



Growth of mixed *E. coli* colonies V.2

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Version 2 ▾

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1 Works for me



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ABSTRACT

Growth of mixed *E. coli* colonies on agar plates and between an agar plate and an agar pad

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Make plates

- 1 Autoclave Lennox LB (10 g/l tryptone / casein digest peptone, 5 g/l NaCl, 5 g/l yeast extract) with 1.5 % agar (w/v).
- 2 **Optional:** Add antibiotics at desired concentration after medium has cooled down sufficiently.
- 3 Pipette **10 mL** of LB and agar in Petri dishes with diameter of **6 cm** .
- 4 Leave plates plates to dry **Overnight** at **Room temperature** . Optionally enclose to avoid nonuniform drying.

Store and prepare plates

- 5 If not used the day after pouring plates, store at 4°C . Enclose in plastic container or bag to avoid further drying of plates.
- 6 Before usage of plates that have been refrigerated, warm up plates at 37°C .

Prepare bacterial cultures

- 7 Grow overnight culture (colony picked or directly from glycerol frozen stocks) in Lennox LB.

Inoculate colonies and incubate

- 8 Mix $900\ \mu\text{l}$ Lennox LB and $50\ \mu\text{l}$ of each of the two overnight cultures in an Eppendorf tube. Vortex.
- 9 Inoculate $1\ \mu\text{l}$ in centre of plate and let dry.
- 10 **Optional:** Cover colonies with agar pad. Cut about $18\ \text{mm} \times 18\ \text{mm}$ agar pad inside plate. Use spatula to lift pad. Place vertically next to colony and let fall upside down onto colony. Use spatula to remove bubbles by pressing on top.
- 11 Place plates into container, together with sufficiently wet paper towels. Incubate for 7-8 days in a dark environment.