MeRes-1 Extend Study

Study Highlights

- Principal Investigator: Dr. Alexandre Abizaid
- The MeRes-1 Extend was a prospective, multicenter, single-arm, trial of MeRes100 sirolimus-eluting bioresorbable vascular scaffold system in patients with de novo native coronary artery lesions
- Three-year clinical follow-up including
 - · QCA analysis at 6 and 36 months
 - OCT analysis at 6 and 36 months
- MeRes-1 Extend study demonstrated the favourable safety and efficacy of MeRes100 BRS at 24 months follow-up



Study Design

A multicentre, single-arm, prospective study



A total of 64 patients were enrolled in Spain, FYR of Macedonia, Brazil, South Africa, Malaysia and Indonesia



Clinical follow-up at 1 month, 6 months, 12 months, 24 months and 36 months post-procedure



Angiographic follow-up at 6 and 36 months Analysed by Cardiovascular Research Centre, Sao Paulo, Brazil



OCT follow-up at 6 and 36 months Analysed by Cardialysis BV, Rotterdam, the Netherlands

Clinical Presentation



Figure 1: Cardiac Status



Figure 2: Lesion Characteristics (ACC/AHA Classification)

Study Results



Figure 3: Late lumen loss at 6-month follow-up

References

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- 5. Presented by Alexandre Abizaid. Implantation of Thin Strut Sirolimus-Eluting Bioresorbable Vascular Scaffold in Patients With de novo Coronary Artery Lesions: Two-year Clinical and Six-month Imaging Outcomes of the MeRes-1 Extend Trial. Accepted in abstract book of Euro Intervention 2019.
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