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Last updated by author(s):	2021.01.07

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 ar	alyses, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.		
n/a	Confirmed			
\boxtimes	The exact	The exact sample size (n) for each experimental group/condition, given as a discrete number and unit of measurement		
\boxtimes	A stateme	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			
	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)			
\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			
	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 <u>statistics for biologists</u> contains articles on many of the points above.				
So	ftware an	d code		
Poli	cy information	about <u>availability of computer code</u>		
Da	ata collection	The databases were queried in pgAdmin 4 v 1.2. Computations were implemented in Matlab R2018b (MathWorks Inc., Natick, MA).		
Da	ata analysis	Matlab R2018b (MathWorks Inc., Natick, MA).		
		g 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 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 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 list of figures that have associated raw data
- A description of any restrictions on data availability

The extraction process has been performed by customized scripts (queries) of Standardized Query Language (SQL) for MIMIC and eICU on the object-relational database system PostgreSQL. The full SQL source code is provided via the dedicated VentAI website: https://ventai.org

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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	Behavioural & so	cial sciences Ecological, evolutionary & environmental sciences	
For a reference copy of t	the document with all sections, see <u>nat</u> u	ure.com/documents/nr-reporting-summary-flat.pdf	
ı :£:		ion	
Life scier	nces study des	<u>ign</u>	
All studies must dis	sclose on these points even wh	en the disclosure is negative.	
Sample size	We included all adult patients from two large intensive care databases, MIMIC-III and eICU-RI. We conducted a secondary analysis of patient data initially collected routinely for patient care.		
Data exclusions	Exclusion criteria were pre-established. We excluded patients younger than 18 years old at the time of ICU admission, patients where mortality was not documented and patients with evidence of withdrawal of treatment. In MIMIC-III, we excluded patients where positive end-expiratory pressure (PEEP), fraction of inspired oxygen (FiO2) and ideal body weight- adjusted tidal volume (Vt) is missing or in case of in sufficient data. In eICU-RI, we excluded ICU readmissions and patient admitted in an ICU with insufficent data collection.		
Replication	The results were reliably reprodu	nced in a large array of sensitivity analyses, as described in Methods and Extended Data.	
Randomization	Not relevant, this was not a randomized controlled study.		
Blinding	Not relevant, this was not a randomized controlled study.		
We require informati system or method list	on from authors about some types ted is relevant to your study. If you	naterials, systems and methods of materials, experimental systems and methods used in many studies. Here, indicate whether each material, are not sure if a list item applies to your research, read the appropriate section before selecting a response.	
Materials & ex	perimental systems	Methods	
n/a Involved in th	,	n/a Involved in the study	
Antibodies		ChIP-seq	
Eukaryotic cell lines		Flow cytometry	
Palaeontology and archaeology MRI-based neuroimaging			
Animals and other organisms Human research participants			
Clinical dat			
Dual use re	esearch of concern		
Human rese	arch participants		
Policy information	about studies involving human	research participants	
Population chara	cteristics The research inv	volved a total of 96,156 unique adult patients admitted to 133 separate intensive care units in the USA. The	

average age was 65 years old and 53% of the subjects were male. All the patients were mechanically ventilated.

Recruitment

There was no recruitment. Only retrsopective data from MIMIC-III database and eICU database

Ethics oversight

Approval of data collection, processing, and release for the MIMIC-III database has been granted by the Institutional Review Boards of Beth Israel Deaconess Medical Center (Boston, MA, USA) and the Massachusetts Institute of Technology (Cambridge, MA, USA). Approval of data collection, processing and release for the eICU database has been granted by the eICU research committee and exempt from Institutional Review Board approval. All data was processed on the computational infrastructure of the Rheinisch Westfälische Technische Hochschule (RWTH) Aachen University and the University Hospital RWTH Aachen in accordance to European Union data protection laws.

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