

<b>iGEM 2016 – SDU</b>	
<b>Title:</b> 2xYT media	<b>Date issued:</b> 2016/08/25
<b>SOP number:</b> SOP0030	<b>Review date:</b> 11/10/2016
<b>Version number:</b> 01	<b>Written by:</b> Joel Vej-Nielsen & Jakob Rønning

## 1. Purpose

Media for growth of cell cultures with high cell density and a longer growth period for *E. coli*

## 2. Area of application

This procedure is valid for all *E. coli* strains

### 3. Apparatus and equipment

Apparatus/equipment	Location (Room number)	Check points	Criteria for approval/rejection
Autoclave	Chemical room		
pH-meter	Chemical room		
Weight	Chemical room		
Weigh dishes			
Single use plastic			

### 4. Materials and reagents – their shelf life and risk labelling

Name	Components (Concentrations)	Manufacturer / Cat. #	Room	Safety considerations
Tryptone			Chemical room	
Yeast extract			Chemical room	
NaCl			Chemical room	
Distilled water				
1 L media bottle			Micro storage	

**5. QC – Quality Control**

**6. List of other SOPs relevant to this SOP**

SOP0001 – ON culture

**7. Environmental conditions required**

**8. Procedure**

1. Measure ~900ml of distilled H<sub>2</sub>O and put in flask 1L.
2. Add 16 g Tryptone, 10 g yeast extract, 5 g NaCl and mix with distilled H<sub>2</sub>O.
3. Adjust pH to 7.0 with 10 M NaOH (~ 200 µL).
4. Adjust volume to 1 liter with Distilled H<sub>2</sub>O.
5. Sterilize by autoclaving and store at room temperature.

## 9. Waste handling

Chemical name	Concentration	Type of waste (C, Z...)	Remarks
Once use plastic			
Weigh dishes			

## 10. Time consumption

- Total-time 3 h.
- Hands-on-time 30 min.

## 11. Scheme of development

Date / Initials	Version No.	Description of changes
16.08.25 / JVN & JR	01	The SOP has been written
16.10.12 / JVN & JR		reviewed

## 12. Appendixes