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Reporting Summary

Nature Research 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 Research policies, see our Editorial Policies and the Editorial Policy Checklist.

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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
X	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.
×	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)
	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>
×	For Bayesian analysis, information on the choice of priors and Markov chain Monte Carlo settings
X	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

No software was used for data collection.

Data analysis

Kaiju v1.6.3 for taxonomic analysis, Unicycler v0.4.6 for genome assembly, stringMLST v0.5.1 for MLST classification, Bowtie2-2.3.4.1 and Bowtie v1.1.2 for short reads alignment, Samtools v0.1.19, Samtools-1.8 and Bedtools-2.25.0 to identify SNPs, IQ-TREE 1.6.12 for phylogenetic trees, Prokka v1.13 to annotate open reading frames, Roary v3.11.2 to generate pangenome, igv v2.8.2 for genome coverage visualization.

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 Research guidelines for submitting code & software for further information.

Data

Policy information about <u>availability of data</u>

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 list of figures that have associated raw data
- A description of any restrictions on data availability

All sequencing data that support the findings of this study have been made available through the SRA repository. Accession numbers are detailed in manuscript Text.

Field-spe	ecific re	porting			
Please select the o	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	□В	ehavioural & social sciences			
For a reference copy of t	the document with a	all sections, see <u>nature.com/documents/nr-reporting-summary-flat.pdf</u>			
Life scier	nces stu	ıdy design			
All studies must dis	sclose on these	points even when the disclosure is negative.			
Sample size	The sample size	size of neonates was determined based on inclusion criteria and informed consent.			
Data exclusions	No data was exc	excluded.			
Replication	We could not re	eplicate this observational study.			
Randomization	We included all	consenting eligible individuals.			
Blinding	No blinding too	No blinding took place.			
We require informati system or method list system or method list Materials & ex n/a Involved in the material i	on from authors a ted is relevant to perimental so ne study	n/a Involved in the study ChIP-seq Flow cytometry MRI-based neuroimaging MRI-based neuroimaging			
Clinical dat Dual use re	esearch of concer	n			
Human rese					
Policy Information Population characte		nvolving human research participants Infants hospitalized in the Rambam hospital NICU were included with parental consent if (a) they were born before the thirty-			
seventh week of pregnancy, and (b) their weight at birth was <1,500 grams.					
Recruitment	cruitment Parents of all hospitalized neonates that fit the inclusion criteria were asked for consent.				

Note that full information on the approval of the study protocol must also be provided in the manuscript.

Ethics oversight

Rambam Medical Center, Haifa, Israel issued IRB RMB-13-0508