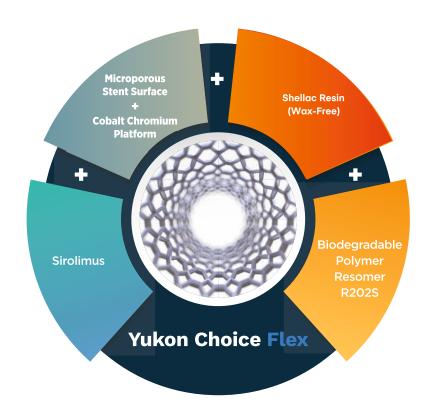
New generation DES providing synergy of biodegradable polymer with microporous surface to enhance optimal performance



Less Polymeric Load Compared To Other DES

One million pores per cm² with average depth of 2 µm ensures optimum drug release with minimal use of polymer

Shellac Resin (Wax-Free) ensures better polymer- drug biding with negligible polymer flaking during stent expansion

Drug and Polymer are co-released in 30 days

Better Endothelialisation & Superior Strut Coverage

Drug polymer matrix coated only on the abluminal side using patented stent coating technology for drug release only to target tissue

No polymer on the luminal side ensures healthy endothelialisation and reduces the incidence of stent thrombosis







Yukon Choice Flex

PRODUCT MATRIX / ORDERING INFORMATION*

| Length | Diameter (mm) | | | | | | | | | | |
|--------|---------------|----------|----------|-----------|-----------|----------|----------|--|--|--|--|
| (mm) | 2.00 | 2.25° | 2.50 | 2.75 | 3.00 | 3.50 | 4.00 | | | | |
| 8.00 | YCFX2008 | YCFX2208 | YCFX2508 | YCFX 2708 | YCFX 3008 | YCFX3508 | YCFX4008 | | | | |
| 12.00 | YCFX2012 | YCFX2212 | YCFX2512 | YOFX 2712 | YCFX 3012 | YCFX3512 | YCFX4012 | | | | |
| 16.00 | YCFX2016 | YCFX2216 | YCFX2516 | YCFX 2716 | YCFX 3016 | YCFX3516 | YCFX4016 | | | | |
| 18.00 | YCFX2018 | YCFX2218 | YCFX2518 | YCFX 2718 | YCFX 3018 | YCFX3518 | YCFX4018 | | | | |
| 21.00 | YCFX2021 | YCFX2221 | YCFX2521 | YOFX 2721 | YCFX 3021 | YCFX3521 | YCFX4021 | | | | |
| 24.00 | YCFX2024 | YCFX2224 | YCFX2524 | YCFX 2724 | YCFX 3024 | YCFX3524 | YCFX4024 | | | | |
| 28.00 | YCFX2028 | YCFX2228 | YCFX2528 | YCFX 2728 | YCFX 3028 | YCFX3528 | YCFX4028 | | | | |
| 32.00 | YCFX2032 | YCFX2232 | YCFX2532 | YCFX 2732 | YCFX 3032 | YCFX3532 | YCFX4032 | | | | |
| 36.00° | - | - | - | YCFX 2736 | YCFX 3036 | YCFX3536 | YCFX4036 | | | | |
| 40.00 | - | - | - | YCFX 2740 | YCFX 3040 | YCFX3540 | YCFX4040 | | | | |
| 44.00° | - | - | - | YCFX 2744 | YCFX 3044 | YCFX3544 | YCFX4044 | | | | |
| 48.00* | - | - | - | YCFX 2748 | YCFX 3048 | YCFX3548 | YCFX4048 | | | | |

Please contact our customer care for available sizes

Sizes Not CE Approved

COMPLIANCE CHART

| | Inflation Pressure (bar/10 ⁵ Pa) | | | | | | | | | | | | | | |
|-------------------|---|------|------|------|------|-----------------------------|------|------|------|------|----------------------------------|------|------|------|------|
| Balloon Ø [mm] | | | | | | NP (Nominal Pressure) | | | | | RBP (Rated Burst Pressure) | | | | |
| | 6 | 7 | 8 | 9 | 10 | 11 | 12 | 13 | 14 | 15 | 16 | 17 | 18 | 19 | 20 |
| Ø 2.00 | 1.83 | 1.87 | 1.90 | 1.93 | 1.96 | 2.00 | 2.03 | 2.06 | 2.10 | 2.13 | 2.16 | 2.20 | 2.23 | 2.26 | 2.29 |
| Ø 2.25 | 2.08 | 2.11 | 2.14 | 2.18 | 2.21 | 2.25 | 2.28 | 2.31 | 2.35 | 2.38 | 2.42 | 2.45 | 2.48 | 2.52 | 2.55 |
| Ø 2.50 | 2.33 | 2.36 | 2.40 | 2.43 | 2.47 | 2.50 | 2.53 | 2.57 | 2.60 | 2.64 | 2.67 | 2.70 | 2.74 | 2.77 | 2.81 |
| | 2.58 | 2.61 | 2.65 | 2.68 | 2.71 | 2.75 | 2.78 | 2.81 | 2.85 | 2.88 | 2.91 | 2.94 | 2.98 | 3.01 | 3.04 |
| Ø 3.00 | 2.81 | 2.85 | 2.89 | 2.92 | 2.96 | 3.00 | 3.04 | 3.07 | 3.11 | 3.15 | 3.18 | 3.22 | 3.26 | 3.29 | 3.33 |
| Ø 3.50 | 3.29 | 3.34 | 3.38 | 3.42 | 3.46 | 3.50 | 3.55 | 3.59 | 3.63 | 3.67 | 3.71 | 3.76 | 3.80 | 3.84 | 3.88 |
| Ø 4.00 | 3.75 | 3.80 | 3.85 | 3.90 | 3.95 | 4.00 | 4.06 | 4.11 | 4.16 | 4.21 | 4.26 | 4.31 | 4.36 | 4.41 | 4.46 |

TECHNICAL SPECIFICATIONS

| Cobalt Chromium Alloy (L605) | | | |
|------------------------------|---------------------|-------------------------|-------------------|
| Crossing profile | 0.035" / 0.889 mm | Entry Profile | 0.016" / 0.406 mm |
| Strut Thickness | 0.0027" / 68μm (SV) | Proximal Shaft Diameter | 1.9 F |
| | 0.0031" / 79μm (MV) | Distal Shaft Diameter | 2.7 F |
| Balloon Marker Material | Platinum / Iridium | Recommended Guide Wire | 0.014" |
| | | Guiding Catheter | min. 5 F |

Translumina Therapeutics LLP
Plot No. #12, Pharmacity, Selaqui, Dehradun 248011 (Uttarakhand) India Manufacturing License No. MFG/MD/2019/000227

Registered Office:

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Under Technological Collaboration With: Translumina GmbH Neue Rottenburger Strasse 50, D-72379 Hechingen, Germany

EC REP

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Email: mmateos@cmcmedical devices.com | web: www.cmcmedical devices.com

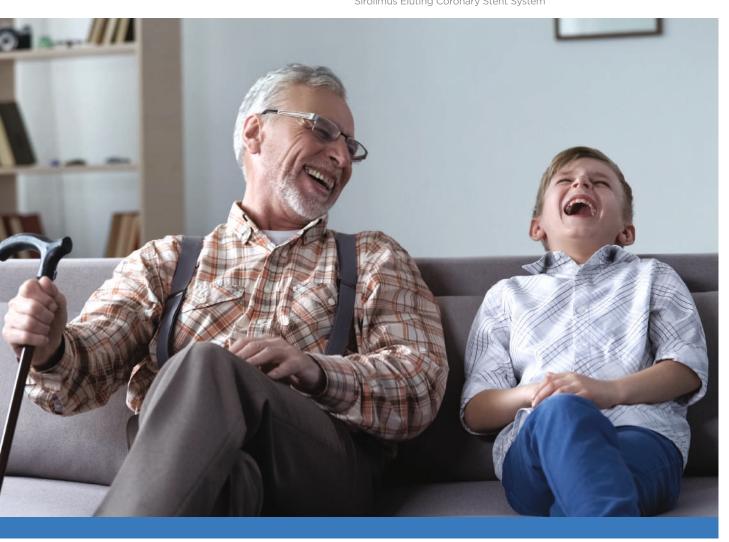
Customer Care No: + 91 11-2874 2874 Email: info@translumina.in

Web: www.translumina.in for more details.

Please refer to the Instructions for Use supplied with these devices for indications, $contraindications, Adverse\ Events, suggested\ procedures, warnings\ and\ precautions.$

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Yukon Choice Flex



Finding ways to the true joys of life.



Advanced platform for redefining flexibility in tortuous anatomy

Ideal Flexible Approach

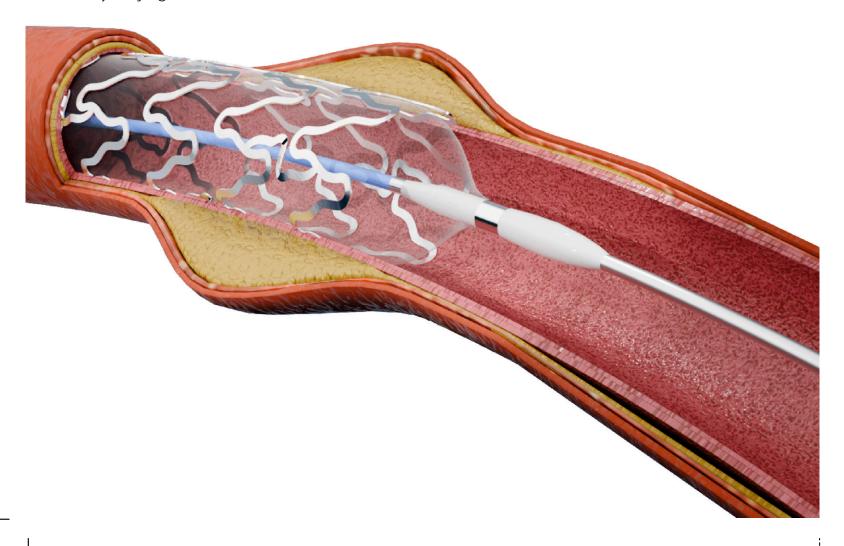
Yukon Choice Flex offers new generation delivery system with "Flexi" platform providing unmatched delivery in most tortuous vessels.

Enhanced Delivery System

The customized 2-Connector stent design of Yukon Choice Flex with thinner structural elements confirms for optimal deliverability.

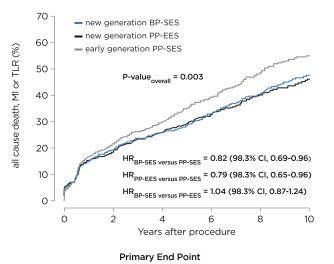
Proprietary Hypotube

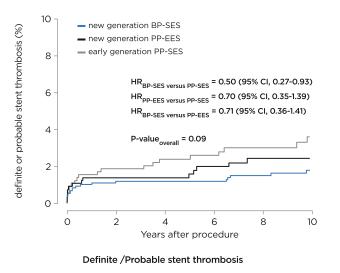
The new shaft design offers optimal force transfer with excellent push-ability and kink resistance allowing high manoeuvrability, justifying its use for the most tortuous vessels.





In this unique long term analysis at 10 years,
Yukon has shown the lowest rate of Definite/ Probable Stent
Thrombosis with a significant risk reduction than Cypher (50%) and
numerically lower TLR rates as compared to Xience (29%) while
maintaining the similar efficacy.





Comparison of clinical outcomes at 10 years in patients treated with new-generation BP-SES versus new-generation PP-EES versus early generation SES.

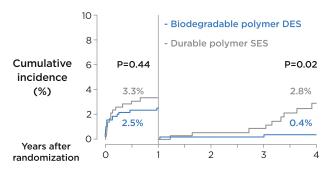


Unmatched Safety- In Complex Patients Subset

Long-term outcomes of biodegradable polymer versus durable polymer drug-eluting stents in **patients with diabetes:** a pooled analysis of individual patient data from 3 randomised trials



At 4 years, Biodegradable Polymer DES Yukon showed significantly lower rates of Stent Thrombosis compared to Durable Polymer SES in patients with Diabetes Mellitus.

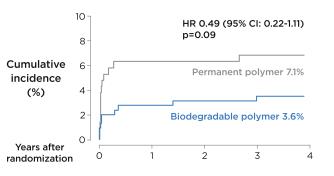


Secondary safety end point: definite or probable stent thrombosi

Long-term outcomes of biodegradable versus durable polymer drug-eluting stents in patients with acute ST-segment elevation myocardial infarction: a pooled analysis of individual patient data from three randomised trials

EuroIntervention

At 4 years, Biodegradable Polymer DES compared to Durable Polymer SES demonstrated improved overall clinical outcome, reduced need for revascularisation as well as lower incidence of cardiac death or MI and reduced stent thrombosis in patients with STEMI.



Definite or probable stent thrombosis for the pooled population in each of the treatment groups. Cl: confidence interval; HR: hazard ratio