

	Technical Data Sheet			
Use in	<ul style="list-style-type: none">Pharmaceutical industryFor industrial, laboratory & research applications onlyBasic medium according to EP 2.6.1 and USP <71>			
Use for	<ul style="list-style-type: none">Sterility testIdentification and growth of fastidious anaerobic micro-organisms as well as aerobic micro-organisms			
Typical composition per liter	Casein peptone	15 g	Glucose-D(+) x H ₂ O	5.5 g
	Yeast extract	5 g	Na-Thioglycolate	0.5 g
	NaCl	2.5 g	Resazurine	1 mg
	Agar/ Gel Agent	0.2 g	L-Cysteine HCl	0.5 g
	This medium can be adjusted / or supplemented according to the performance criteria required.			
Filling volume	<ul style="list-style-type: none">200 mL			
Bottle format	<ul style="list-style-type: none">220 mL screw capType II glassBottle opening about 31 mmColour of cap: blueGL40 screw cap with 2 integrated septa			
Bottles per tray	<ul style="list-style-type: none">12 bottles on a plastic tray wrapped with shrink foil			
Shelf life	<ul style="list-style-type: none">8 months from production date			
Storage conditions	<ul style="list-style-type: none">Recommended storage temperature: 2 - 25 °CShould be stored at temperatures as stable as possibleStore protected from light exposure			
Label	<ul style="list-style-type: none">On the sideContain autoclave indicator			
Label information	<ul style="list-style-type: none">Product name: FTM clear 200 mLExpiry date: YYYYMMDD → MMM in letters (e.g.: 2023Nov04)Lot-numberIndividual numberBarcode			
Barcode	<ul style="list-style-type: none">2-dimensional (data matrix), 20 digits:Digits 1-3: Art.-No.Digits 4-9: Lot-NumberDigits 10-14: Individual-NumberDigits 15-20: Date (YYMMDD)			
Delivery	<ul style="list-style-type: none">Temperature controlled delivery on requestFor shipments of larger amounts plastic pallets in Euro-size can be used			

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Bottle information	<ul style="list-style-type: none"> Label contains autoclaving indicator (brown → green) Bottles are incubated at 25 – 35 °C for at least 48 hours after autoclaving and then packed Bottles are not touched any more by hand after autoclaving
Resazurine	<p>Resazurine is an indicator for anaerobic conditions. The indicator is colourless under anaerobic condition and turns into pink colour under aerobic conditions. Due to the composition of the medium 2 relatively stable zones can be clearly separated:</p> <ol style="list-style-type: none"> 1. an aerobic, pink zone on the top 2. an anaerobic, yellowish zone on the bottom. <p>Acc. to EP/USP not more than the upper third of the medium should be pink coloured. If a larger zone of the medium shows pink colour, the medium can be restored once by heating for a period not longer than 20 minutes. To avoid excess pressure the bottle should be vented with an aeration cannula during heating as well as cooling down phase</p>
Place of production	PharmaMedia Dr. Müller GmbH Gustav-Throm-Str. 1, 69181 Leimen - Germany

Quality control, Certificates					
Certificates	Each lot of product can be obtained with a certificate of analysis (CoA):				
	Physico-chemical test parameters:				
	Appearance	Clear, yellowish			
	pH value	6.9 – 7.3			
	Filling volume	196 – 208 mL			
	Growth Promotion test: 10-100 CFU				
	<i>S.aureus</i>	ATCC 6538	30 – 35 °C	≤ 3 days	Good growth
	<i>P.paraeruginosa</i>	ATCC 9027	30 – 35 °C	≤ 3 days	Good growth
	<i>C.sporogenes</i>	ATCC 11437	30 – 35 °C	≤ 3 days	Good growth
	<i>C.sporogenes</i>	ATCC 19404	30 – 35 °C	≤ 3 days	Good growth
	<i>C.acnes</i>	ATCC 11827	30 – 35 °C	≤ 5 days	Good growth
	Sterility control				
≥ 7 days at 30 – 35 °C, no growth					
Release of negative pressure in media bottles	During the autoclaving process, chemical reactions inside the bottles may result in a slight vacuum. Please assure that the vacuum is released without contaminating the bottle. Ideally, the vacuum is released by puncturing the septum with an aeration needle equipped with a sterile filter prior to opening a bottle.				

Quality control, Certificates	
Certificate of origin	<p>All media lots produced by PMM can be obtained with a Certificate of Origin (CoO). All animal derived raw materials are specified as follows:</p> <ul style="list-style-type: none"> • Raw material • Tissue • Animal source • Country of origin • Infectivity category (acc. to TSE guideline: EMA/410/01 current version)
BSE policy	<ul style="list-style-type: none"> • In compliance with the current note for guidance on minimizing the risk of transmitting animal spongiform encephalopathy via human or veterinary medicinal products, we check the CoO of raw material in respect to the specified animal source, the country of origin and the infectivity category. We neither store or process ruminant raw materials obtained from high infectivity tissues (IA) nor ruminant raw materials whose animal source originates from countries or regions with an undetermined risk (cat C/GBR IV).

Safety Data	
Toxic ingredients	<ul style="list-style-type: none"> • None
Basic composition	<ul style="list-style-type: none"> • See typical composition
Solvent content	<ul style="list-style-type: none"> • None
Safety data sheet required	<ul style="list-style-type: none"> • Not mandatorily required