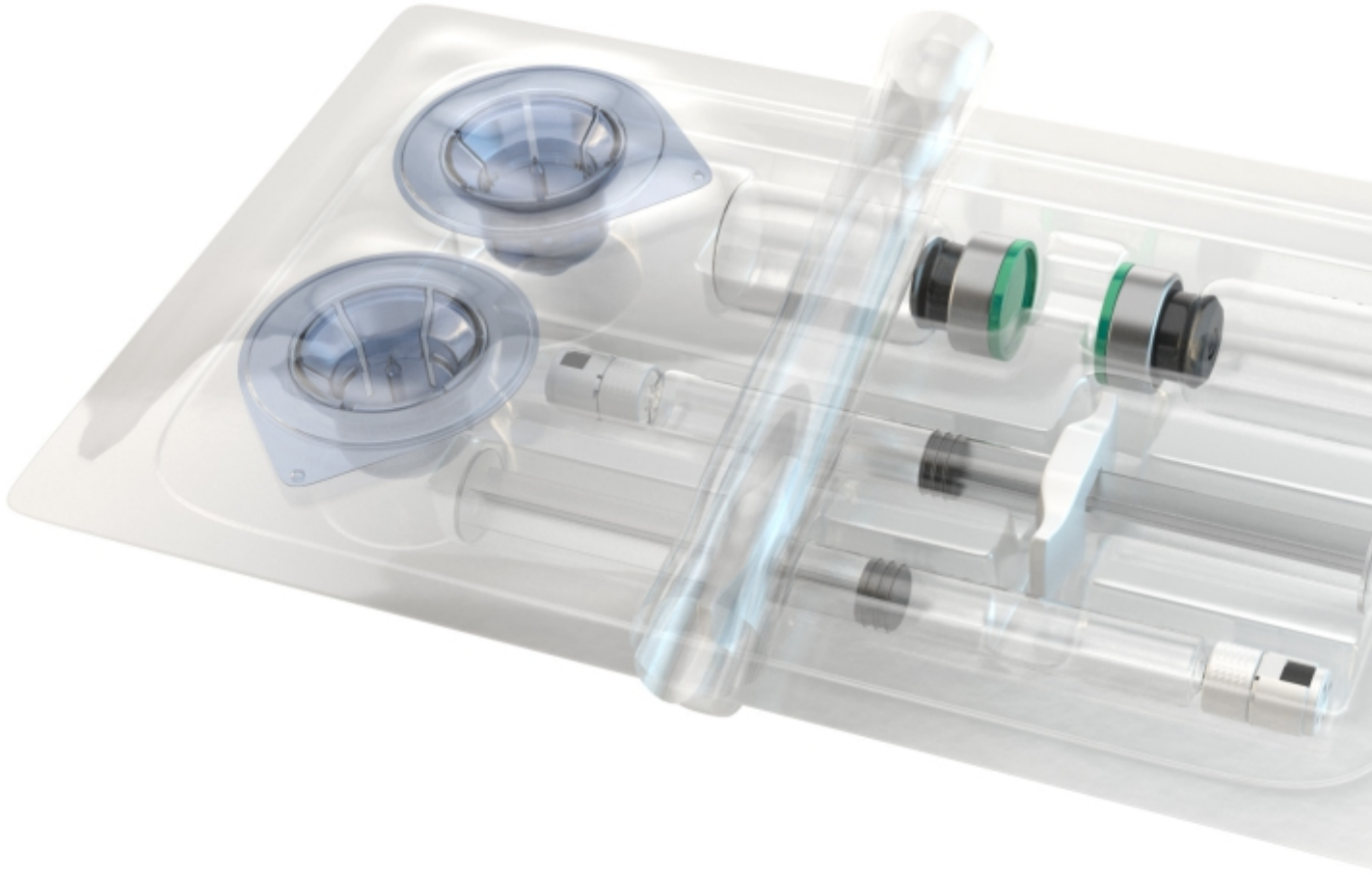
Your pharmaceutical product

packaged by SteriPack



SteriPack™
CONTRACT MANUFACTURING

01

Tablets & capsules
blistering & bottling

02

Kitting & serialisation
of pharmaceutical products

03

Medical devices
for pharmaceutical kits

04

Drug delivery devices
moulding & assembly

05

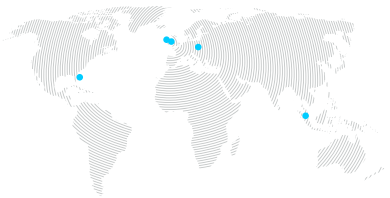
Local market
customisation

06

Rapid inspection,
re-packing & component replacement

Quality & Compliance

- Pharmaceutical Manufacturing Authorization for primary and secondary packaging (cGMP)
- ISO 13485:2016 Compliant to MDD 93/42/EEC, Annex II
- FDA registered and audited 21 PART 820 Quality System Regulation compliant
- Accredited to JPAL and CAN/CSA
- CAN/CSA ISO 13485:2016
- Accredited to MHLW Japanese Ministerial Ordinance No. 169
- CE marking
- 510K submission
- ISO 17025 Accredited Laboratory
- Class ISO 7 and 8 Cleanroom production
- Cleanrom Class D
- Humidity and temperature controlled manufacturing environment



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