

Antibiotics gradient assay for V. natriegens

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Abstract

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Protocol

Step 1.

Inoculate preculture of *V. natriegens* in LB3 medium (3% NaCl) and incubate overnight at 37°C, shaking.

Preparation of gradient plates

Step 2.

Preparation of the first layer:

Place 11.5x11.5 cm plates in an inclient position. The angle of inclination is such that the agar layer deminishes to nothing at one edge of the plate. Pour 30 ml of LB3 with chosen antibiotics of concentrations to be tested into the plate. Let the agar solidify.

Preparation of gradient plates

Step 3.

Switch position of the plate to an even surface and pour 30 ml of LB3 agar **without** any antibiotics to onto the first layer. Let the agar solidify.

Spotting of V. natriegens

Step 4.

Diltute the preculture to an OD550 of 0.1.

Spotting of V. natriegens

Step 5.

Pipette 8-10 spots of 5 μl of the culture along the gradient.

Spotting of V. natriegens

Step 6.

Let the spots dry and incubate the plates upside-down at 37°C over night or at room temperature over the weekend.

Measurement

Step 7.

Measure the distance from the edge of the agar plate to the last point of growth.

Measurement

Step 8.

Calculate the highest concentration were *V. natriegens* can survive by using the following equation:

$$c_{antibioticH} = c_{antibioticmax} * d_{growth}/d_{plate}$$

c_{antibioticH}: highest antibiotic concentration *V. natriegens* can survive

 $c_{\mbox{\scriptsize antibioticmax}}$: maximum atibiotics concentration (concentration used for the lower layer of the plate)

 $d_{\text{growth}}\!\!:$ distance from the edge of the agarplate to the last point of growth $d_{\text{plate}}\!\!:$ length of the plate (11.5 cm)