

Technical Data Sheet

Pull1Aid® Blood Stopper 4-in-1

1. Identification of the product and the company

1.1. Product identifier and intended use

Trade Name:	Pull1Aid® Blood Stopper 4-in-1 Mini / Pull1Aid® Blood Stopper 4-in-1
Intended Use:	Plum Pull1Aid® Blood Stopper 4-in-1 is intended to be used as a mechanical barrier and compression.
Expiry date:	Printed on the sealed package.

1.2. Details of the supplier of the data sheet

Distributor	
Company:	Plum Safety ApS
Address:	Mandelalleen 1
Zip code:	5610
City:	Assens
Country:	Denmark
E-mail	info@plum.eu
Phone	+45 6916 9600
Fax	+45 6916 9630

1.3 Legislative information

General information:	As the products are class 1s sterile medical devices according to <i>Council Directive 93/42/EEC of 14 June 1993</i> concerning medical devices, they are certified by the Notified Body, Intertek Semko AB, Kista, Sweden. No Safety Data Sheet is required for medical devices.
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AKLA AB, Parkgatan 15, 696 33 Askersund, Sweden is the legal manufacturer of Pull1Aid® Blood Stopper 4-in-1 Mini and Pull1Aid® Blood Stopper 4-in-1.

Plum Safety ApS is certified according to *ISO 13485:2016 Medical devices – Quality management systems – Requirements for regulatory purposes*.

2. General information about the product

Material requirements:	The materials in these products have been judged biocompatible according to the requirements of <i>ISO 10993-1:2018 Biological evaluation of medical devices – Part 1: Evaluation and testing within a risk management process</i> .
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Physical properties and composition:

	Pull1Aid® Blood Stopper 4-in-1 Mini Art. No. 5153 (REF. 92109)	Pull1Aid® Blood Stopper 4-in-1 Art. No. 5154 (REF 92111PM)
Material	Nonwoven dressing, small, in CELLOSOFT-material, 3-layer combination of: 100% LDPE-HDPE 30g/m ² micro perforated. 100% Viscose 200 g/m ² +/- 10% 100% LDPE-HDPE 30g/m ² micro perforated non-adhesive, stitched on elastic bandage 2m x 6cm.	Nonwoven dressing, large, in CELLOSOFT-material: 70% Viscose. 30% Polyethylene non-adhesive, stitched on elastic bandage 4m x 6cm, Extra bandage 3m x 6cm Bandage: Polyamid 60%/Viscose 40%.
Primary Packing (Pouch)	Cold sealed paper	Cold sealed paper
Secondary Packing (Box)		
Absorption	≥ 6g/g	≥ 800g/m ²
Dimensions	7,5 x 12cm (3 pcs)	17 x 17 cm
Special feature	Suitable for hand, finger and toes	Suitable as pressure bandage, protective dressing.

Technical Data Sheet

Pull1Aid® Blood Stopper 4-in-1

3. Side effects and malfunction

Side effects and malfunction:

The products are designed for skin contact and first aid use involving wounds.

If side effects or malfunctions are experienced, please report these to Plum Safety ApS, info@plum.eu or phone +45 6916 9600, so that necessary remedial measures can be initiated.

Reporting of accidents and incidents. Vigilance. Serious incidents involving products made available on the EU market must be reported immediately to the National Competent Authority. All serious incidents must be reported no later than 15 days after discovery, in the event of a serious threat to public health within 2 days. In the event of death or an unexpected and serious deterioration of a person's state of health, reporting must take place within 10 days.

4. Storage

Shelf life: 5 years from production date

5. Other information

Do not reuse



Do not use if package is damaged



Version history and indication of changes

Version	Revision date	Responsible	Changes
1.1	08-07-2025	MAW	1.3 + 2
1.0	20-06-2022	CL	

Other information Additional information can be obtained by contacting info@plum.eu or by telephone +45 6916 9600.

TDS prepared by:

Company: Plum Safety ApS
Address: Mandelalleen 1
Zip code: 5610
City: Assens
Country: DENMARK
E-mail: info@plum.eu
Phone: +45 6916 9600
Fax: +45 6916 9630

Document Language: English

Technical Data Sheet

Pull1Aid® BurnGel

1. Identification of the product and the company

1.1. Product identifier and intended use

Trade Name:	Pull1Aid® Burn Gel
Intended Use:	Plum Pull1Aid® Burn Gel is a sterile dressing with cooling gel intended to be used for burns.
Expiry date:	Printed on the sealed package.

1.2. Details of the supplier of the data sheet

Distributor	
Company:	Plum Safety ApS
Address:	Mandelalleen 1
Zip code:	5610
City:	Assens
Country	Denmark
E-mail	info@plum.eu
Phone	+45 6916 9600
Fax	+45 6916 9630

1.3 Legislative information

As the products are class IIb sterile medical devices according to *Council Directive 93/42/EEC of 14 June 1993* concerning medical devices, they are certified by the Notified Body, Intertek Semko AB, Kista, Sweden.
No Safety Data Sheet is required for medical devices.



Plum Safety ApS is certified according to *ISO 13485:2016 Medical devices – Quality management systems – Requirements for regulatory purposes*.

AKLA AB, Parkgatan 15, 696 33 Askersund, Sweden is the legal manufacturer of Pull1Aid® BurnGel.

2. General information about the product

Material requirements:	The materials in these products have been judged biocompatible according to the requirements of <i>ISO 10993-1:2018 Biological evaluation of medical devices – Part 1: Evaluation and testing within a risk management process</i> .
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Physical properties and composition:

	Pull1Aid® BurnGel Art. No. 5156 (REF. 94373)
Material	<i>Polyurethane, foam compress with 50g gel.</i> <i>Ingredients: Purified water, Melaleuca Oil (1%), Emulsifiers and preservatives</i>
Primary Packing (Multi layered pouch)	
Secondary Packing (Box)	
Dimensions	<i>10 x 10 cm</i>
Special feature	<i>Can be used as first aid on all degrees of thermal burns.</i> <i>Suitable as first aid on all skin areas. In case of eye contact rinse with water!</i>

Technical Data Sheet

Pull1Aid® BurnGel

3. Side effects and malfunction

Side effects and malfunction:

The products are designed for skin contact and first aid use involving burns.

If side effects or malfunctions are experienced, please report these to Plum Safety ApS, info@plum.eu or phone +45 6916 9600, so that necessary remedial measures can be initiated.

Reporting of accidents and incidents. Vigilance. Serious incidents involving products made available on the EU market must be reported immediately to the National Competent Authority. All serious incidents must be reported no later than 15 days after discovery, in the event of a serious threat to public health within 2 days. In the event of death or an unexpected and serious deterioration of a person's state of health, reporting must take place within 10 days.

4. Storage

Shelf life: 5 years from production date

5. Other information

Do not reuse



Do not use if package is damaged



Version history and indication of changes

Version	Revision date	Responsible	Changes
1.1	08-07-2025	MAW	1.3 + 2
1.0	20-06-2022	CL	

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Document Language: English

Technical Data Sheet

Pull1Aid® Protect Kit

1. Identification of the product and the company

1.1. Product identifier and intended use

Trade Name:	Pull1Aid® Protect Kit
Intended Use:	Plum Pull1Aid® Protect Kit to limit contact between the first aid provider and the injured person.
Expiry date:	Printed on the sealed package, when needed.

1.2. Details of the supplier of the data sheet







Distributor	
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1.3 Legislative information

General information:	REF 97106K, Respiratory sheet: Are class 1 medical devices according to <i>Council Directive 93/42/EEC of 14 June 1993</i> concerning medical devices. <i>No Safety Data Sheet is required for medical devices.</i> Legal manufacturer is FIRSTAR HEALTHCARE Co. Ltd. (Guangzhou), Rm 901, Building No. 2, Headquarters Center, Tian'an High-tech Ecological Park, Panyu, 511400, Guangzhou, P.R. China Distributor: AKLA AB, Parkgatan 15, 696 33 Askersund, Sweden Commercial contact: Plum Safety ApS, Mandelalleen 1, 5610 Assens, Denmark REF 94416ST, Disinfection wipe: Reference to MSDS REF 2080809001, Protective Gloves: Not classified
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2. General information about the product

Physical properties and composition:

Pull1Aid®Protect Kit. Art no. 5155	REF. 97106K, Respiratory sheet	REF. 94416ST, Disinfection wipe	REF. 2080809001, Protective Gloves (1 pair)
Material	Valve= PP, Sheet=PE	Nonwoven tissue, Cellulose, Viscose, Co-polyester/Polyefine saturated with 70% alcohol solution (alcohol, aqua, glycerine)	Powder-free vinyl gloves, unsterile, transparent, smooth
Primary packing			
Secondary packing			

Technical Data Sheet

Pull1Aid® Protect Kit

Dimensions	-	17 x 22 cm	Size 8-9 (Ø105xL230mm), Thickness 0,05 mm.
Special features	For mouth-to-mouth ventilation	For disinfection in case of first aid.	For hand protection in case of first aid.

3. Side effects and malfunction

Side effects and malfunction:

The products are designed for skin contact and first aid use involving burns. If side effects or malfunctions are experienced, please report these to Plum Safety ApS, info@plum.eu or phone +45 6916 9600, so that necessary remedial measures can be initiated.

Reporting of accidents and incidents. Vigilance. Serious incidents involving products made available on the EU market must be reported immediately to the National Competent Authority. All serious incidents must be reported no later than 15 days after discovery, in the event of a serious threat to public health within 2 days. In the event of death or an unexpected and serious deterioration of a person's state of health, reporting must take place within 10 days.

4. Storage

Shelf life: 5 years from production date

5. Other information

Do not reuse

**Version history and indication of changes**

Version	Revision date	Responsible	Changes
1.1	08-07-2025	MAW	1.3 + 2
1.0	20-06-2022	CL	

Other information Additional information can be obtained by contacting info@plum.eu or by telephone +45 6916 9600.

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