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Reporting Summary

Data are available under https://osf.io/4gdy2/.

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Statistics			
For	all statistical ar	nalyses, 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 stateme	ent 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 descrip	tion of all covariates tested	
	A descrip	tion 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 Bayes	sian analysis, information on the choice of priors and Markov chain Monte Carlo settings	
	For hiera	rchical and complex designs, identification of the appropriate level for tests and full reporting of outcomes	
\square 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.	
So	ftware an	d code	
Poli	cy information	about availability of computer code	
Da	ata collection	Code provided under https://osf.io/4gdy2/.	
Da	ata analysis	Code provided under https://osf.io/4gdy2/.	
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Da	ta		
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 description of any restrictions on data availability - For clinical datasets or third party data, please ensure that the statement adheres to our policy		

Human research participants

Policy information about studies involving human research participants and Sex and Gender in Research.

Reporting on sex and gender

61 participants of the female sex (16 adult, 45 newborn) and 73 participants of the male sex (14 adult, 59 newborn) were considered in our study. Findings apply to both sexes. The factor of sex was considered in order to obtain a balanced design. Information was self-reported for the adult participants, and reported by the parents for the newborns. Information on gender was not collected. Sex- or gender-specific analyses were not performed as the factor was not implicated in our research questions.

Population characteristics

Healthy young adults with no self-reported indications of psychological and neurological disorders or acute or chronic heavy respiratory diseases that may prevent the participant from sitting still during the EEG recording. We initially measured participants' hearing thresholds between 1 and 12.5 kHz to ensure that they deviated not more than 20 dB from their age mean. Twenty-nine participants fulfilled this requirement and took part in the study (15 female sex: 25.0 ± 2.60 years old; 14 male sex: 25.1 ± 2.77 years old; reported as mean \pm standard deviation). We recruited 104 healthy, full-term newborns (0–4 days of age, 59 male sex, 45 female sex). 46 were firstborns, 34 were second, 14 were third child and 6 had more than 3 siblings. None of them were twins. Mean gestational age was 40.17 ± 1 weeks and mean birth-weight 3787 ± 373 g. All newborns had normal hearing as indicated by successfully completing a Brainstem Evoked Response Audiometry (BERA) test prior to the experiment. No data on race / ethnicity was collected.

Recruitment

Adult participants were recruited through the institute's database of voluntary study participants. Recruitment of newborn participants took place at the maternity ward of the hospital. Pregnant women were contacted by a research nurse employed by RCNS.

Ethics oversight

This study followed Good Clinical Practice (GCP) standards and was conducted in full compliance with the Declaration of Helsinki (2000) and all applicable national and international laws. Adult testing was performed in accordance with the Austrian Universities Act of 2002 and the newborn protocol was also approved by the National Public Health Center, Hungary. Informed consent was obtained from adult participants and from either one or both parents of the newborn participants.

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.			
Life sciences	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>

Behavioural & social sciences study design

All studies must disclose on these points even when the disclosure is negative.

Study description

Quantitative study including two age groups (adults and newborns tested cross-sectionally) with two within subject factors manipulated either in passive or active conditions (only passive presentation was feasible for newborns).

Research sample

For young adult group, twenty-nine participants were included (15 female sex: 25.0 ± 2.60 years old; 14 male sex: 25.1 ± 2.77 years old; denoted as mean \pm standard deviation). We aimed for a sample balanced over sex/gender. The young adults are representative for Austrian university students. For the newborn group, we recruited 104 healthy, full-term neonates (0–4 days of age, 59 male, 45 female). The sample is representative of the full-term population in terms of gestational age and birth weight. The hospital has a large catchment area (ca. 2600 births/year); the patient pool is representative of the population in Budapest. We aimed for a sample balanced over sex and representative of socioeconomic status.

Sampling strategy

To determine the appropriate sample size for our study, an a priori power analysis was conducted for a 2-sided mixed effect design at a 0.05 alpha level and desired power level of 0.80 with medium effect sizes (0.40). This resulted in an estimated sample size of N= 28 for each of the two age groups. To compensate for potential exclusions, 30 adults and 90 newborns were finally recruited.

Data collection

Recruitment of newborns was carried out at the maternity ward of the hospital by a research nurse employed by RCNS.

Timing

Data collection of the adult data was performed continuously in spring 2021. Newborn data was collected continuously over 2021-2022 (occasionally interrupted due to COVID restrictions or due to other studies).

Data exclusions

The exclusion criteria for all studies included a history of psychiatric or neurological symptoms; the usage of recreational drug or medication with any psychoactive effects; and the existence of non-removable metallic parts that would prevent MRI scanning. Adult participants were screened to have pure-tone thresholds within normal age-dependent limits. All newborns had normal hearing as indicated by successfully completing a Brainstem Evoked Response Audiometry (BERA) test. Exclusion of newborn data was based on

		ion of useful trials: after data preprocessing, 33 subjects maintained less than 60 % of the trials and were therefore 1 newborn data sets were finally analyzed for the present study.		
Non-participation Not red		corded		
Randomization F	Participants	ants were allocated into two groups based on their age (newborn, young adult).		
Reporting for	rspe	cific materials, systems and methods		
'		t some types of materials, experimental systems and methods used in many studies. Here, indicate whether each material, study. If you are not sure if a list item applies to your research, read the appropriate section before selecting a response.		
Materials & experimen	ntal syste	ms Methods		
n/a Involved in the study		n/a Involved in the study		
Antibodies Continue		ChIP-seq Flow cytometry		
Palaeontology and arc	chaeology	MRI-based neuroimaging		
Animals and other org	ganisms			
Clinical data				
Dual use research of c	concern			
Magnetic resonand	ce ima	ging		
Experimental design				
Design type		Resting state		
Design specifications		Structural anatomical data was recorded only for EEG source localization purposes.		
Behavioral performance measures		none		
Acquisition				
Imaging type(s)		Structural		
Field strength		3 Tesla		
Sequence & imaging param	neters	not relevant		
Area of acquisition		not relevant		
Diffusion MRI U	Used Not used			
Preprocessing				
Preprocessing software	Preprocessing software MRIcroGL 1.2.20201102, Freesurfer reconstruction			
Normalization	not relevant			
Normalization template	late not relevant			
Noise and artifact removal	and artifact removal not relevant			
Volume censoring	ensoring not relavant			
Statistical modeling & ir	nference			
Model type and settings	del type and settings none			
Effect(s) tested	fect(s) tested none			
Specify type of analysis: Whole brain ROI-based Both				
	Anatomic	al location(s) Desikan-Killiany at las as implemented in Prainctorm		

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Statistic type for inference (See <u>Eklund et al. 2016</u>)	none
Correction	none

Models & analysis

n/a	Involved in the study
X	Functional and/or effective connectivity
X	Graph analysis
\times	Multivariate modeling or predictive analysis