

## **Culturomics using microfluidics**

**Keywords** : Microfluidic, culture

### **Context**

Sequencing does not completely replaces culture approaches, which are still laying the basis needed when it comes to interpret sequences. However it is very time-consuming.

Here we aim to use the work of a postdoc working in microfluidics to generate hundreds of droplets containing a single bacterial cell, effectively generating hundreds of micro-cultures.

### **Aims of the master student**

With the help of other INRA partners, the planning for the master student is the following :

- 1. Setup the microfluidic device in the NED team*
- 2. Generate 30 um droplets containing a single bacterial cell*
- 3. Try 5 different media and count the droplets displaying growth as an indicator of the percentage of cultivable cells through this technique*
- 4. Sequence the growing cells using the Illumina technology and the pipelines available in the lab (no prior bioinformatics knowledge required)*

Once the proof of concept achieved (estimated time=3 months), the student will continue the work by one or more of the following :

- automatisation with a pipetting robot for high throughput
- check for antibiotic resistance

### **Abilities**

The student should understand english and be highly motivated. Previous knowledge about handling microbial communities would be appreciated but is not necessary.

### **Contact for more information**

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### **Practical details**

- 6 months from Jan to Jun 2016
- INRA Toulouse – UMR GenPhySE à Auzeville (31)
- paid about 550 €/mois – (lunch available on site for ~ 2.50 eur)