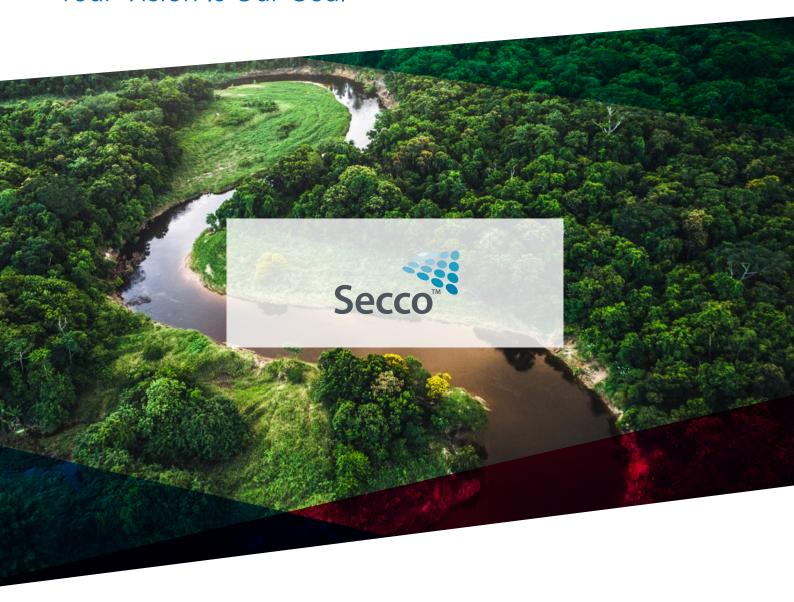
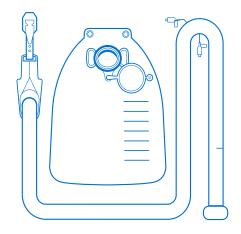
### Your Vision Is Our Goal





# Secco Protect™ Safer and Easier to Use

Contain faecal contamination, maintain a healthy patient environment



## Secco Protect™: Safer and Easier to Use

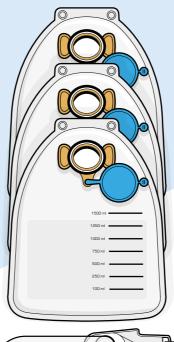
34.5% of hospital-acquired pressure ulcers (PU) occur in patients with medical devices<sup>1</sup> Patients with medical devices are 2.4 times more likely to develop PUs of any kind<sup>1</sup>

Secco Protect™ is a closed catheter system for the management of episodic faecal incontinence. Small pressure ulcers or 'hot-spots' aren't uncommon in patients with a bowel catheter, and can lead to longer hospitalisations, increased clinical intervention, incident reporting, and potential discomfort and anxiety for the patient.

- Silicone tube parylene coated for low surface friction 'easy flow' and reduction of odour transmission
- Patient Safety Valve provides inflation protection at max 45ml water
- Twin finger Easy Insertion Cover locator
- Super-absorbent insert pad in waste bag to gel content and reduce cross infection by spillage
- 1.5 litre capacity
- · 'L' shaped elbow to take fluid into bag reduces tube twisting
- Sealing cap for each full bag pre-attached
- Odour proof bag film and flatus filters

## **Contain, Maintain and Control**

Secco Protect<sup>™</sup> - the easy to use Faecal Management System that reduces skin breakdown and pressure ulcer development whilst preventing cross contamination. With its unique super absorbency technology, it contains spillages and odours as well as maintaining patient dignity.



### CONTAIN

Helps prevent the spread of infection for patients and staff with its closed catheter system and super-absorbent technology. This results in less risk of infection and spillages, whilst maintaining a healthy patient environment.

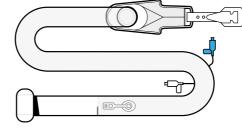
MAINTAIN

Maintain greater patient

Maintain greater patient comfort due to its
Thinwall Technology. Secco Protect™ can reduce
skin breakdown and development of pressure
ulcers commonly seen in bowel catheter patients.

CONTROL

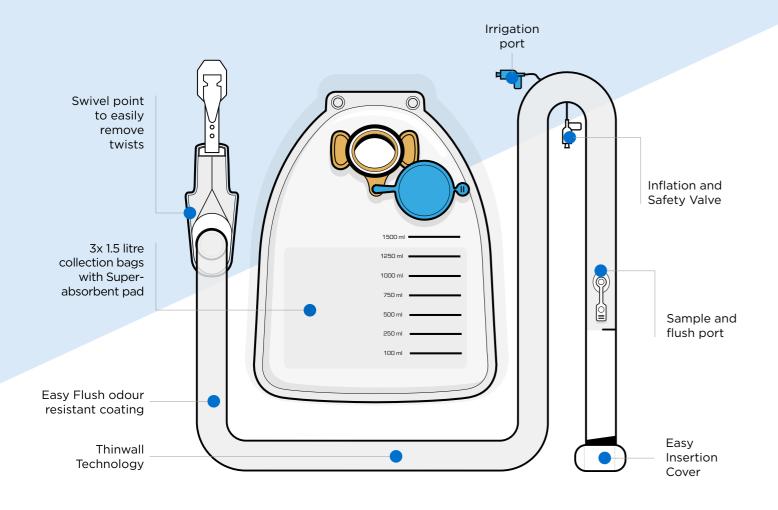
Control odours and improve patient dignity. The EasyFlush odour resistant coating along with the deodorising filter ensures a more pleasant ward environment whilst helping to protect patient dignity. Added with its 1.5 litre collection bag means 50% fewer replacement bags required.

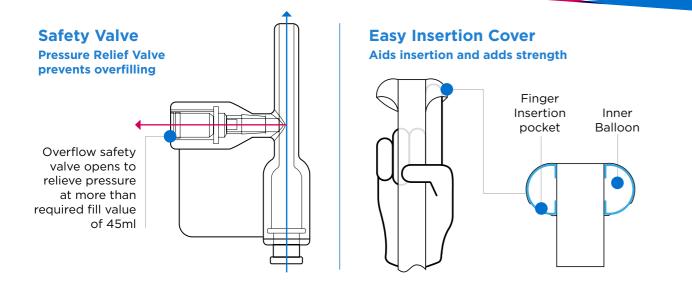


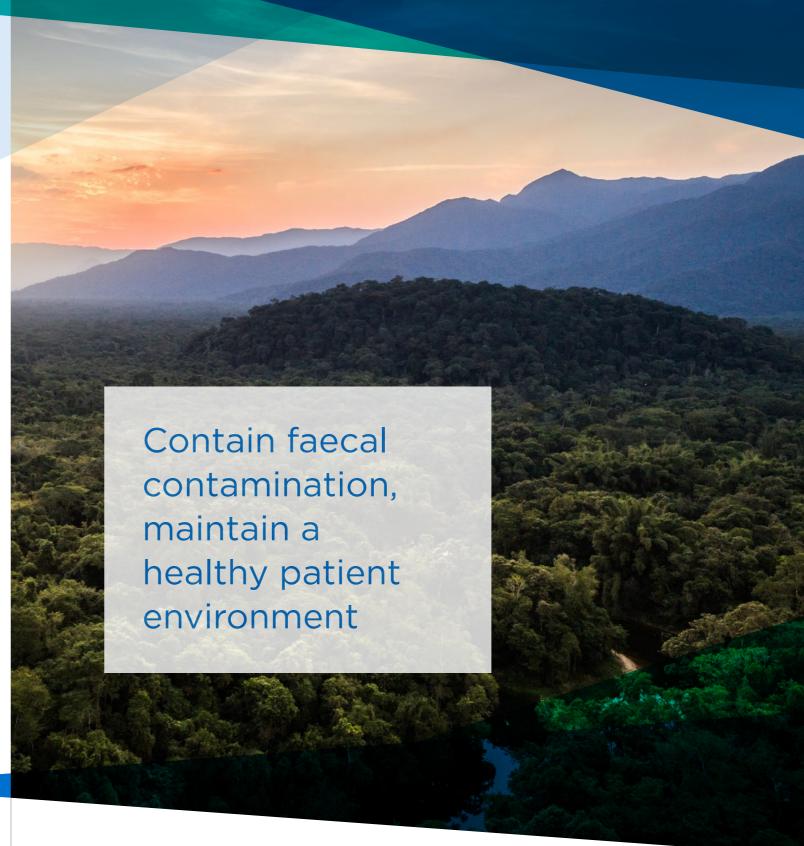
Click to watch the Secco Protect™ animation



# Improved design







#### Features prevent overfilling and aids insertion

- Extra safety, prevents overfilling balloon cuff
- Attempted water overfilling is indicated by overflow safety valve, which opens to relieve pressure at more than required fill value of 45ml
- Easy insertion balloon cover aids product insertion into anus and adds strength to help restrict overfilling. This added feature prevents patient injury from an over inflated balloon cuff
- Improved nurse user feedback during testing on application

# Secco Protect™ saves more than just patient dignity

- The cost of a longer stay in hospital due to C. difficile associated disease is estimated to be £4,000 per case<sup>2</sup>
- Patients with faecal incontinence are three times more likely to develop pressure ulcers than those who are continent<sup>3</sup>

- 1. Black JM, Cuddigan JE, Walko MA, et al (2010) Medical device related pressure ulcers in hospitalized patients. Int Wound J7(5)
- 2. M.H. Wilcox, W.N. Fawley, N. Wigglesworth, P. Parnell, P. Verity, J. Freeman. Comparison of the effect of detergent versus hypochlorite cleaning on environmental contamination and incidence of Clostridium difficile infection. Journal of Hospital Infection 2003 54, 109-114
- 3. Theaker C, Mannan M, Ives N, Soni N. Risk factors for pressure sores in the critically ill. Anaesthesia. 2000 Mar;55(3):221-4



P. J. Dahlhausen & Co. GmbH, Emil-Hoffmann-Strasse 53, 50996 Köln, Germany

#### Distributed by





Suite 303, Highland House, 165 The Broadway, Wimbledon, SW19 1NE, UK

**L** +44 (0)20 8944 7585

www.prosysinternational.co.uk

The products meets the provisions (or requirements) of the Medical Device Directive 93/42/EEC.