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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.					
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n/a	Cor	nfirmed			
	\boxtimes	The exact sample size (n) for each experimental group/condition, given as a discrete number and unit of measurement			
	\boxtimes	A statement on whether measurements were taken from distinct samples or whether the same sample was measured repeatedly			
	\boxtimes	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.			
\boxtimes		A description of all covariates tested			
\boxtimes		A description of any assumptions or corrections, such as tests of normality and adjustment for multiple comparisons			
	\boxtimes	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)			
	\boxtimes	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>			
\boxtimes		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			
\boxtimes		Estimates of effect sizes (e.g. Cohen's d, Pearson's r), indicating how they were calculated			

Our web collection on <u>statistics for biologists</u> contains articles on many of the points above.

Software and code

Policy information about availability of computer code

Data collection

Commercial softwares licensed by material-characterization companies were utilized, including SmartLab Studio (Rigaku SmartLab), Fluoracle (FLS980, Edinburgh), FESEM system (TESCAN MAIA3), GO Optical system (STOmics microscope), Image Lab (Biorad) and HPLC 1260 Infinity II system (Agilent).

Data analysis

HPLC 1260 Infinity II software (Agilent) was used to analyze yield and step efficiency for synthesized oligonucleotides. Imagel 1.52v was used for size measurement in SEM images. Data is processed and visualized by OriginPro 9 (9.0.0). The raw sequencing data from each library were initially processed using fastp (v0.23.2) for quality control, followed by alignment of the high-quality reads to their respective reference sequences through bwa mem (v0.7.17-r1188) with parameter -B 2. Codon frequency quantification was performed through alignment analysis at all NNK-encoded positions.

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 policy

The DNA sequencing data of variant libraries that support the findings of this study have been deposited in the CNSA (CNGB Nucleotide Sequence Archive) under accession number CNP0007102. All data supporting the findings of this study are available within the manuscript file and its Supplementary Information files.

Research involving human participants, their data, or biological material

•	studies with <u>human participants or human data</u> . See also policy information about <u>sex, gender (identity/presentation),</u> d race, ethnicity and racism.					
Reporting on sex and ge	ender N/A					
Reporting on race, ethn other socially relevant groupings	icity, or N/A					
Population characteristi	cs N/A					
Recruitment	N/A					
Ethics oversight	N/A					
Note that full information on the approval of the study protocol must also be provided in the manuscript.						
Field-specific reporting						
Please select the one below that is the best fit for your research. If you are not sure, read the appropriate sections before making your selection.						
☐ Behavioural & social sciences ☐ Ecological, evolutionary & environmental sciences						
For a reference copy of the document with all sections, see <u>nature.com/documents/nr-reporting-summary-flat.pdf</u>						
Life sciences study design						
All studies must disclose c	on these points even when the disclosure is negative.					
	In oligonucleotide synthesis experiments, 100 to 10,000 microchips per run were employed to synthesize target sequences. These synthesis tests were conducted with a minimum of 10 independent replicates to verify process reproducibility of mMPS system.					
Data exclusions No da	o data was excluded in this study.					
charac	For SEM imaging, 3 replicates were performed for each chip and 3-5 different and random areas in each chip were taken in imaging. Other characterizations on chips and synthesized oligonucleotides were repeated 3-6 times independently. Biological triplicates were performed to study the effect of oligonucleotide mix yielded on gene assembly success rate.					
Randomization All the	the microchips utilized in qualification and quantification analysis were randomly picked up.					
Rlinding Blindin	Blinding is not relevant because no group allocation was involved in this study.					

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		ethods		
n/a Involved in the study	n/a	Involved in the study		
Antibodies	\boxtimes	ChIP-seq		
Eukaryotic cell lines	\boxtimes	Flow cytometry		
Palaeontology and archaeology		MRI-based neuroimaging		
Animals and other o	organisms			
Clinical data				
Dual use research or	Dual use research of concern			
	□ Plants			
Plants				
Seed stocks	N/A			
Novel plant genotypes N/A				
Authentication	N/A			
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