	Technical Data Sheet			
Use in	 Pharmaceutical Industry in clean rooms and isolators For industrial, laboratory & research applications only Basic medium according to EP 2.6.12, 2.6.13 and USP <61>, <62> 			
Use for	 Detection of aerobic and anaerobic micro-organisms Contact sampling, personnel monitoring, as well as active air monitoring Isolation and growth of fastidious bacteria, yeasts and moulds Universal neutralization of residues of disinfectants The medium should be applied with a uniform and steady pressure to the surface for few seconds. After sampling the surface must be cleaned to remove residues of the medium. 			
Typical composition per liter	Casein peptone 15 g Lecithin (L) 0.7 g Soy peptone 5 g Polysorbate 80 (T) 5 g NaCl 5 g Histidine (H) 0.5 g Agar 15 g Thiosulfate (T) 0.1 g Neutralizer PLUS This medium can be adjusted / or supplemented according to the performance criteria required.			
Irradiation	Irradiated at 9-20 kGy			
Filling volume	• 16-19 mL			
Packaging	 Triple bagged, staples of 10 plates Transparent High barrier foil for H₂O₂ as well as for water-vapor 10 staples of 10 plates per packaging unit Temperature isolated handle-bag in the cardboard-boxes 			
Plates per box	100 (10 staples with 10 plates each)			
Shelf life	12 months from production date			
Storage conditions	 Recommended storage temperature: 15-25 °C Should be stored at temperatures as stable as possible Store protected from light exposure Before use: it is recommended to keep the plates upright (agar on the lower part, lid on the upper part) to avoid formation of extra condensation After use: it is recommended to keep the plates upside down (agar on the upper part, lid on the lower part) to reduce the risk of accumulation of condensation during incubation which can affect colony formation 			
Label	On the side, at the bottom			



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Label information	 Product name: TSA - U+ Expiry date: YYYYMMMDD → MMM in letters (e.g.: 2023Nov04) Lot-number Individual number Barcode 		
Barcode	 2-dimensional (data matrix), 20 digits: Digits 1-3: ArtNo. Digits 4-9: Lot-Number Digits 10-14: Individual-Number Digits 15-20: Date (YYMMDD) 		
Delivery	 Temperature controlled delivery on request For shipments of larger amounts plastic pallets in Euro-size can be used 		
Petri dish	 Locking-lid plate, made from polystyrene Inner diameter: ~ 56.5 mm, thus providing an area of ~25 cm² Outer diameter: ~ 66 mm Bottom part with 1 cm² square grid for facilitated evaluation Incubations in vent and closed position possible Specific design to improve binding of agar to plate Easy handling due to increased handling area 		
Lid positions	 All plates are delivered in the non-locked position The plate contains 2 locked positions. If turning the lid clockwise the locked positions are in the following order: 1. Vent position 2. Closed position 		
Aerobic incubation	 The closed position provides ideal incubation conditions for aerobic microorganisms Limits the dehydration of the agar during incubation For long incubation of aerobic microorganisms, the closed position is recommended To lock the lid in the closed position turn the lid clockwise into the final stop position 		
Anaerobic incubation	 The vent position is ideal for anaerobic incubations, as it allows an easy and effective removal of oxygen under anaerobic incubation conditions Incubate in anaerobic incubator, anaerobic jar or suitable equipment First option: Turn the lid clockwise into the final stop position Turn the lid one click counter-clock-wise to the vent position Second option: Turn the lid clockwise directly into the first locked position 		



	Technical Data Sheet
Place of	PharmaMedia Dr. Müller GmbH
production	Gustav-Throm-Str. 1, 69181 Leimen - Germany

	Quality control, Certificates				
	Each lot of product can be obtained with a certificate of analysis (CoA):				
	Physico-chemical	test parameter	rs:		
	Appearance	Slightly turbid,			
	pH value	7.1 – 7.5			
	Filling volume	16 – 19 mL			
	Irradiation	9-20 kGy			
	Growth Promotion	n test: 10-100 C	FU		
Coutificates	S. aureus	ATCC 6538	30-35 °C	1 day	50-200%
Certificates	E. coli	ATCC 8739	30-35 °C	1 day	50-200%
	P. paraeruginosa	ATCC 9027	30-35 °C	1 day	50-200%
	B. spizizenii	ATCC 6633	30-35 °C	1 day	50-200%
	C. albicans	ATCC 10231	20-25 °C	3-5 days	50-200%
	A. brasiliensis	ATCC 16404	20-25 °C	3-5 days	50-200%
	Neutralizer PLUS	test: 10-100 CF			
	B. spizizenii	ATCC 6633	30-35 °C	1 day	50-200%
	Sterility control				No growth
Certificate of origin	All media lots produced by PMM can be obtained with a Certificate of Origin (CoO). All animal derived raw materials are specified as follows: Raw material Tissue Animal source Country of origin Infectivity category (acc. to TSE guideline: EMA/410/01 current version)				
BSE policy	 In compliance with transmitting animal medicinal product specified animals. We neither store infectivity tissues originates from collins. 	al spongiform e ts, we check the source, the cour or process rum (IA) nor rumina	ncephalopa e CoO of rantry of origin hinant raw rantraw rantraw	thy via hun w material and the inf naterials ob terials whos	nan or veterinary in respect to the fectivity category. stained from high se animal source



Quality control, Certificates Art. 101.0100 has been exposed to temperature stress conditions (3 days at 2-8 °C as well as 3 days at 30-35 °C) and has passed shelf-life testing **Temperature** at least 30 days after the assigned expiry date. Shelf-life testing comprises stress all regular tests of the normal release test of this article except for sterility control (see CoA). The disinfection of surfaces is crucial for maintaining an adequate environment for the production of sterile pharmaceutical drugs. To guarantee the best possible success of the disinfection process many pharmaceutical companies do perform a regular rotation of the disinfectants used. Quite often at least one of the disinfectants used contains quaternary ammonium compounds, benzalkonium compounds, biguanides or even a combination of these substances. The advantages of such disinfectants are the well proved bactericidal activity against microorganisms even if used in relatively low concentrations. However, the disadvantages are the residues which remain on treated surfaces, if not removed by a suitable cleaning step. The removal or inactivation of residues of disinfectants is critical for the reliable detection of viable and cultivable microorganisms. If highly active residues remain on surfaces, these will be picked up with contact plates or swabs when performing environmental monitoring tests. Then these residues can interfere with the growth of potential contaminants and this could finally result in false negative results. Whereas some residues of disinfectants can be neutralized with the standard neutralizers LTHT (Lecithin, Tween 80, Histidine and Thiosulfate - please see product description of art. 100.0100) especially the residues of quaternary **Neutralization** ammonium compounds, benzalkonium compounds as well as biguanides are not of residues of sufficiently inactivated by these neutralizers. disinfectants To overcome this unsatisfactory inactivation of these residues, media manufacturers have tried to develop special neutralizer media. However, most of the media offered so far had different drawbacks: turbidity, precipitation, short shelf-life, low recovery rates on Gram positive strains and quite high price - and due to these disadvantages, such media have not been really accepted. PMM now offers a newly designed plate without showing these drawbacks. TSA U+ plates look-like a regular TSA plate and are free of precipitation throughout the shelf-life of more than 9 months. However, the outstanding inactivation of all typically used disinfectants including even high concentrations of quaternary ammonium compounds, benzalkonium compounds and biguanides really is the big step forward in obtaining reliable results for the environmental monitoring. TSA U+ plates were tested with respect to the inactivation of disinfectants using the worst-case approach by directly inoculating defined amounts of disinfectant on the agar plates. Typically, 20 µL, 50 µL or 100 µL of disinfectant was used. 100 µL of disinfectant applied to a contact plate of about 25 cm² surface correspond to about 40 mL of disinfectant used to disinfect an area of one square meter, a concentration typically used in the pharmaceutical industry. After a period of 15 to 20 min the test organisms were applied to the treated plates.

Quality control, Certificates

Test organisms used for such neutralization tests could be for example *B. spizizenii* ATCC 6633, *S. aureus* ATCC 6538 and *S. epidermidis* ATCC 14990 as well as *E. coli* ATCC 8739, *P. paraeruginosa* ATCC 9027, *C. albicans* ATCC 10231 and *A. brasiliensis* ATCC 16404. However, as Gram positive microorganisms are typically more sensitive to quaternary ammonium compounds, it is recommended to perform the tests with Gram positive microorganisms.

As reference, plates not treated with disinfectant are used.

Specifications: for sufficient inactivation of disinfectants the amount of 50 μ L of a disinfectant applied to a contact plate must be inactivated, resulting in a recovery rate of more than 50%.

Results: Beside the disinfectants inactivated already by our standard plate (see product description of art. 100.0100) **TSA U+ plates** are as well inactivating quite high concentrations of quaternary ammonium compounds, biguanides and benzylalkoniumchlorides. Disinfectants tested were Amphospray 41 IP, Gigasept AF (4%), Hexanios G+R, Hexaquart forte (2%), Incidin plus (2%), Biocide A, Biocide B, Lysoformin 3000 (2%), Melsept SF (2%), Microbac forte (2%) and Terralin protect (2%).

Results obtained with the above listed disinfectants show recovery rates of more than 70% if 20 or 50 μL of the disinfectant was applied directly on **TSA U+ plates**. Even when applying 100 μL most recovery rates were above 70%, only few recovery rates dropped to values between 30 to 50%. In comparison to these results standard TSA plates with neutralizers did not show any or very low recovery rates even if only 20 μL of these disinfectants were applied. As a conclusion **TSA U+ plates** can be considered as the universal media for performing environmental monitoring, delivering reliable results independent from the disinfectant used.

	Safety Data	
Toxic ingredients	• None	
Basic composition	See typical composition	
Solvent content	• None	
Safety data sheet required	Not mandatorily required	