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Procedural and early clinical outcome of Xience Prime LL Everolimus-eluting 38mm-long stent

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Purpose: The SPRIT PRIME trail demonstrated excellent deliverability and clinical outcome of Xience Prime LL Everolimus—eluting 38mm—long stent (XP—38), the longest drug—eluting stent available in Japan. However, it is unclear in the real—world setting. To address the issue, we evaluated deliverability and early clinical outcome of XP—38 implantation in our hospital. Methods: From May 2012 to June 2012, consecutive 23 patients with 24 lesions treated by at least one XP—38 were enrolled. The procedural and 30—days clinical outcome was retrospectively evaluated. Results: The mean of age was 63±12 years old; 17.4% of female and 21.7% of diabetes. Of 24 lesions, all lesions were type C lesion with more than 20mm of lesion length. 26.1% was severe calcified, and 4.3% was chronic total occlusion. The lesion was located in the LMT in 8.7%, LAD in 52.2%, LCX in 8.7%, and RCA in 30.4%. The number of XP38 per patient was 1.3±0.6. Total number of used XP38 was 30. Stent implantation was successfully performed in all cases; one patient required "two—wire technique", and two did "mother—child catheter technique". During 30—days after stent implantation, none had cardiac death, myocardial infarction, target vessel failure and stent thrombosis. Conclusion: Xience Prime LL Everolimus—eluting 38mm—long stent showed favorable deliverability and early clinical outcome in the real—world setting. Further studies are needed to clarify long—term clinical outcome.