

Aug 24, 2024

Solid Growth Medium - Yeast

DOI

<https://dx.doi.org/10.17504/protocols.io.j8nlk85x5l5r/v1>



Mathias Hammer¹, Ammeret Rossouw¹, Azra Lari², Ben Montpetit³, David Grunwald¹

¹UMass Chan Medical School, RNA Therapeutics Institute, Worcester, MA, USA;

²University of Alberta, Department of Cell Biology, Edmonton, AB, Canada;

³University of California, Department of Viticulture and Enology, Davis, CA, USA



Mathias Hammer

UMASS Chan Medical School/RTI

Create & collaborate more with a free account

Edit and publish protocols, collaborate in communities, share insights through comments, and track progress with run records.

Create free account

OPEN  ACCESS



DOI: <https://dx.doi.org/10.17504/protocols.io.j8nlk85x5l5r/v1>

Protocol Citation: Mathias Hammer, Ammeret Rossouw, Azra Lari, Ben Montpetit, David Grunwald 2024. Solid Growth Medium - Yeast. protocols.io <https://dx.doi.org/10.17504/protocols.io.j8nlk85x5l5r/v1>

License: This is an open access protocol distributed under the terms of the **[Creative Commons Attribution License](#)**, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited

Protocol status: Working

We use this protocol and it's working

Created: April 06, 2024

Last Modified: August 24, 2024

Protocol Integer ID: 97849

Keywords: yeast grow medium, yeast solid grow medium, solid culture medium for saccharomyces cerevisiae, solid culture medium, solid growth medium, saccharomyces cerevisiae, yeast this protocol, yeast

Funders Acknowledgements:

NSF

Grant ID: 1917206

Disclaimer

DISCLAIMER – FOR INFORMATIONAL PURPOSES ONLY; USE AT YOUR OWN RISK

The protocol content here is for informational purposes only and does not constitute legal, medical, clinical, or safety advice, or otherwise; content added to **[protocols.io](#)** is not peer reviewed and may not have undergone a formal approval of any kind. Information presented in this protocol should not substitute for independent professional judgment, advice, diagnosis, or treatment. Any action you take or refrain from taking using or relying upon the information presented here is strictly at your own risk. You agree that neither the Company nor any of the authors, contributors, administrators, or anyone else associated with **[protocols.io](#)**, can be held responsible for your use of the information contained in or linked to this protocol or any of our Sites/Apps and Services.

Abstract

This protocol describes the steps to prepare solid culture medium for *Saccharomyces cerevisiae*.



Materials

Chemicals:

SC-Ura Powder

Sunrise Science Products

Cat#: 1306-030

Lot#: 23K3083

Exp: 10/2027

Yeast Nitrogen Base Without Amino Acids

Sigma Life Science

Cat#: Y0626-250G

Lot#: SLBG0555V

Glucose

Sunrise Science Products

Cat#: 1907-1kg

Lot#: 3A0036

Agar

Sunrise Science Products

Cat#: 1910-500

Lot#: 3B0104

Sodium Hydroxide

Fisher Scientific

Cat#: S318-500

Lot#: 130802

Deionized Water

Equipment:

500 ml laboratory bottle with screw cap

1ml pipette

50 ml pipette

stirring hot plate

magnetic stirring bar

micro scales

autoclave

thermometer

10cm polystyrene Petri dishes (20 pieces)



4°C fridge

Before start

Have the following solutions premixed:

Glucose 20% 500 ml solution:

Concentration: 200 g/l

mix 100 g Glucose in 500 ml deionized water (ddH₂O)

Sodium Hydroxide 1M solution 50 ml:

Concentration: 39.997 g/l

mix 1.99985 g NaOH into 50 ml ddH₂O

Optional:

SC-xx 10x 100ml solution:

Concentration: 19.2 g/l

mix 1.92 g into 100 ml ddH₂O

YNB 20x 100ml solution:

Concentration: 134.4 g/l

mix 13.44 g into 100 ml ddH₂O



1 Compound medium for autoclave

STEP CASE

Medium preparation with pre-resolved components

12 steps

This version of the protocol shows the preparation of the medium from SC-XX 10x and YNB 20x solutions.

1.1 Fill a 500 ml flask with  374 mL ddH₂O.

Add a magnetic stirring bar and place the flask on a stirring hot plate.

1.2 Add  25 mL YNB 20x solution (Yeast Nitrogen Base with Ammonium Sulfate without Amino Acids).

1.3 Add  50 mL SC-XX 10x solution.

Note

In regard to cover all optional dropout media the amino acid base holds the notification - xx, where xx stand for the amino acid(s) that is as selection factor, missing in the medium.

1.4 Add  10 g Agar.

1.5 Add  1 mL NaOH 1 molar solution.

Note

This is essential for the solidification of the medium!

2 Autoclave for  00:15:00 at  121 °C .

Note

Remove the stirring bar before going to autoclave.



3 Cooling and plating

3.1 Add a sterile stirring bar and thermometer. Let the medium stir to homogenize its temperature, while cooling.

3.2 When the medium cooled down to around  80 °C add  50 mL sterile Glucose 20%.

3.3 When the medium cooled down to  60 °C , plate  25 mL per  10 cm Petri dish (~20 dishes).

3.4 Let the medium solidify and cool down to  Room temperature .

4 Seal the prepared dishes in a plastic bag to prevent condensation and store them in  4 °C fridge.

Note

The agar plates can be store in the 4°C fridge for 2 to 3 months.