iGEM2014 – Microbiology – BMB – SDU	
Title: Ligation	Date issued: 2013.06.19
SOP number: SOP0015_v01	Review date: 2015.09.13
Version number: 01	Written by: ASF

1. Purpose

To ligate pieces of DNA

2. Area of application

Cloning

3. Apparatus and equipment

Apparatus/equipme nt	Location (Room number)	Check points	Criteria for approval/rejection
Vortex	Laboratory 1. Floor	•	
Pipettes (p20, p10)	Micro Storage	•	
		•	
		•	
		•	
		•	
		•	

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

Name	Componen	Supplier / Cat. #	Room (hallway	Safety
	ts		storage)	considerations
Purple pipette		Contact	Micro storage	
tips		lab-manager		
Green pipette		Contact	Micro storage	
tips		lab-manager		
Eppendorftube		Contact	Micro storage	
S		lab-manager		
Distilled water		Contact	Micro storage	
		lab-manager		
Ligasebuffer		Agilent	Freezer at 1. Floor	
		Technologies		
Ligase			Freezer 1. Floor	

DNA piece 1		Refrigiator 1. Floor	
DNA piece 2		Refrigiator 1. Floor	

5. QC – Quality Control

6. List of other SOPs relevant to this SOP

7. Environmental conditions required

8. Procedure

- 1. Prepare the ligation mixture and mix by pipetting up and down
- 2. Leave the mixture overnight at 16°C
- 2a. If there is no time leave the ligation solution at 22.5°C for 30 mins. Then denature the ligase at 65°C for 10min.
- 3. Use ligation solution for transformations

Reagents	Volume
10x T4 DNA ligase buffer	2 μL
T4 DNA ligase (add last!)	1 μL
PCR product (cut) of each brick which is to be ligated – or 1 part plasmid and 5 part bricks	5 μL or 10 fmol Plasmid, 0, 10 and 20 fmol PCR
H2O	to reach a total volume of $20\mu L$

9. Waste handling

Chemical name	Concentration	Type of waste (C, Z)	Remarks
One use Plastic		GMO	Yellow GMO Trash

10. Time consumption

- 3 Hours
- 1 Hour + Ligation overnight

11. Scheme of development

Date / Initials	Version	Description of changes
	No.	
13.06.19 / ASF	01	The SOP has been written
13.06.26 /PRA	01	The SOP has been approved
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12. Appendixes