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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 Editorial Policies and the Editorial Policy Checklist.

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	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
x	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
×	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)
x	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>
x	For Bayesian analysis, information on the choice of priors and Markov chain Monte Carlo settings
×	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 availability of computer code

Data collection EMBL Nucleotide Sequence Database (ENA) database under Bioproject PRJEB35712

Data analysis Softwares and version used are: Virsorter version 2.0, CheckV v1.01, PPR-Meta v1.1, geNomad v1.7.1, Vcontact version 2.0, NCBI BLAST ncbi-

blast-2.15.0+, WISH v1.1, Protscale Expasy tool (https://web.expasy.org/protscale/),

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 <u>data availability statement</u>. 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 original sequenced metagenomic and metatranscriptomic raw and assembled data including contigs mined and identified here as viruses along with single-amplified genomes and metagenome-assembled genomes, were available at EMBL Nucleotide Sequence Database (ENA) database under Bioproject PRJEB35712. Data on viral contigs (fasta nucleotide and amino acid sequences) are also available in the Supplementary Dataset that support and complement the manuscript. This comprehensive Supplementary dataset additionally contain data on viral classification, viral network analysis, virus-host assignment, transcriptional activity of

		on. The following public databases were used in this study: geNomad database 7. IMG_VR_2022-09-20_7.1 - IMG/VR v4.1 (viral database) https://genome.jgi.doe.gov/portal/IMG_VR/IMG_VR.home.html,	
Research in	volving hu	ıman participants, their data, or biological material	
Policy information	about studies	with human participants or human data. See also policy information about sex, gender (identity/presentation), ethnicity and racism.	
Reporting on sex a		(n/a	
Reporting on race, ethnicity, or other socially relevant groupings		n/a	
Population characteristics		n/a	
Recruitment		n/a	
Ethics oversight		n/a	
Note that full inform	nation on the app	roval of the study protocol must also be provided in the manuscript.	
Field-spe	ecific re	porting	
lease select the c	one below that	is the best fit for your research. If you are not sure, read the appropriate sections before making your selection.	
X Life sciences		Sehavioural & social sciences	
		all sections, see <u>nature.com/documents/nr-reporting-summary-flat.pdf</u>	
_ife scie	nces st	udy design	
All studies must di	isclose on these	points even when the disclosure is negative.	
Sample size	We do not con	duct sampling in this survey. Sampling and sample size is described previously in Martinez-Perez et al 2022 (Nat Comm)	
Data exclusions	No data were	excluded from the analysis	
Replication	Replication for	sequencing data was successful and described previously in Martinez-Perez et al 2022 (Nat Comm).	
Randomization		e Antarctic continent in the Ross Ice Shelf for performing a drilling campaign was not at random since it was selected based on set by a multidisciplinary and trans-continetal team of researchers. See location of sampling in a previous paper by Martinez-2 (Nat Comm).	
Blinding	sampling and I	ling in the Antarctic continent in the Ross Ice Shelf for performing a drilling campaign does not make any sense because ocation is based on scientific, technical and logistic criteria set by a multidisciplinary and trans-continetal team of researchers. f sampling in a previous paper by Martinez-Perez et al 2022 (Nat Comm).	
Reportir	ng for s	pecific materials, systems and methods	
		about some types of materials, experimental systems and methods used in many studies. Here, indicate whether each material your study. If you are not sure if a list item applies to your research, read the appropriate section before selecting a response.	
Materials & ex	perimental s	systems Methods	
n/a Involved in t	he study	n/a Involved in the study	
Antibodie	ies ChIP-seq		
	Eukaryotic cell lines		
	ology and archaed	———————————————————————————————————————	
Animals a	nd other organisr	ns	

Clinical data
 Dual use research of concern
 Plants