



Aug 03, 2022

Culturing Xanthomonas

Murray Grant¹, Shannon Greer¹, Joana Vicente²

¹University of Warwick; ²Fera Science

1 Works for me



dx.doi.org/10.17504/protocols.io.ewov1nr92gr2/v1

Xanthomonas genomics



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 <u>protocols.io</u> is not peer reviewed and may not have undergone a formal approval of any kind.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 <u>protocols.io</u>, 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 is the protocol for culturing strains of Xanthomonas species in preparation for extracting genomic DNA for sequencing.

DOI

dx.doi.org/10.17504/protocols.io.ewov1nr92gr2/v1

PROTOCOL CITATION

Murray Grant, Shannon Greer, Joana Vicente 2022. Culturing Xanthomonas. **protocols.io**

https://dx.doi.org/10.17504/protocols.io.ewov1nr92gr2/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

CREATED

Aug 03, 2022

LAST MODIFIED

Aug 03, 2022

PROTOCOL INTEGER ID

68145

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only and does not constitute legal, medical, clinical, or safety advice, or otherwise; content added to protocols.io is not peer reviewed and may not have undergone a formal approval of any kind.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.

- 1 Streak isolates onto King's B plates. Incubate at 28°C for ~48 hours.
- 2 Pick a single colony and inoculate to 10ml King's B liquid media in 30ml universal tubes. Incubate over night at 28°C and 220rpm.
- 3 Pellet 1.8ml of liquid culture in a 2ml Eppendorf tube by centrifuging at 5000g for 10mins. Pour off and discard the supernatant.
- 4 Pipette another 1.8ml culture into the tube and repeat step 3. This will pellet 4ml of culture in a 2ml Eppendorf tube.

protocols.io

2

5	Spin the tubes for 2mins at 5000g. Remove the supernatant by pipetting and discard.
6	Proceed to the extraction process or flash freeze the pellets in liquid nitrogen and store at -20C until use.