	Technical Data Sheet		
Use in	 Pharmaceutical Industry in clean rooms and isolators For industrial, laboratory & research applications only Basic medium according to EP 2.6.13 and USP <62> 		
Use for	 Isolation and growth of yeasts and molds Contact sampling, personnel monitoring, as well as active air monitoring Inhibits the growth of most bacteria The medium should be applied with a uniform and steady pressure to the surface for a few seconds. After sampling the surface must be cleaned to remove residues of the medium. 		
Typical composition per liter	Casein peptone5 gLecithin (L)0,7 gMeat peptone5 gPolysorbate 80 (T)5,0 gGlucose-D(+)*H2O44 g*Histidine (H)0,5 gAgar15 gThiosulfate (T)0,1 gThis medium can be adjusted / or supplemented according to the performance criteria required.*Glucose-D(+)+H2O = Glucose monohydrate*Glucose-D(+)+H2O = Glucose monohydrate*44 g Glucose monohydrate = 40 g Glucose = 40 g Dextrose		
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 		
Units per pack	100 plates		
Shelf life	12 months from production date		
Storage conditions	 Recommended storage temperature: 15-25 °C Should be stored at temperatures as stable as possible Before use: it is recommended to keep the plates upright with the agar always on the bottom For incubation: it is recommended to keep the plates upside down for reducing the risk of condensation dropping on the agar surface, thus affecting colonies growing on the surface 		
Label	On the side of the bottom part of the dish		





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Label information	 Product name: SDA + LTHT 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 (Pink Plates)	 Incubations in vent and closed position possible Specific design to improve binding of agar to plate Easy handling due to increased handling area SDA plates are produced in pink dishes for better differentiation from TSA plates 	
Locking lid	 Locking-lid contact plate, made from polystyrene Inner diameter: 56.5 mm, thus providing an area of about 25 cm² Outer diameter: 67.5 mm Bottom part with 1 cm² square grid for facilitated evaluation 	
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: Vent position Closed position For long incubation of aerobic microorganisms, the closed position is recommended 	
Aerobic incubation (Closed position)	 Turn the lid clockwise to the right to the end into the final stop position The lid locks in the closed position Ideal incubation condition for aerobic micro-organisms Limits the dehydration of the agar during incubation 	



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Anaerobic incubation (Vent Position)	 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 1. First option: Turn the lid clockwise to the right to the end into the final stop position Turn the lid one click counter-clock-wise to the vent position 2. Second option: Turn the lid clockwise directly into the first locked position
Place of production	PharmaMedia Dr. Müller GmbH Gustav-Throm-Str. 1, 69181 Leimen - Germany

	Quality control, Certificates				
	Each lot of produ	ct can be obtaine	ed with a cer	tificate of ar	nalysis (CoA):
	Physico-chemical test parameters:				
	Appearance	Slightly turbid,	yellowish		
	pH value	5,4 - 5,8			
Opertification	Filling volume	16 – 19 mL			
Certificates	Irradiation	9-20 kGy			
	Growth Promo	tion test: 10-100	CFU		
	C. albicans	ATCC 10231	20-25 °C	2-3 days	50-200%
	A. brasiliensis	ATCC 16404	20-25 °C	3-5 days	50-200%
	Sterility contro				No growth
Certificate of origin	Origin (CoO). • Raw material • Tissue • Animal source • Country of orig		d raw materi	als are spec	cified as follows:
BSE policy	transmitting an medicinal proc specified anim We neither sto infectivity tissu	nimal spongiform ducts, we check al source, the co ore or process ru ues (IA) nor rum	encephalop the CoO of ountry of orig uminant raw inant raw m	bathy via hu raw materia in and the in materials o aterials who	nimizing the risk of iman or veterinary al in respect to the nfectivity category. obtained from high ose animal source ed risk (cat C/GBR



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	Quality control, Certificates
Temperature stress	 Art. 120.0060 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 at least 30 days after the assigned expiry date. Shelf-life testing comprise all regular tests which are part of the normal release test of this article (see CoA).
Neutralization of residues of disinfectants	The inactivation of residues of disinfectants is critical for the detection of viable and cultivable microorganisms in pharmaceutical production environments. For this purpose, different neutralizer combinations are added to the medium used for environmental monitoring. Most commercially available media contain Lecithin, Tween 80, Histidine and Thiosulfate. However, other neutralizers like Saponin, Cysteine and Glycine may be used as well. The composition as well as the concentration of single components are crucial for an effective detection of the residuals of disinfectants and therefore for the effective detection of microorganisms. The addition of different neutralizing components and concentrations to media has to be evaluated thoroughly. Besides the inactivation of residues of disinfectants neutralizers may have an inhibiting effect on the growth of microorganisms if used in higher concentrations thus making the detection of certain microorganisms difficult to impossible. Today most media used for environmental monitoring are using at least Lecithin and Tween in more or less identical concentrations: $-$ Lecithin: 0,7 g/L $-$ Tween: 5 g/L $-$ Tween: 5 g/L $-$ Tween: 5 g/L $-$ Tween: 5 g/L $-$ Na-Thiosulfate: 0,05 to 0,5 g/L $-$ 10 (sinfectant was used $-$ 10 (disinfectant sugness the explaied to a contact plate of about 25 cm ² 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. Test organisms B. <i>spizzenii</i> ATCC 6633, S. <i>aureus</i> ATCC 6538 and S. <i>epidermidis</i> ATCC 14990 as well as <i>E.coli</i> ATCC 8739, <i>P. paraeruginosa</i> ATCC 9027, <i>C. albicans</i> ATCC10231 and <i>A. brasiliensis</i> ATCC 16404. As reference, plates without disinfectant were inoculated with the test strains. Spe



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Quality control, Certificates
 TSA plates w. LTHT (Artcode 100.0100) were able to inactivate the following groups of disinfectant: Alcohols (ethanol, propanol, iso-propanol) Hydrogen peroxide (Biocide C) Peracetic acids (Incidin active2%, Perform sterile PAA) Mg-peroxyphtalate (Dismozon 4%) K-peroxymonosulfate (Perfom con. OXY 1%) Aldehydes like Glutaraldehyd, Formaldehyde (Aldasan 4%) Combinations of alcohol, hydrogen peroxide and peracetic acid (Actril) Combinations of alcohol, hydrogen peroxide and peracetic acid (Actril) Combinations of aldehydes + alcohols (Aerodesin 2000, Bacillol Plus) However, TSA plates w. LTHT were only able to inactivate quite low concentrations of quaternary ammonium compounds, biguanides and benzalkonium chloride. As these components are normally used in higher concentrations in disinfectants, they do not degrade by themselves and they are not volatile, it is required to clean such surfaces after disinfection with sterile water or sterile alcohol. Whereas the cleaning/rinsing may work properly on flat surfaces it seems likely that on other surfaces residues may remain or eventually even may be concentrated.

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



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