**pCrU6.4-SaCas9 cloning/aphVIII (pPH339)**

Deposited by Andre Greiner, Peter Hegemann lab, Humboldt-University Berlin, October 2017

Cloning vector for *Staphylococcus aureus* Cas9 (SaCas9) guide RNA transcription, controlled by the U6 snRNA promoter #4 (3’ of [Cre15.g640800](https://phytozome.jgi.doe.gov/pz/portal.html#!gene?search=1&crown=1&detail=1&method=0&searchText=transcriptid:30783541)). The 20 bp target site can be inserted in a cut-ligation reaction as annealed oligos into an *Esp*3I cloning site following the protocol of Ann Ran ([Ran et al. 2013](https://doi.org/10.1038/nprot.2013.143)). The immediate 4-bp sequence upstream of the transcriptional start site was changed to ACTT in all U6 constructs to simplify the cloning procedures.

Vector contains an *aphVIII* cassette for selection on paromomycin.

Selection in *C. reinhardtii*: paromomycin

Selection in *E.coli*: ampicillin

host strain: DH5α

Sequence map: <https://benchling.com/s/seq-tw570ykR6MRUqx0FGeaF>

**Reference**

Greiner, A., Kelterborn, S., Evers, H., Kreimer, G., Sizova, I., and Hegemann, P.Targeting of photoreceptor genes via zinc-finger nucleases and CRISPR/Cas9 in *Chlamydomonas reinhardtii.* Plant Cell 30 (2017). <https://doi.org/10.1105/tpc.17.00659>

**Overview of all CRISPR/Cas9 plasmids from the Hegemann lab**

<https://www.chlamycollection.org/hegemann_lab>

Visit [www.chlamy.de](http://www.chlamy.de) for more info or contact [CRISPR@chlamy.de](mailto:CRISPR@chlamy.de)