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## CPEC Protocol

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**Protocol status:** Working

**We use these protocols in our group and they work.**

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- 1 Measure the DNA concentration (ng/ml) of each assembly piece.
- 2 Add 100 ng of the linearized vector backbone and equimolar amounts of the other assembly pieces to a 25 ml total volume assembly reaction mixture as follows:

linearized vector backbone (100 ng)  
+ each additional assembly piece (to equimolar with backbone)  
+ 5 ml 5X HF Phusion Reaction Buffer  
+ 1 ml 10 mM dNTPs  
+ 0.75 ml DMSO  
+ 0.5 ml 2U/ml Phusion Polymerase  
+ \_\_\_\_\_dH<sub>2</sub>O to  
25 ml

- 3 Perform the assembly reaction in a thermocycler as follows:

30 sec @ 98 C    1 cycle  
10 sec @ 98 C }  
30 sec @ 55 C }    1 to 15 cycle(s)\*\*  
length\* (kb) x 15 sec @ 72 C }  
10 min @ 72 C    1 cycle

\*The total length of the assembled product (in kb)

\*\*The number of repeated cycles should exceed the number of assembly pieces

Transform 5 ml of the assembly reaction into 100 ml of competent E. coli and/or run a diagnostic agarose gel to check for successful assembly.