

Cloning in Asaia



Swiss Federal Institute of Technology Lausanne iGEM team 2010, Dupont Thibault, Gerweck Nadia, Grädel Nadine, Helfer Jonas, Lisowski Wiktor, Monnot Gwennaëlle, Perrudet Christian, Richter Solange, Varricchio Stefano, Vokali Efthymia, Déneraud Nicolas, Gubelmann Carine, Niederholtmeyer Henrike Marie, Opota Onya, Deplancke Bart, Maerkl Sebastian.

Introduction

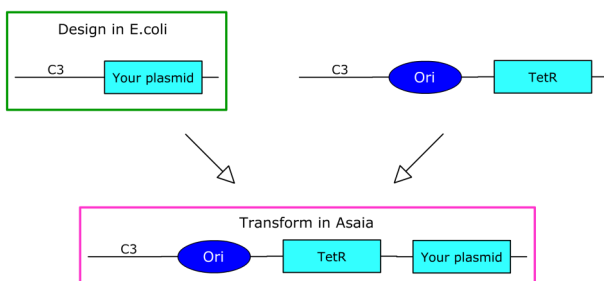
You can't do cloning with Asaia like in E.coli. You have to follow some rules. You can find those rules here.

This document contains the indispensable plasmid you need to work with Asaia. You will also find all the plasmids we have created until now.

We have given what we tried to do without success, so maybe you will manage to triumph in those tasks, good luck.

WORKING WITH ASAIA

We recommend you to work with E. coli when possible, as its doubling time is twice as short. Typically, design and create your plasmid in E. coli and then cut out the part you are interested in and ligate it into a plasmid containing the Asaia Origin. You can order C3+Asaia origin+TetR or C3+Asaia origin+KanR in the parts registry.



Once the cloning is done you can transform it into Asaia and E. coli and proceed with your experiments. We recommend you to make GLY stock with E.coli so, in the case you have to do a liquid culture to do mini-prep, it will take only one day and not 2 days.

Note that the Asaia origin is compatible with E. coli (though the inverse is not true.) You can therefore design a plasmid with the Asaia origin and work with it in E. coli.

ACQUISITION OF THOSE PARTS

If you want any of the parts mentioned below, you can just order it from the web site : <http://partsregistry.org>

PARTS WE CREATED

This section contains all the parts we designed and created, based on pSB1C3, followed by a short description. Details can be obtained by following the links associated to each plasmid description.

Each parts description are structured this way :

[BIOBRICKS ID] // [SEQUENCE IN BLOCK]

[Image of the plasmid]

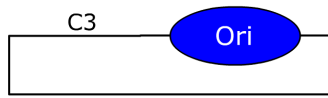
[Description of the plasmid]

[link to the registry]

Here are the abbreviations we used

Abbreviation	Full name
C3	pSB1C3
ori	Asaia origin
Strong	Strong promoter
KanR	Kanamycin Resistance
AmpR	Ampicillin Resistance
TetR	Tetracyclin Resistance
Immuno	Immunotoxin expression
p25	sequence expressing p25 protein
p28	sequence expressing p28 protein

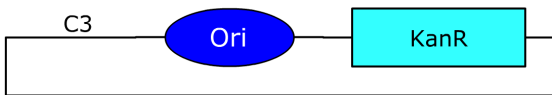
BBA_K320000 // C3+ORI



This plasmid only contain Asaia origin, you can use it to just cut out this origin.

[\[link to the registry\]](#)

BBA_K320004 // C3+ORI+KANR



This part is very useful when you want to adapt your plasmid for Asaia. Just cut this plasmid with SpeI and PstI and your plasmid with XbaI and PstI and ligate them to create your Asaia compatible plasmid.

[\[link to the registry\]](#)

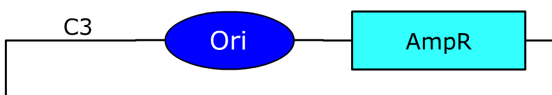
BBA_K320003 // C3+ORI+TETR



This part is very useful when you want to adapt your plasmid for Asaia. Just cut this plasmid with SpeI and PstI and your plasmid with XbaI and PstI and ligate them to create your Asaia compatible plasmid.

[\[link to the registry\]](#)

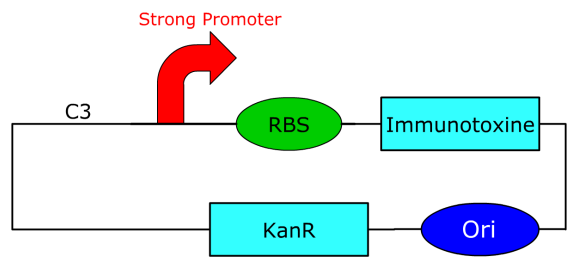
BBA_K320011 // C3+ORI+AMPR



This part is very useful when you want to adapt your plasmid for Asaia. Just cut this plasmid with SpeI and PstI and your plasmid with XbaI and PstI and ligate them to create your Asaia compatible plasmid. Caution : read the "Growing Asaia" sheet to see that Asaia is naturally resistant to Ampicillin

[\[link to the registry\]](#)

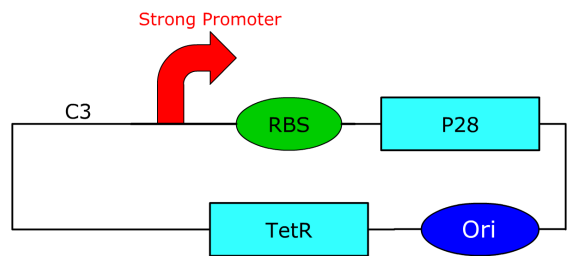
BBA_K320008 // C3+STRONG+IMMUNO+ORI+KANR



This plasmid is an Asaia compatible plasmid expressing the immunotoxine. It was one aim of our project. You will found more information on the web site : http://2010.igem.org/Team:EPF_Lausanne

[\[link to the registry\]](#)

BBA_K320009 // C3+STRONG+P28+ORI+TETR



This plasmid is an Asaia compatible plasmid expressing the p28 protein. It was another aim of our project. You will found more information on the web site : http://2010.igem.org/Team:EPF_Lausanne

[\[link to the registry\]](#)