

# Cellulophaga growth reading

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## Abstract

For [One-step growth curves for Cellulophaga phages](#) protocol and [Transcriptomics During One-Step Growth Curves for Cellulophaga Phages](#) protocol.

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## Protocol

### Step 1.

Pipet 200 µl of MLB media into wells A1 and A2 of a white microtiter plate

#### 📌 NOTES

**VERVE Team** 24 Aug 2015

This is your 'blank'.

#### ■ ANNOTATIONS

**Bonnie Poulos** 15 Mar 2016

Make sure the microtiter plate you are using is clean inside and out, with no scratches or spots on its surface, as it will interfere with the light reading.

**Bonnie Poulos** 15 Mar 2016

For a determination of growth, an absorbance reading at 595nm will be taken of the culture.

### Step 2.

Pipet 200 µl of sample (the new culture you just inoculated) into wells B1 and B2 of the same plate

#### 📌 NOTES

**VERVE Team** 24 Aug 2015

Ensure that there are no bubbles in the wells, as they will affect your readings. Pipet away any bubbles.

### Step 3.

Read the plate on the plate reader

#### ■ ANNOTATIONS

**Bonnie Poulos** 15 Mar 2016

Take absorbance reading at 595nm.