRODUCT2 - Used for development		TST-101 5/15/2023		12/31/2001									•
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accept that your electronic signature is the legally binding equivalent of your