













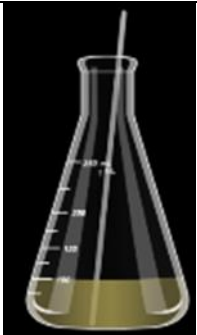



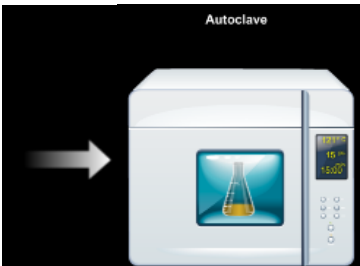


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|--|--|--|----|--|------------|-----------|---|-----------------------------------|
| <u>FLOW SHEET</u> # | 01 | SCENARIO # | 01 | DATE | 11/02/2016 | PERSONNEL | Svitlana Hnatush, Yaryna Kolisnyk, Andriy Halushka, Julia Turchyn | |
| TITLE | Isolation and identification of Candida sp. from the oral cavity. | | | | | | | |
| SOURCES/ REFS | | | | | | | | |
| VIDEO TIME (min:s) | ACTION | OBJECTS | | PHOTOS OF OBJECTS | | | POSSIBLE ERRORS | NOTES/ SCREENTEXT /GRAPHICS |
| SCENE # | 1 | PREPARATION OF NUTRITION MEDIA | | | | | | |
| 0:00 | Introduction: health & safety, principles and methods. Gather supplies | Avatar wearing lab coat & gloves. Materials and equipment: Work bench | |  | | | Failure to wear lab coat and/or gloves. | Links to codes of practice: |




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| | | <p>The jar/bottle of nutrient agar media</p> <p>Measuring cylinder</p> <p>Bottle of distilled water</p> <p>Neochlore disinfecting solution</p> |     | | |
|--|--|--|---|--|--|



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|--|--|-------------------------------|--|--|--|
| | | Electronic laboratory balance |  | | |
| | | Erlenmeyer flask |  | | |
| | | Glass stirring rod |  | | |
| | | Chemical spoon |  | | |

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|--|--|---|--|--|--|
| | | <p>Sterile, empty Petri plates</p>  | | | |
| | | <p>Bunsen burner</p>  | | | |
| | | <p>Autoclave</p>  | | | |

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| | Avatar measures distilled water into clean flask | | |  | | Failure to measure the correct amount of water | |
| | Avatar measures the components of the medium into a flask containing 9/10 volume of the solvent. Uses a clean chemical spoon for every measurement | | |  | | Failure to measure the correct amount of the medium | |
| | Avatar dissolves the solid components and fills with the remaining solvent up to final volume | | |  | | Failure to completely dissolve components of the medium | |

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| | Avatar closes the flask with cotton plug and covers with aluminium foil, puts into the autoclave and starts a sterilisation cycle | | |   | Failure to sterilize correctly | |
| 0:19 | Avatar prepares box to the work | washes surfaces with Neochlore disinfecting solution, turns on the UV lamp for 20 min | |  | Omitting preparation a suitable work area Failure to allow sufficient time for sterilizing box Forget to turn off the lamp before entering the box | |
| | Avatar cools flask with the media about 20-30 minutes | | |  | | If the medium is too cool, it will start to solidify into plates. If it is too hot, it will leave excess condensation on the lids. |

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| 0:30 | Avatar adjusts Bunsen burner | | |  | | Failure to adjust bunsen to give blue zone | The neck of the flask is not sterilized and contamination occurs |
| 0:57 | Avatar takes out the cotton plug and flames the neck of the flask over a Bunsen burner | | | | | | |
| 1:07 | Avatar pours the medium into sterile, empty Petri plates | (15-20 mL into each Petri plate) | |  | | розливає середовище далі, ніж за 15 см від полум'я; наливає в чашку замало середовища (не покрито дно); | |
| 1:31 | Avatar closes the flask with cotton plug | | | | | | |
| 1:37 | Avatar keeps the plates with medium on table to solidify | Turn dishes upside-down and stack them up for storage | |  | | перевертає або переносить чашку до того, як застигло | |

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|------|---|---|--|---|-------------|--|
| | | | | | середовище. | |
| 1:43 | Avatar extinguishes the Bunsen burner | | | | | |
| | After several hours avatar labels the plates according to the type of the medium and the date | In case of longer storage, Petri plates must be placed into plastic bags or boxes to avoid drying out | |  | | Do not store medium containing agar-agar under 4-5°C as it destroys its structure! |
| | Avatar performs a sterility test: | incubate the test plates at 28°C for 24 hours, and check for sterility | |  | | |

QUESTIONS FOR SCENARIO

(Establish the correspondence between the names of media and their description:)

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|-----------------------------------|---|
| 1. Differential-diagnostic | a) enable the fast growth of certain group of microorganisms with common physiological properties |
| 2. Transportation | b) contain components that inhibit the growth of some microorganisms, but do not have the influence on another |
| 3. Elective | c) contain conserving compounds and are used to prevent the death of microorganisms during the transportation of the studied material to the microbiological laboratory |
| 4. Selective | d) are used for the determination of fermentative properties of microorganisms, which is important during their biochemical identification |
| | e) are used for the accumulation of sufficient amount of cells for the analysis |

Establish the correct sequence of nutrition media preparation stages (put the photos in the correct order) :

