



Earnings Quality Insurance Protection for Medical DevicesKey Features

Corporate Insurance Partner

EQuIP is an innovative non-damage business interruption cover. EQuIP protects Medical Device Manufacturers from losses arising from a suspension of manufacture due to regulatory non-compliance at Named Own and Named Supplier facilities.

Fully customizable dependent on client profile but with core key features as follows:

Coverage triggers

- Regulatory Order issued or instigated by a Defined Regulatory
 Agency (DRA*) including withdrawal of CE Mark or equivalent by
 an approved Notified Body resulting in suspension of manufacture
 due to irregularities in the manufacturing process
- Voluntary Suspension of manufacture to pre-empt a Regulatory
 Order including where due to use in manufacture of non-compliant inwards components, from a non-regulated named Supplier facility
- An order for Prohibition of Sale or Supply applied by a DRA to a named facility located in a non-DRA country (optional sub limited coverage)

Basis of indemnity

- Loss of Gross Margin and direct Increased Costs of Working including Remediation, Recall Costs, On-site Repair and Start-Up Costs
- Extra Expenses including Regulator Expert Fees and Product
 Destruction Costs (including compliance with European WEEE Regulations)

Indemnity period

 Up to 24 months (includes ongoing loss of Market Share AFTER re-start of manufacture)

Key exclusions

- Products and Prototypes not yet approved for sale
- Physical damage events (as covered in standard PDBI policies)
- Cyber and Network Outage Incidents (as covered in specialist cyber policies)

*DRA – authorities that are covered as standard in respect of enforcement of good manufacturing practices are USA, Canada, EU, Switzerland, Norway, Israel, Japan, Taiwan, South Korea, Singapore Australia, New Zealand.

For further details please contact:

Jenny Yu Senior Underwriter Tel.: +44 20 30 03-7456 jyu@munichre.com



Petra Mates Underwriter Tel.: +44 20 30 03 72 93 Mobile: +44 7976 824705 pmates@munichre.com



Visitor address

Münchener Rückversicherungs-Gesellschaft Berliner Strasse 95 80805 München Germany

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