SELUTION SLR™ DEB

DRUCELUTION DONE BETTER

SUMMARY OF

SELUTION4ISR TRIAL'

A prospective, multi-center, randomized, single blind, controlled, non-inferiority clinical trial to demonstrate safety and efficacy of SELUTION SLR DEB vs. Standard of Care (SOC) in ISR.

418
PATIENTS

48 SITES

SELUTION SLR DEB

(N=210)



US Canada Brazil Europe

STANDARD OF CARE

(N=208)

80% DES **20**% POBA

365

STUDY RESULTS -

SELUTION SLR™ DEB IS AN ALTERNATIVE TO STANDARD OF CARE IN ISR, MINIMIZING METAL.



NON-INFERIOR: SELUTION SLR DEB VS SOC*

SOC 13.5%

180 270

Time Post-Procedure (days)

15.2%

TARGET LESION FAILURE SELUTION SLR DEB

VS

13.5%

TARGET LESION FAILURE STANDARD OF CARE

90

50

40

30

20

10

0

0

Cumulative TLF Rate (%, ITT)



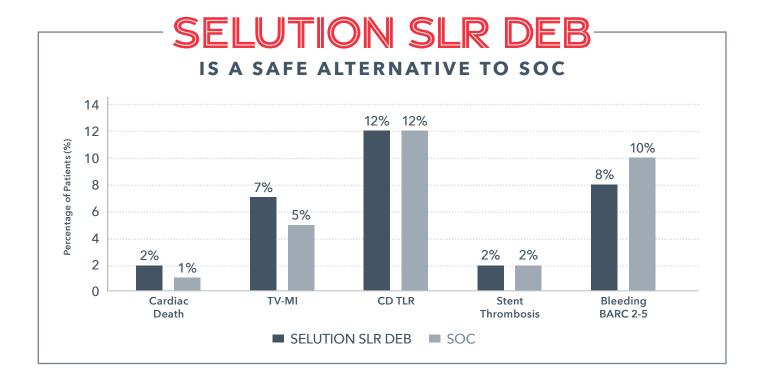
^{1.} SELUTION4ISR Trial, Oral Presentation, TCT 2025

^{*}Per protocol results (16.2% vs. 14.5%)

SELUTION SLR DEB

IS AN ALTERNATIVE TO SOC ACROSS ALL SUBGROUPS

SUBGROUP ANALYSIS OF THE PRIMARY ENDPOINT (ITT) Group Subgroup DEB SOC **Relative Risk Difference** RR 95% CI Interaction N=210 N=208 p-value 7/44 (16) >75 10/60 (17) 0.95 (0.39 - 2.31) Age 0.632 <= 75 25/166 (15) 18/148 (12) 1.24 (0.7 - 2.18) 6/39 (15) 7/55 (13) 1.21 (0.44 - 3.32) Female 0.882 Male 26/171 (15) 21/153 (14) 1.11 (0.65 - 1.89) Medical 11/119 (9) 0.9 (0.4 - 2.04) No Diabetes 10/120 (8) History of Diabetes 0.464 Diabetes 22/90 (24) 17/89 (19) 1.28 (0.73 - 2.24) Reference 1.67 (0.69 - 4.04) > 2.75 11/67 (16) 7/71 (10) 0.302 Vessel 0.96 (0.55 - 1.67) 21/143 (15) 21/137 (15) Diameter <= 2.75< 10mm 14/93 (15) 17/99 (17) 0.88 (0.46 - 1.68) 0.438 Lesion Length 1.56 (0.69 - 3.53) 10 - 20mm 14/92 (15) 8/82 (10) > 20 mm 4/24 (17) 2/24 (8) 2 (0.4 - 9.91) One 22/167 (13) 16/168 (10) 1.38 (0.75 - 2.54) Number of 0.242 10/43 (23) 12/40 (30) 0.78 (0.38 - 1.59) Two .001 10 100 **FAVORS SELUTION SLR DEB FAVORS SOC**



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For Healthcare Professionals Only.

Important information: Prior to use, refer to the instructions for use supplied with this device for indications, contraindications, suggested procedure, warnings and precautions. As part of its continuous product development policy, Cordis reserves the right to change product specifications without prior notification. Please contact your Cordis representative for additional product availability information. SELUTION SLR Drug-Eluting Balloon is manufactured by MedAlliance, LLC and its affiliates. SELUTION SLR is a trademark of M.A. Med Alliance SA. MedAlliance, LLC and M.A. Med Alliance SA are Cordis companies. CORDIS and Cordis LOGO are Trademarks of Cordis and may be registered in the US and/or in other countries. All other marks are the property of their respective owners.



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