| | Technical Data Sheet | | | |
|-------------------------------------|--|--|--|--|
| Use in | Pharmaceutical Industry Medical Device Industry Cosmetic Industry For industrial, laboratory & research applications only Basic medium according to EP 2.6.13 und USP <62> Test for specified micro-organisms | | | |
| Use for | Diluent for sample preparation | | | |
| Typical composition per liter | NaCl 4.3 g KH ₂ PO ₄ 3.6 g Na ₂ HPO ₄ x 2H ₂ O 7.2 g Caseine peptone 1 g Lecithin 3 g Polysorbate 80 30 g Histidine 1 g This medium can be adjusted / or supplemented according to the performance criteria required. | | | |
| Filling volume | • 200 mL | | | |
| Bottle format | 220 mL screw cap Type II glass Bottle opening about 31 mm Colour of cap: blue GL40 screw cap with 2 integrated septa | | | |
| Bottles per tray | 12 bottles on a plastic tray wrapped with shrink foil | | | |
| Shelf life | 12 months from production date | | | |
| Storage conditions | Recommended storage temperature: 2 - 25 °C Should be stored at temperatures as stable as possible Store protected from light exposure | | | |
| Label | On the sideContain autoclave indicator | | | |
| Label information | Product name: NPB+LTH 200 mL 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) | | | |



| | Technical Data Sheet |
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| Delivery | Temperature controlled delivery on request For shipments of larger amounts plastic pallets in Euro-size can be used |
| Bottle information | 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 |
| Place of production | PharmaMedia Dr. Müller GmbH Gustav-Throm-Str. 1, 69181 Leimen - Germany |

| | Quality control, Certificates | | | | | |
|---|--|--|---------------------------|----------------------------|--|-------------|
| | Each lot of product can be obtained with a certificate of analysis (CoA): | | | | | |
| | Physico-chemic | al test parame | eters: | | | 1 |
| | Appearance | Clear to slightly turbid, slightly yellowish | | | | |
| Certificates | pH value | 6.8 – 7.2 | | | | |
| | Filling volume | 196 – 208 mL | | | | |
| | Growth Promotic | on test: 200-2 | ,000 CFU/mL | * | | |
| | S.aureus | ATCC 6538 | 20-25 °C | 1 hour ±15 min | no change in CFU number | ĺ |
| | E.coli | ATCC 8739 | 20-25 °C | 1 hour ±15 min | no change in CFU number | ĺ |
| | P.paraeruginosa | ATCC 9027 | 20-25 °C | 1 hour ±15 min | no change in CFU number | ĺ |
| | Sterility control | Sterility control | | | | |
| | ≥ 7 days at 30-35 | ≥ 7 days at 30-35 °C, no growth | | | | |
| | *In case of a di ensure that suf | | | | | |
| Release of negative pressure in media bottles | During the autocl result in a slight vaccontaminating the septum with an aca bottle. | acuum. Please bottle. Ideally | assure that the vacuun | the vacuun n is release | n is released with ed by puncturing | nout the |



| | Quality control, Certificates |
|---------------------------------|--|
| Aeration for direct inoculation | During the autoclaving process, the majority of the oxygen inside the bottles is consumed in a chemical reaction, thus resulting in a medium not suitable for direct inoculation of aerobic microorganisms. When using such bottle for the growth of aerobic microorganisms without opening, please assure that the bottle is aerated throughout the complete incubation process by an aeration needle equipped with a sterile filter. Additionally, air filtered through a sterile filter may be pressed into the bottle using a syringe. |
| | Example for aeration: For bottles containing larger volumes of medium, e.g. TSB in 500 or 1000 mL, puncture the bottle lid (stopper) by a cannula of at least 1.6 mm diameter equipped with a sterile filter. Equilibrate with the cannula for not less than three days at 20 to 25 °C prior to inoculation. |
| 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 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). |
| Temperature stress | Art. 574.B200 has not been exposed to a temperature stress study so far. However, Art. 572.B090 (90 mL filling of NPB + LTH) has been exposed to temperature stress conditions (3 days at 30-35 °C) and has passed shelf-life testing at least 30 days after the assigned expiry date. Shelf-life testing comprises all regular tests which are part of the normal release test of this article except for sterility control (see CoA). As the medium in art. 574.B200 is identical to the one in 572.B090 in respect to the composition, it can be assumed, that NPB + LTH in the 200 ml filling is insensitive to such temperature conditions as well. |

| | Safety Data |
|-------------------|-------------------------|
| Toxic ingredients | • None |
| Basic composition | See typical composition |



| | Safety Data |
|----------------------------|--------------------------|
| Solvent content | • None |
| Safety data sheet required | Not mandatorily required |