# nature portfolio

Corresponding author(s):	Joseph Bondy-Denomy			
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### **Reporting Summary**

Nature Portfolio wishes to improve the reproducibility of the work that we publish. This form provides structure for consistency and transparency in reporting. For further information on Nature Portfolio policies, see our <u>Editorial Policies</u> and the <u>Editorial Policy Checklist</u>.

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FOr	all statistical analyses, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.
n/a	Confirmed
	The exact sample size (n) for each experimental group/condition, given as a discrete number and unit of measurement
	A statement on whether measurements were taken from distinct samples or whether the same sample was measured repeatedly
	The statistical test(s) used AND whether they are one- or two-sided Only common tests should be described solely by name; describe more complex techniques in the Methods section.
X	A description of all covariates tested
X	A description of any assumptions or corrections, such as tests of normality and adjustment for multiple comparisons
	A full description of the statistical parameters including central tendency (e.g. means) or other basic estimates (e.g. regression coefficient AND variation (e.g. standard deviation) or associated estimates of uncertainty (e.g. confidence intervals)
	For null hypothesis testing, the test statistic (e.g. <i>F</i> , <i>t</i> , <i>r</i> ) with confidence intervals, effect sizes, degrees of freedom and <i>P</i> value noted <i>Give P values as exact values whenever suitable.</i>
$\times$	For Bayesian analysis, information on the choice of priors and Markov chain Monte Carlo settings
$\boxtimes$	For hierarchical and complex designs, identification of the appropriate level for tests and full reporting of outcomes
	Estimates of effect sizes (e.g. Cohen's <i>d</i> , Pearson's <i>r</i> ), indicating how they were calculated
	Our web collection on statistics for biologists contains articles on many of the points above.

#### Software and code

Policy information about  $\underline{availability\ of\ computer\ code}$ 

Data collection

NIS-Elements AR 5.02.00 64-bit, Image Lab (BioRad) software version 6.0.1, BioTek LogPhase 600 plate reader version 1.08, MiSeq Control Software version 3.1.0.13.

Data analysis

NIS-Elements AR 5.02.00 64-bit, Python 3.8, Geneious Prime® 2021.2.2. The custom python codes are available from public repositories at https://zenodo.org/record/6324407#.YiAjrejMI2w

For manuscripts utilizing custom algorithms or software that are central to the research but not yet described in published literature, software must be made available to editors and reviewers. We strongly encourage code deposition in a community repository (e.g. GitHub). See the Nature Portfolio guidelines for submitting code & software for further information.

#### Data

Policy information about availability of data

All manuscripts must include a data availability statement. This statement should provide the following information, where applicable:

- Accession codes, unique identifiers, or web links for publicly available datasets
- A description of any restrictions on data availability
- For clinical datasets or third party data, please ensure that the statement adheres to our <u>policy</u>

All relevant data are included in the paper and/or the Supplementary/Source Data files. The complete genome sequence of OMKO1 was deposited in GenBank under accession number ON631220. All strains and plasmids are available from the corresponding author upon request.

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Please select the or	ne below that is the best fit for your research. If you are not sure, read the appropriate sections before making your selection.
Life sciences	Behavioural & social sciences Ecological, evolutionary & environmental sciences
For a reference copy of t	the document with all sections, see <u>nature.com/documents/nr-reporting-summary-flat.pdf</u>
Life scier	nces study design
All studies must dis	close on these points even when the disclosure is negative.
Sample size	Sample sizes refer to the number of at least 500 bacterial cells in the dataset of each microscopic experiment.
Data exclusions	No data were excluded for analysis.
Replication	Plaque assays and time-lapse microscopic experiments were independently performed at least twice. Microplate liquid assays were performed in three independent replicates. One-step growth curves were independently performed at least three times.
Randomization	Randomization is not relevant to this work. The goal of this work is to develop a Cas13a-mediated genetic tool to engineer bacteriophages. It was sufficient to include an appropriate internal control (e.g. expression of a non-targeting Cas13a crRNA) and compare the effects with the gene-editing samples. In this respect, randomization is not necessary as we were investigating the possibility of Cas13-based genome editing in different bacteriophage strains.
Blinding	Blinding was not relevant in this study, as the effects of genome editing were individually tested by PCR or sequencing for each engineered phage strains.

## Reporting for specific materials, systems and methods

We require information from authors about some types of materials, experimental systems and methods used in many studies. Here, indicate whether each material, system or method listed is relevant to your study. If you are not sure if a list item applies to your research, read the appropriate section before selecting a response.

Materials & experimental systems		Methods		
n/a	Involved in the study	n/a	Involved in the study	
$\boxtimes$	Antibodies	$\boxtimes$	ChIP-seq	
$\boxtimes$	Eukaryotic cell lines	X	Flow cytometry	
$\boxtimes$	Palaeontology and archaeology	$\boxtimes$	MRI-based neuroimaging	
$\boxtimes$	Animals and other organisms			
$\boxtimes$	Human research participants			
$\boxtimes$	Clinical data			
$\boxtimes$	Dual use research of concern			