

5 year clinical outcomes of a polymerfree sirolimus-eluting stent versus a polymer-based paclitaxel-eluting stent: The ISAR-TEST trial

Lamin A King, Julinda Mehilli, Stefanie Schulz, Robert A Byrne, Albert Schömig, Adnan Kastrati and

Intracoronary Stenting and Angiographic
Restenosis – Test Equivalence Between Two DrugEluting Stents (ISAR-TEST) Trial Investigators

Deutsches Herzzentrum, Technische Universität, Munich, Germany











No potential conflicts of interest

Speaker's name: Dr Lamin King

☐ I do not have any potential conflict of interest











ISAR-TEST: TRIAL DESIGN

DESIGN:

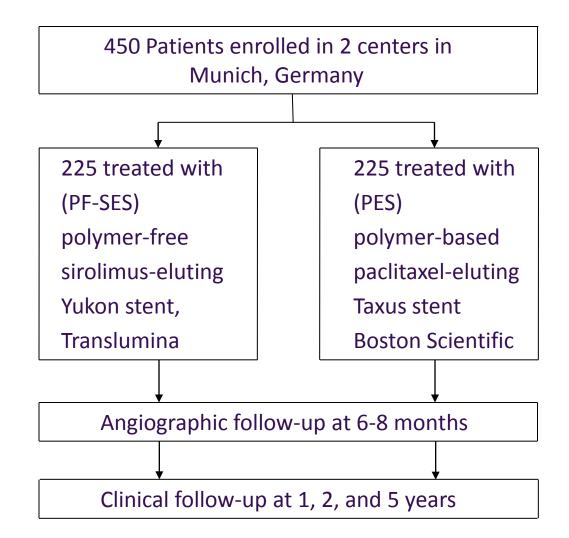
Prospective, randomized, two-center clinical trial

INCLUSION:

De novo native coronary artery stenosis ≥50% AND symptoms or objective evidence of ischaemia

EXCLUSION:

Left main stem disease
MI within 48 hours
Contraindication to study drugs





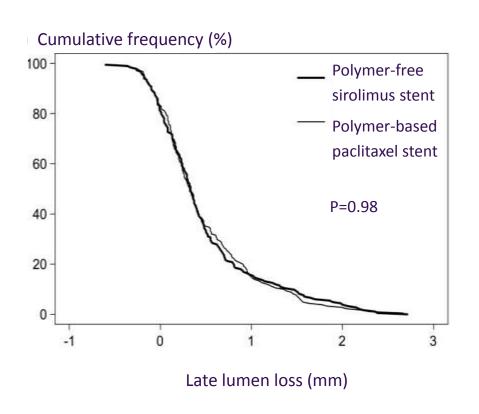


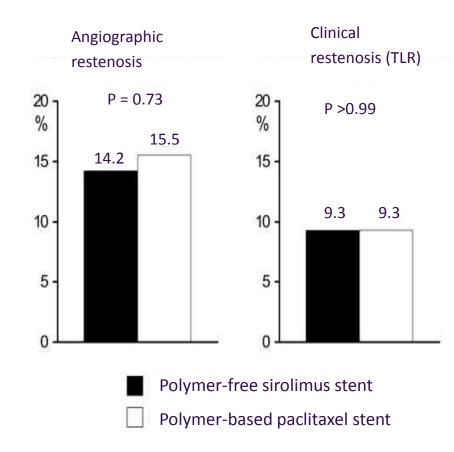






ISAR-TEST Primary Outcomes: Equivalence at 9 months















ISAR-TEST: 5 YEAR DATA



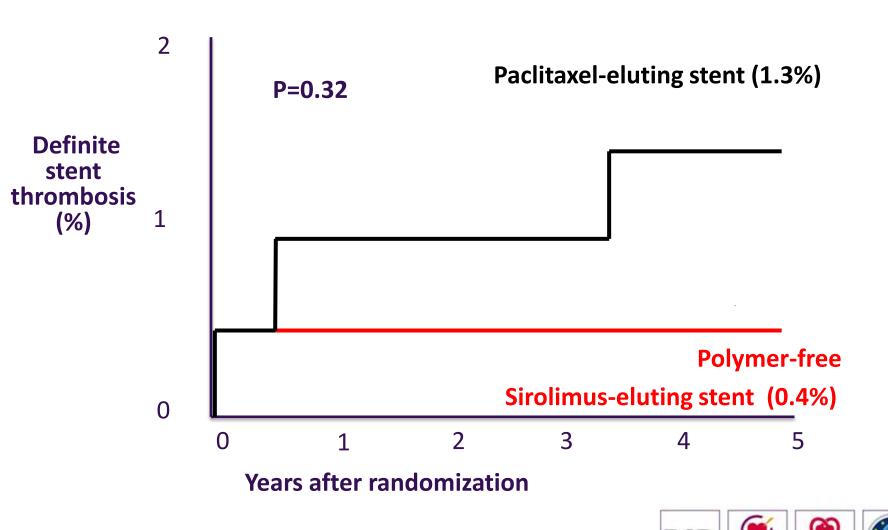






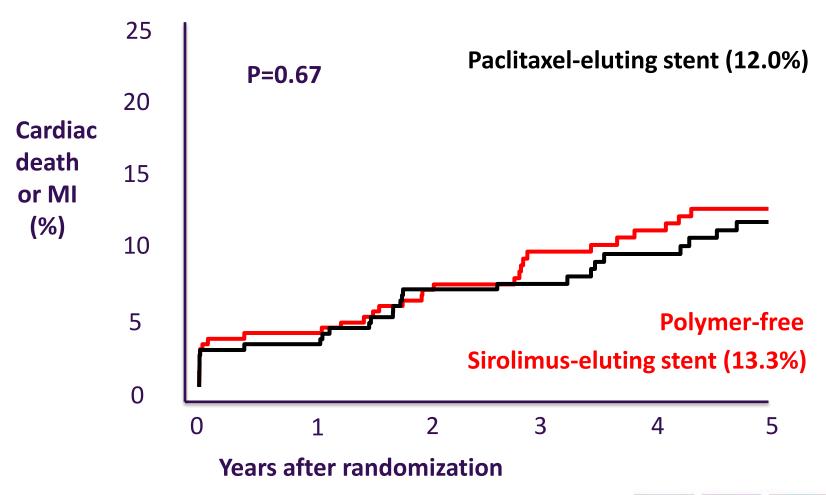


ISAR-TEST: Definite Stent Thrombosis





ISAR-TEST: Cardiac death or MI





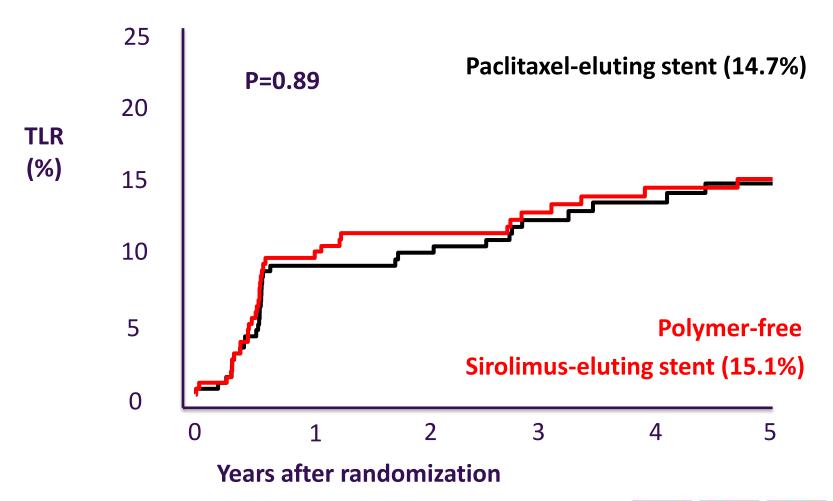








ISAR-TEST: TLR





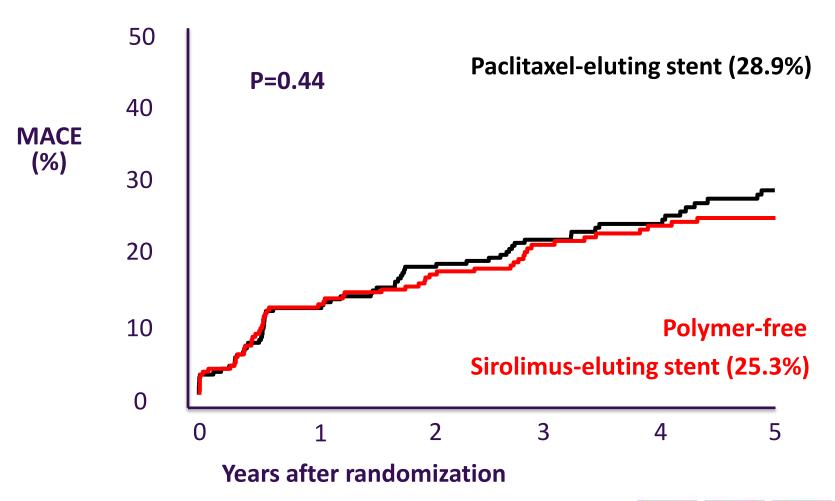








ISAR-TEST: MACE













ISAR-TEST: Conclusion

Low incidence of adverse events between year 1 and year 5 for both stent platforms

Polymer-free sirolimus-eluting stent demonstrates over 5 years

- ☐ durability of efficacy
- durability of safety

Comparable to a polymer-based stent platform







