

Biological system reduces amount of freight for delivery to orbit

1 July 2016, by Steven Burgess

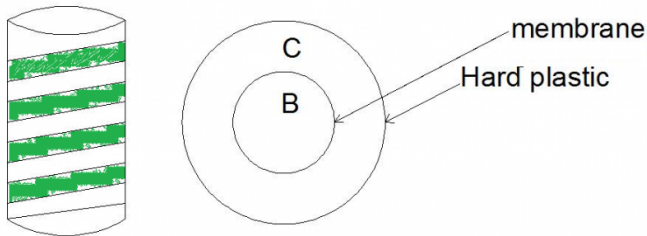


Figure 1: Overview of the co-culture system used in the project. It will have a design that allows for a high modularity, with the cyanobacterium found in compartment C and *B. subtilis* in compartment B. Compartment should then be de-attached from the system and attached again to change the *B. subtilis*. The membrane will most likely be a dialysis bag.

Ever wondered why humans have not colonized Mars, or traveled further into space? We have. It is not because the technology is lacking – in fact, using the technology we have today it might be possible to achieve these aims (Wall 2013). The main limiting factor is the cost. For instance, the Curiosity mission cost 2.8 billion US dollars. During the mission, a rover was sent to Mars to investigate the composition of the soil. The cost of this was not insanely high, but with a budget for space exploration that keeps being cut by governments of developed countries around the globe (Luxton 2016); it is hard to find the money for longer, manned missions in space.

One of the main costs is the external fuel tank (Giges 2014). The more you have to load on the [space](#) rocket, the more fuel it needs to launch, and the more the mission will cost. Therefore, it would be desirable to decrease the amount of fuel that is needed to have on the [space rocket](#) prior launching. [CosmoCrops](#) has a solution to this!

Engineered *Bacillus subtilis* for the rescue!

What we are working on this summer, is to make a biological system that can reduce the amount of material you need to have on board on the space rocket prior to launching. This is done by using two different organisms that will work together in a co-culture. The idea is that when the astronauts or settlers on other planets need food, drugs or any organic material, they can simply produce it themselves. The co-culture will be designed in such a way that it allows for a high degree of modularity (Fig. 1), allowing the astronauts or settlers to take out one component and add another engineered microorganism. This means that the co-culture system could be used to produce almost everything – as long as you have a microorganism that can produce it.

The two organisms we will use are the cyanobacterium *Synechococcus elongatus* and *Bacillus subtilis*. The cyanobacterium will be the driving force of the system, using sunlight and carbon dioxide to produce sucrose. This sucrose will then be secreted by the sucrose transporter *cscB* (Ducat et al. 2012). Hereafter, the secreted sucrose will be used by *B. subtilis* to produce whatever compound is desired. *B. subtilis* can form spores, meaning that prior to launching it will be possible to have different engineered forms of *Bacillus* spores on the shelf. Here, they can be stored for a long time without the need of any nutrition. When the astronauts/settlers need something, they find the corresponding engineered spores and put them into the co-culture system.

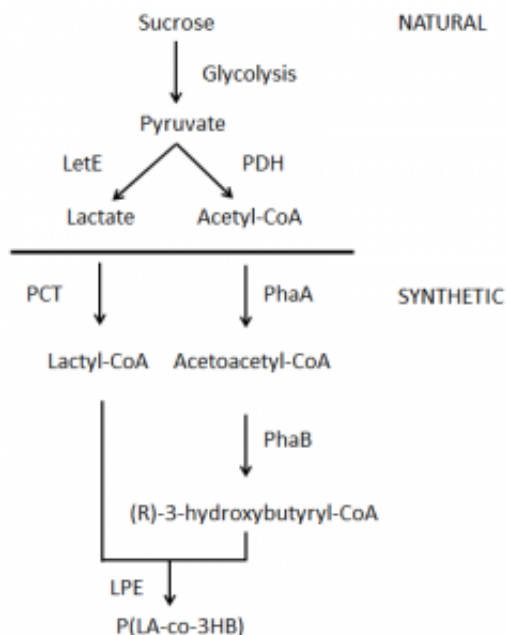


Figure 2: How *B. subtilis* will be engineered to produce P(LA-co-3HB), by introducing four new enzymes. The overview is divided in the “Natural” and “synthetic” part, where Natural is what is going on in the WT strain and Synthetic is what will be happening in the engineered strain that produces the bioplastic.

The Genetic Approach

As a proof of concept, we will engineer *B. subtilis* to produce the bioplastic P(LA-co-3HB) that can be used in the [3D printer currently on the International Space Station](#). To do this, we introduce four enzymes to *B. subtilis* (Taguchi et al. 2008; Jung et al. 2010; Yamada et al. 2010) (Fig. 2). These four enzymes are a propionate-CoA transferase (PCT) from *Clostridium propionicum*, β -ketothiolase, (PhaA) and acetoacetyl-CoA reductase (PhaB) both from *Ralstonia eutropha* and lactate polymerizing enzyme (LPE). LPE is originally a Pha synthase that has been mutated to use lactyl-CoA as a substrate (Taguchi et al. 2008). Together, these four genes introduce a new sink pathway for carbon. Because of this it might be necessary to make some genomic alterations to optimize the yield, which has been done in *Escherichia coli* (Jung et al. 2010).

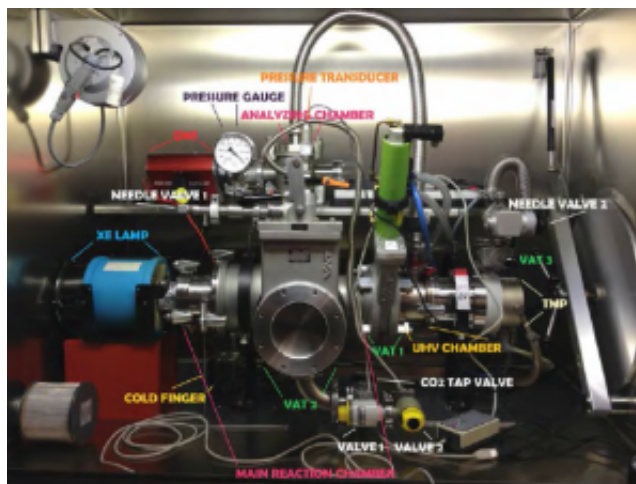
The Jens Martin Mars Chamber

Besides the genetic approach, one aspect of this project is also to test the *B. subtilis* strain in the

Jens Martin Mars Chamber (Fig. 3) at the Niels Bohr Institute. This is a chamber that can simulate the environment in space or on Mars by controlling factors such as temperature, pressure, UV radiation and the atmospheric composition (Kajtár 2014).

The chamber is essentially a doubly sealed environment (several airtight layers) with industrial vacuum pumps, UV-lamps, oven elements, Peltier panels and water cooling. All this allows for experiments in pressure as low as around the Moon, from Arctic temperature to several hundred degrees with or without UV radiation. For normal pressures, it also allows altered atmospheric compositions.

By testing our *B. subtilis* strain in this chamber, we hope to stimulate some mutations that can make the strain more viable in extreme environments as space or Mars. This will be done by inducing mutations in the genome of the bacterium by UV mutagenesis. If viable cells are observed after being tested for one or several of the parameters, one approach will be to test it more thoroughly with all the parameters, to see if it can survive. If the cells do survive then whole genome sequencing will be carried out and bioinformatics will be used to find the mutations that allow the cells to survive.



Future Perspectives

With our project, the future is brighter for space

exploration. When the co-culture system is functional, it can be used to produce not only bioplastic, but a variety of different compounds that can make space missions more feasible. Furthermore, it will make it cheaper to perform space missions, because there is no need to have so many materials on board on the rocket before launching, simply just the co-culture, spores of *B. subtilis* and a 3D printer – then everything else can be made out in space. It will also be used for future settlements on other planets, so they are not dependent on Earth for materials, but can produce what they need by themselves. Thereby, making the lives of new settlers easier than it otherwise would be.

The CosmoCrops

CosmoCrops is an interdisciplinary team from University of Copenhagen that consists of 10 undergraduate and graduate students. In June CosmoCrops won the Nordic iGEM Conference held in Stockholm, Sweden by having the best presentation and project among the Nordic teams. We can be followed on Facebook and Twitter if you want to stay tuned!

More information: Ducat D.C., Avelar-Rivas J.A., Way J.C. & Silvera P.A. (2012) Rerouting carbon flux to enhance photosynthetic productivity. *Applied and Environmental Microbiology* 78, 2660–2668.

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