July 1, 2024

Innovative Health, LLC.
Amanda Babcock
Regulatory Affairs Manager
1435 North Hayden Road
Suite 100
Scottsdale, Arizona 85257

Re: K231015
Trade/Device Name: Reprocessed Cordis Super Torque and Super Torque Plus Diagnostic Angiographic Catheters
Regulation Number: 21 CFR 870.1200
Regulation Name: Diagnostic Intravascular Catheter
Regulatory Class: Class II
Product Code: NLI
Dated: April 7, 2023
Received: April 10, 2023

Dear Amanda Babcock:

We have reviewed your section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (the Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. Although this letter refers to your product as a device, please be aware that some cleared products may instead be combination products. The 510(k) Premarket Notification Database available at https://www.accessdata.fda.gov/scripts/cdrh/cfdocs/cfpmn/pmn.cfm identifies combination product submissions. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration. Please note: CDRH does not evaluate information related to contract liability warranties. We remind you, however, that device labeling must be truthful and not misleading.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Additional information about changes that may require a new premarket notification are provided in the FDA guidance documents entitled "Deciding When to Submit a 510(k) for a Change to an Existing Device" (https://www.fda.gov/media/99812/download) and "Deciding When to Submit a 510(k) for a Software Change to an Existing Device" (https://www.fda.gov/media/99785/download).

Your device is also subject to, among other requirements, the Quality System (QS) regulation (21 CFR Part 820), which includes, but is not limited to, 21 CFR 820.30, Design controls; 21 CFR 820.90, Nonconforming product; and 21 CFR 820.100, Corrective and preventive action. Please note that regardless of whether a change requires premarket review, the QS regulation requires device manufacturers to review and approve changes to device design and production (21 CFR 820.30 and 21 CFR 820.70) and document changes and approvals in the device master record (21 CFR 820.181).

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); medical device reporting (reporting of medical device-related adverse events) (21 CFR Part 803) for devices or postmarketing safety reporting (21 CFR Part 4, Subpart B) for combination products (see https://www.fda.gov/combination-products/guidance-regulatory-information/postmarketing-safety-reporting-combination-products); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820) for devices or current good manufacturing practices (21 CFR Part 4, Subpart A) for combination products; and, if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR Parts 1000-1050.

Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21 CFR 807.97). For questions regarding the reporting of adverse events under the MDR regulation (21 CFR Part 803), please go to https://www.fda.gov/medical-devices/medical-device-safety/medical-device-reporting-mdr-how-report-medical-device-problems.

For comprehensive regulatory information about medical devices and radiation-emitting products, including information about labeling regulations, please see Device Advice (https://www.fda.gov/medical-devices/device-advice-comprehensive-regulatory-assistance) and CDRH Learn (https://www.fda.gov/training-and-continuing-education/cdrh-learn). Additionally, you may contact the Division of Industry and Consumer Education (DICE) to ask a question about a specific regulatory topic. See the DICE website (https://www.fda.gov/medical-devices/device-advice-comprehensive-regulatory-assistance/contact-us-division-industry-and-consumer-education-dice) for more information or contact DICE by email (DICE@fda.hhs.gov) or phone (1-800-638-2041 or 301-796-7100).

Sincerely,

Ariel G. Ash-shakoor -S

Digitally signed by Ariel G. Ash-shakoor -S
Date: 2024.07.01 09:21:23 -04'00'

For

Gregory O'Connell

Assistant Director
DHT2C: Division of Coronary
and Peripheral Intervention Devices
OHT2: Office of Cardiovascular Devices
Office of Product Evaluation and Quality
Center for Devices and Radiological Health

Enclosure

DEPARTMENT OF HEALTH AND HUMAN SERVICES
Food and Drug Administration

Indications for Use

Form Approved: OMB No. 0910-0120
Expiration Date: 07/31/2026
See PRA Statement below.

510(k) Number *(if known)*
K231015

Device Name
Reprocessed Cordis Super Torque and Super Torque Plus Angiographic Diagnostic Catheters

Indications for Use *(Describe)*
Reprocessed Cordis Catheters are designed to deliver radiopaque contrast medium to selected sites in the vascular system.

Type of Use *(Select one or both, as applicable)*

☒ Prescription Use (Part 21 CFR 801 Subpart D) ☐ Over-The-Counter Use (21 CFR 801 Subpart C)

CONTINUE ON A SEPARATE PAGE IF NEEDED.

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510(k) SUMMARY

Submitter's Name and Address:
Innovative Health, LLC.
1435 N. Hayden Road, Suite 100
Scottsdale, AZ 85257

Contact Name and Information:
Meerna Muradvich
Regulatory Affairs Engineer
Innovative Health, LLC.
(480) 692-7176 (office)
(888) 965-7706 (fax)
mmuradvich@innovative-health.com

Amanda Babcock
Regulatory Affairs Manager
Innovative Health, LLC.
(480) 525-5911 (office)
(888) 965-7706 (fax)
ababcock@innovative-health.com

Date prepared:
06/26/2024

Device Information:

Trade/Proprietary Name:	Reprocessed Cordis Super Torque and Super Torque Plus Diagnostic Angiographic Catheters
Common Name:	Angiographic Catheter
Classification Name:	Catheter, Angiography, Reprocessed
Classification Number:	Class II, 21 CFR 870.1200
Product Code:	NLI

Predicate Device:

510(k) Number	Device	Manufacturer
K915836	Cordis 5.0 French Super Torque Catheters	Cordis Corp.
K914007	Cordis 5.2 French Super Torque Plus	Cordis Corp.

Device Description:

The reprocessed Cordis Catheters are available in a broad variety of French sizes and configurations. These catheters combine an atraumatic tip with a braided body.

The item numbers included in the scope of this submission are as follows:

Description	Item Number	French Size	Length	Side holes	Curve
	532-410T	4	65	8	PIG
	532-420T	4	65	8	straight
	532-462	4	65	2	RDCS

Description	Item Number	French Size	Length	Side holes	Curve
Reprocessed Cordis Super Torque Diagnostic Angiographic Catheter	532-444	4	65	0	C3
	532-445	4	65	2	C3
	532-473	4	65	0	SHK 1
	532-474	4	65	0	SHK 2
	532-447	4	65	0	JC 1
	532-448	4	65	0	JC 2
	532-449	4	65	0	JC 3
	SR3740	4	80	0	JL 1.5
	SRD6054	4	80	0	JL 2
	SR4533	4	80	0	JL 2.5
	SRD6920	4	80	4	PIG 145°
	532-432	4	80	2	multipurpose small
	532-478	4	80	2	RDCS
	532-411T	4	90	8	PIG
	532-421T	4	90	8	straight
	SRD5816	4	100	0	IM-MOD-5
	532-412T	4	100	8	PIG
	532-422T	4	100	8	straight
	532-430	4	100	0	multipurpose small
	532-482	4	100	2	multipurpose small
	532-414	4	100	0	SIM 1
	532-415	4	100	0	SIM 2
	532-461	4	100	0	H1
	532-470	4	100	0	MAN
	532-405	4	100	0	BERN
	532-436	4	100	0	JB1
	532-437	4	100	0	JB2
	532-438	4	100	0	JB3
	532-497	4	100	0	VERT
	532-413T	4	110	8	PIG
	532-524V8	5	65	8	PIG
	532-564V8	5	65	8	straight
	532-576	5	65	0	multipurpose (subintimal recanalization)
	532-510	5	65	2	RDCS
	532-514	5	65	0	C3
	532-517	5	65	2	C3
	532-519	5	65	0	SHK 1
	532-522	5	65	0	JC 2
	532-523	5	65	0	JC 3
	532-579	5	80	0	MPA

Description	Item Number	French Size	Length	Side holes	Curve
Reprocessed Cordis Super Torque Diagnostic Angiographic Catheter	532-511	5	80	2	RDCS
	532-509	5	80	0	RDCA
	532-567	5	80	0	SHK 2
	532-539F8	5	90	8	PIG
	532-555H8	5	100	8	PIG
	532-569H8	5	100	8	straight
	532-578	5	100	0	MPA
	532-501	5	100	0	SIM 1
	532-546	5	100	2	SIM 1
	532-502	5	100	0	SIM 2
	532-547	5	100	2	SIM 2
	532-503	5	100	0	SIM 3
	532-504	5	100	0	H1
	532-520	5	100	0	HN4
	532-571	5	100	0	MAN
	532-541H0	5	100	0	JB1
	532-543H0	5	100	0	JB2
	532-549H0	5	100	0	VERT
	455-610T	6	65	12	PIG
	455-672	6	65	0	C3
	455-689	6	80	2	RDCA
	455-698	6	80	4	NIH
	532-506	6	80	2	multipurpose small
	455-623	6	80	0	MPA
	455-636	6	80	2	MPA
	455-686	6	80	0	RDCA
	455-611T	6	90	12	PIG
	532-618	6	100	0	JL 3.5
	532-620	6	100	0	JL 4
	532-627	6	100	0	JL 4.5
	532-622	6	100	0	JL 5
	532-624	6	100	0	JL 6
	532-645	6	100	0	AL 1
	532-646	6	100	0	AL 2
	532-647	6	100	0	AL 3
	532-619	6	100	0	JR 3.5
	532-621	6	100	0	JR 4
	532-623	6	100	0	JR 5
	532-625	6	100	0	JR 6
	532-648	6	100	0	AR MOD
	532-642	6	100	2	MPA 2

Description	Item Number	French Size	Length	Side holes	Curve
Reprocessed Cordis Super Torque Diagnostic Angiographic Catheter	532-649	6	100	2	MPB 2
	532-640	6	100	0	SON 1
	532-664	6	100	4	SON 2
	532-672	6	100	0	LCB
	532-670	6	100	0	RCB
	532-660	6	100	0	IM
	455-637	6	100	2	MPA
	455-660	6	100	0	SIM 1
	455-660D	6	100	2	SIM 1
	455-661	6	100	0	SIM 2
	455-662	6	100	0	SIM 3
	455-663	6	100	0	SIM 4
	455-665	6	100	0	H1
	455-666	6	100	0	H1
	532-650S	6	110	6	PIG
	532-652S	6	110	6	PIG 145°
	532-654S	6	110	6	PIG 145° MOD
	455-613E	6	110	8	PIG
Reprocessed Cordis Super Torque Plus Diagnostic Angiographic Catheter	533-562	5.2	65	2	MPA 2
	533-579	5.2	80	0	MPA
	SR3216	5.2	100	0	JL 3
	533-551	5.2	100	0	JL 3.5
	533-553	5.2	100	0	JL 4
	533-527	5.2	100	0	JL 4.5
	533-559	5.2	100	0	JL 5
	533-561	5.2	100	0	JL 6
	533-542	5.2	100	0	AL 2
	533-544	5.2	100	0	AL 3
	SR4615	5.2	100	0	XB 3.5 ST
	533-550	5.2	100	0	JR 3.5
	533-552	5.2	100	0	JR 4
	533-528	5.2	100	0	JR 4 MOD
	533-563	5.2	100	0	JR 4 ST
	533-558	5.2	100	0	JR 5
	533-560	5.2	100	0	JR 6
	SR1924	5.2	100	0	MPA
	533-582	5.2	100	2	MPA 2
	533-584	5.2	100	0	CAS 1
	533-585	5.2	100	0	CAS 2
	599500J2	5.2	100	2	RBL-JK
	599500T1	5.2	100	1	RBL-TG

Description	Item Number	French Size	Length	Side holes	Curve
Reprocessed Cordis Super Torque Plus Diagnostic Angiographic Catheter	533-572	5.2	100	0	LCB
	533-570	5.2	100	0	RCB
	533-580	5.2	100	0	IM
	SR4658	5.2	100	0	IM-MOD
	SR4268	5.2	100	0	IM-MOD-2
	SR4685	5.2	100	0	IM-MOD-3
	533-533	5.2	110	6	PIG
	533-534A	5.2	110	6	PIG 145°
	533-537S	5.2	110	6	PIG 145° MOD
	533-533A	5.2	110	6	PIG 155°
	533-633	6	80	0	MPA 1
	533-629	6	80	2	MPA 2
	533-682	6	80	4	CAS 2
	533-635	6	80	0	NIH
	533-618	6	100	0	JL 3.5
	533-620	6	100	0	JL 4
	533-627	6	100	0	JL 4.5
	533-622	6	100	0	JL 5
	533-624	6	100	0	JL 6
	533-645	6	100	0	AL 1
	533-646	6	100	0	AL 2
	533-647	6	100	0	AL 3
	533-619	6	100	0	JR 3.5
	533-621	6	100	2	JR 3.5
	533-687	6	100	0	JR 4 CLASSIC
	533-689	6	100	0	JR 4 RECESSED BRAIDING
	533-628	6	100	0	JR 4 MOD
	533-626	6	100	0	JR 4 ST
	533-623	6	100	0	JR 5
	533-625	6	100	0	JR 6
	533-648	6	100	0	AR MOD
	533-641	6	100	0	AR 1 MOD
	533-643	6	100	0	AR 2 MOD
	533-640	6	100	0	MPA 1 COURNAD
	533-642	6	100	2	MPA 2
	533-649	6	100	2	MPB 2
	533-634	6	100	6	MPB 3
	533-630	6	100	4	SON 1
	533-631	6	100	0	SON 2
	533-632	6	100	4	SON 3

Description	Item Number	French Size	Length	Side holes	Curve
Reprocessed Cordis Super Torque Plus Diagnostic Angiographic Catheter	533-684	6	100	0	CAS 1
	533-685	6	100	0	CAS 2
	533-686	6	100	0	CAS 3
	599600R401	6	100	1	RBL 4.0
	599600R451	6	100	1	RBL 4.5
	599600R501	6	100	1	RBL 5.0
	599600J2	6	100	2	RBL-JK
	599600T1	6	100	1	RBL-TG
	533-637	6	100	0	EGB 1
	533-638	6	100	0	EGB 2
	533-672	6	100	0	LCB
	533-670	6	100	0	RCB
	533-660	6	100	0	IM
	533-636	6	100	0	NIH
	533-650F	6	110	4	PIG
	533-650S	6	110	6	PIG
	533-650E	6	110	8	PIG
	533-652S	6	110	6	PIG 145°
	533-653S	6	110	6	PIG 145° MOD
	533-654S	6	110	6	PIG 155°
	533-655S	6	110	6	PIG 155° MOD

Table 1: Device Scope

Indications for Use:
Reprocessed Cordis Catheters are designed to deliver radiopaque contrast medium to selected sites in the vascular system.

Technological Characteristics:
The purpose, design, materials, function, and intended use of the Reprocessed Cordis Super Torque and Super Torque Plus diagnostic angiographic catheters are identical to the predicate devices. There are no changes to the claims, clinical applications, patient population, performance specifications, or method of operation. In addition, Innovative Health's reprocessing of these devices includes removal of visible soil and decontamination. Each device is inspected and function tested prior to packaging and labeling.

Functional and Safety Testing:
Bench and laboratory testing was conducted to demonstrate substantially equivalent performance of the Reprocessed Cordis Super Torque and Super Torque Plus Catheters. This included the following:

- Biocompatibility
    - Cytotoxicity
    - Hemocompatibility
    - Sensitization
    - Irritation
    - Systemic Toxicity
- Cleaning Validation
- Sterilization Validation
- Functional testing
    - Visual Inspection
    - Dimensional Verification
    - Radiopacity Testing
    - Simulated Use Testing
    - Leak Testing
    - Torsion Testing
    - Tip Buckling Testing
    - Tensile Testing
    - Coating Coverage
    - Particulate Testing
- Packaging Validation

The Reprocessed Cordis Super Torque and Super Torque Plus Catheters are reprocessed no more than one (1) time. Each device is marked, serialized and tracked. After the device has reached the maximum number of reprocessing cycles, the device is rejected from further reprocessing. Reprocessing is performed only by Innovative Health. Innovative Health restricts its reprocessing to exclude devices previously reprocessed by other reprocessors.

Conclusion:
Innovative Health concludes that the Reprocessed Cordis Super Torque and Super Torque Plus Diagnostic Angiographic catheters do not raise new questions of safety and effectiveness and supports substantial equivalence to the predicate devices described herein.

The item numbers included in the scope of this submission are as follows:

Description	Item Number	French Size	Length	Side holes	Curve
Reprocessed Cordis Super Torque Diagnostic Angiographic Catheter	532-410T	4	65	8	PIG
	532-420T	4	65	8	straight
	532-462	4	65	2	RDCS
	532-444	4	65	0	C3
	532-445	4	65	2	C3
	532-473	4	65	0	SHK 1
	532-474	4	65	0	SHK 2
	532-447	4	65	0	JC 1
	532-448	4	65	0	JC 2
	532-449	4	65	0	JC 3
	SR3740	4	80	0	JL 1.5
	SRD6054	4	80	0	JL 2
	SR4533	4	80	0	JL 2.5
	SRD6920	4	80	4	PIG 145°
	532-432	4	80	2	multipurpose small
	532-478	4	80	2	RDCS
	532-411T	4	90	8	PIG
	532-421T	4	90	8	straight
	SRD5816	4	100	0	IM-MOD-5
	532-412T	4	100	8	PIG
	532-422T	4	100	8	straight
	532-430	4	100	0	multipurpose small
	532-482	4	100	2	multipurpose small
	532-414	4	100	0	SIM 1
	532-415	4	100	0	SIM 2
	532-461	4	100	0	H1
	532-470	4	100	0	MAN
	532-405	4	100	0	BERN
	532-436	4	100	0	JB1
	532-437	4	100	0	JB2
	532-438	4	100	0	JB3
	532-497	4	100	0	VERT
	532-413T	4	110	8	PIG
	532-524V8	5	65	8	PIG
	532-564V8	5	65	8	straight
	532-576	5	65	0	multipurpose (subintimal recanalization)
	532-510	5	65	2	RDCS

Description	Item Number	French Size	Length	Side holes	Curve
Reprocessed Cordis Super Torque Diagnostic Angiographic Catheter	532-514	5	65	0	C3
	532-517	5	65	2	C3
	532-519	5	65	0	SHK 1
	532-522	5	65	0	JC 2
	532-523	5	65	0	JC 3
	532-579	5	80	0	MPA
	532-511	5	80	2	RDCS
	532-509	5	80	0	RDCA
	532-567	5	80	0	SHK 2
	532-539F8	5	90	8	PIG
	532-555H8	5	100	8	PIG
	532-569H8	5	100	8	straight
	532-578	5	100	0	MPA
	532-501	5	100	0	SIM 1
	532-546	5	100	2	SIM 1
	532-502	5	100	0	SIM 2
	532-547	5	100	2	SIM 2
	532-503	5	100	0	SIM 3
	532-504	5	100	0	H1
	532-520	5	100	0	HN4
	532-571	5	100	0	MAN
	532-541H0	5	100	0	JB1
	532-543H0	5	100	0	JB2
	532-549H0	5	100	0	VERT
	455-610T	6	65	12	PIG
	455-672	6	65	0	C3
	455-689	6	80	2	RDCA
	455-698	6	80	4	NIH
	532-506	6	80	2	multipurpose small
	455-623	6	80	0	MPA
	455-636	6	80	2	MPA
	455-686	6	80	0	RDCA
	455-611T	6	90	12	PIG
	532-618	6	100	0	JL 3.5
	532-620	6	100	0	JL 4
	532-627	6	100	0	JL 4.5
	532-622	6	100	0	JL 5
	532-624	6	100	0	JL 6
	532-645	6	100	0	AL 1
	532-646	6	100	0	AL 2

Description	Item Number	French Size	Length	Side holes	Curve
Reprocessed Cordis Super Torque Diagnostic Angiographic Catheter	532-647	6	100	0	AL 3
	532-619	6	100	0	JR 3.5
	532-621	6	100	0	JR 4
	532-623	6	100	0	JR 5
	532-625	6	100	0	JR 6
	532-648	6	100	0	AR MOD
	532-642	6	100	2	MPA 2
	532-649	6	100	2	MPB 2
	532-640	6	100	0	SON 1
	532-664	6	100	4	SON 2
	532-672	6	100	0	LCB
	532-670	6	100	0	RCB
	532-660	6	100	0	IM
	455-637	6	100	2	MPA
	455-660	6	100	0	SIM 1
	455-660D	6	100	2	SIM 1
	455-661	6	100	0	SIM 2
	455-662	6	100	0	SIM 3
	455-663	6	100	0	SIM 4
	455-665	6	100	0	H1
	455-666	6	100	0	H1
	532-650S	6	110	6	PIG
	532-652S	6	110	6	PIG 145°
	532-654S	6	110	6	PIG 145° MOD
	455-613E	6	110	8	PIG
Reprocessed Cordis Super Torque Plus Diagnostic Angiographic Catheter	533-562	5.2	65	2	MPA 2
	533-579	5.2	80	0	MPA
	SR3216	5.2	100	0	JL 3
	533-551	5.2	100	0	JL 3.5
	533-553	5.2	100	0	JL 4
	533-527	5.2	100	0	JL 4.5
	533-559	5.2	100	0	JL 5
	533-561	5.2	100	0	JL 6
	533-542	5.2	100	0	AL 2
	533-544	5.2	100	0	AL 3
	SR4615	5.2	100	0	XB 3.5 ST
	533-550	5.2	100	0	JR 3.5
	533-552	5.2	100	0	JR 4
	533-528	5.2	100	0	JR 4 MOD
	533-563	5.2	100	0	JR 4 ST

Description	Item Number	French Size	Length	Side holes	Curve
Reprocessed Cordis Super Torque Plus Diagnostic Angiographic Catheter	533-558	5.2	100	0	JR 5
	533-560	5.2	100	0	JR 6
	SR1924	5.2	100	0	MPA
	533-582	5.2	100	2	MPA 2
	533-584	5.2	100	0	CAS 1
	533-585	5.2	100	0	CAS 2
	599500J2	5.2	100	2	RBL-JK
	599500T1	5.2	100	1	RBL-TG
	533-572	5.2	100	0	LCB
	533-570	5.2	100	0	RCB
	533-580	5.2	100	0	IM
	SR4658	5.2	100	0	IM-MOD
	SR4268	5.2	100	0	IM-MOD-2
	SR4685	5.2	100	0	IM-MOD-3
	533-533	5.2	110	6	PIG
	533-534A	5.2	110	6	PIG 145°
	533-537S	5.2	110	6	PIG 145° MOD
	533-533A	5.2	110	6	PIG 155°
	533-633	6	80	0	MPA 1
	533-629	6	80	2	MPA 2
	533-682	6	80	4	CAS 2
	533-635	6	80	0	NIH
	533-618	6	100	0	JL 3.5
	533-620	6	100	0	JL 4
	533-627	6	100	0	JL 4.5
	533-622	6	100	0	JL 5
	533-624	6	100	0	JL 6
	533-645	6	100	0	AL 1
	533-646	6	100	0	AL 2
	533-647	6	100	0	AL 3
	533-619	6	100	0	JR 3.5
	533-621	6	100	2	JR 3.5
	533-687	6	100	0	JR 4 CLASSIC
	533-689	6	100	0	JR 4 RECESSED BRAIDING
	533-628	6	100	0	JR 4 MOD
	533-626	6	100	0	JR 4 ST
	533-623	6	100	0	JR 5
	533-625	6	100	0	JR 6
	533-648	6	100	0	AR MOD

Description	Item Number	French Size	Length	Side holes	Curve
Reprocessed Cordis Super Torque Plus Diagnostic Angiographic Catheter	533-641	6	100	0	AR 1 MOD
	533-643	6	100	0	AR 2 MOD
	533-640	6	100	0	MPA 1 COURNAD
	533-642	6	100	2	MPA 2
	533-649	6	100	2	MPB 2
	533-634	6	100	6	MPB 3
	533-630	6	100	4	SON 1
	533-631	6	100	0	SON 2
	533-632	6	100	4	SON 3
	533-684	6	100	0	CAS 1
	533-685	6	100	0	CAS 2
	533-686	6	100	0	CAS 3
	599600R401	6	100	1	RBL 4.0
	599600R451	6	100	1	RBL 4.5
	599600R501	6	100	1	RBL 5.0
	599600J2	6	100	2	RBL-JK
	599600T1	6	100	1	RBL-TG
	533-637	6	100	0	EGB 1
	533-638	6	100	0	EGB 2
	533-672	6	100	0	LCB
	533-670	6	100	0	RCB
	533-660	6	100	0	IM
	533-636	6	100	0	NIH
	533-650F	6	110	4	PIG
	533-650S	6	110	6	PIG
	533-650E	6	110	8	PIG
	533-652S	6	110	6	PIG 145°
	533-653S	6	110	6	PIG 145° MOD
	533-654S	6	110	6	PIG 155°
	533-655S	6	110	6	PIG 155° MOD

Table 1: Device Scope