



**STANDARD OPERATING PROCEDURE**

<b>Department:</b> Microbiology	<b>SOP No.:</b>
<b>Title:</b> Preparation of Culture Media Plates and Slants	<b>Effective Date:</b>
<b>Supersedes:</b> Nil	<b>Review Date:</b>
<b>Issue Date:</b>	<b>Page No.:</b>

**1.0 OBJECTIVE:**

To lay down a procedure for Preparation of culture media plates and slants.

**2.0 SCOPE:**

This SOP is applicable for Preparation of culture media plates and slants in Microbiology Laboratory

**3.0 RESPONSIBILITY:**

Officer / Executive - Microbiology

**4.0 ACCOUNTABILITY:**

Head QC

**5.0 ABBREVIATION:**

°C	Degree centigrade
IPA	Isopropyl Alcohol
LAF	Laminar Air Flow
Ltd.	Limited
ML	Microbiology Laboratory
No.	Number
Pvt.	Private
QA	Quality Assurance
USP	United State Pharmacopeia
SOP	Standard Operating Procedure

**6.0 PROCEDURE:**

**6.1 PREPARATION OF MEDIA PLATES:**

- 6.1.1** Prepare the required quantity of media as per SOP of "Receipt, Approval and Preparation of Culture Media" SOP.
- 6.1.2** Heat to dissolve the media if required, during heating shake the media regularly to avoid the overheating.
- 6.1.3** After heating dispense the media as per requirement in flasks/Bottles.
- 6.1.4** Keep the media flasks/ Bottles in autoclave and sterilize the slants as per validated time and temperature.
- 6.1.5** After sterilization cycle completion takes out the media from autoclave.
- 6.1.6** The sterilized Agar media and sterilized petriplates kept in pass box provided to concern area.



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**6.1.7** Enter in to the concern area and clean the LAF bench with 0.22 $\mu$  filtered 70% IPA.

**6.1.8** Pour approx 20-25 ml agar media (cool up to 45°C) in each petriplate and allow to solidifying.

**6.1.9 Pre-incubation:** Pre incubate the agar media plates at 30-35°C for 24 -48 hours.

**6.2 PREPARATION OF SLANTS:**

**6.2.1** Prepare the required quantity of media as per SOP of “Receipt, Approval and Preparation of Culture Media” SOP.

**6.2.2** Heat to boil the medium and mix properly.

**6.2.3** Dispense about 10-12 ml agar media in glass test tubes and plugged with cotton/cap.

**6.2.4** Keep the slant tubes in autoclave and sterilize the slants as per validated time and temperature.

**6.2.5** After completion of sterilization cycle take out the slants from autoclave and allows solidifying by tilting it at about 30°C and placing in incubator for pre incubation at 30-35°C for 24-48 hours.

**7.0 ANNEXURES:**

Not Applicable

**ENCLOSURES:** SOP Training Record

**8.0 DISTRIBUTION:**

- Controlled Copy No. 01                      Quality Assurance
- Controlled Copy No. 02                      Microbiology Laboratory
- Master Copy                                      Quality Assurance

**9.0 REFERENCES:**

USP Chapter <1117> Microbiological Best Laboratory Practices

**10.0 REVISION HISTORY:**

**CHANGE HISTORY LOG**

Revision No.	Change Control No.	Details of Changes	Reason for Change	Effective Date	Updated By