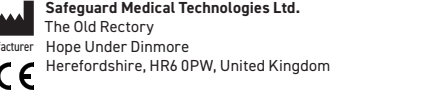


SWAT-T™

Stretch, Wrap, and Tuck Tourniquet





Safeguard Medical Technologies Ltd.
The Old Rectory
Hope Under Down
Herefordshire, HR6 0PW, United Kingdom

UDI Device Identifier
5464422011515

REF Reference Number
P0164

CE REP
Procamed AG
Schulstrasse 5
CH-8355 Aadorf
Switzerland

Intended use
The SWAT-T™ is a non-pneumatic tourniquet designed to stop severe traumatic bleeding before or during transport to a care facility.

User profile
The intended users are first responders, to include self-care, laypersons, emergency medical personnel and health care professionals. Special training is not required.

Patient profile
To be used on those patients with severe bleeding from a limb.

Clinical benefits
SWAT-T™ controls bleeding from an extremity by circumferentially compressing the injured limb, at least 5 cm above the bleed site, to an extent where arterial occlusive pressure is reached.

Performance characteristics

- Is available in two colours: black and orange
- Can remain secure when wet or dry
- Is highly portable

Product claims
Device has been designed to control bleeding from an extremity by circumferentially compressing the injured limb, above the bleed site, to an extent where arterial occlusive pressure is reached.

Contraindications
N/A

RESIDUAL RISKS/ WARNINGS: Safe for 2 hours. Longer application may cause serious loss of extremity. Reassess for loosening. Dispose of as clinical waste. Do not reuse.

1. Apply between wound and heart.

2. Wrap tightly while pulling tension.

3. To secure: Tuck end under itself.

4. Tourniquet fits: Ovals/ rectangles turn to circles/square.

Reporting: Any serious incident that has occurred in relation to this device should be reported to manufacturer (vigilance@safeguardmedical.com) and applicable national competent authority.

Reporting: Qualsiasi incidente grave che si è verificato in relazione al dispositivo deve essere comunicato al produttore (vigilance@safeguardmedical.com) e all'autorità nazionale competente.

Reporting: Ethver alvorlig hendelse, der har hendes sted forbindelse med denne anordning, skal indberettes til producenten (vigilance@safeguardmedical.com) og de relevante nationale kompetente myndighed.

Reporting: Kijakni svakaý incidenc, ke ktrému dolož v súvislosti s týmto prostredkom, je nutné nahlásiť výrobcovi (vigilance@safeguardmedical.com) a príslušnému národnému orgánu.

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Reporting: Kijakni svakaý incidenc, ke ktrému dolož v súvislosti s týmto pomôckou, sa musí nahlásiť výrobcovi (vigilance@safeguardmedical.com) a príslušnému národnému orgánu.

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