

The amplitude of fNIRS hemodynamic response in the visual cortex unmasks autistic traits in typically developing children

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39 Abstract

40 Autistic traits represent a continuum dimension across the population, with
 41 autism spectrum disorder (ASD) being the extreme end of the distribution.
 42 Accumulating evidence shows that neuroanatomical and neurofunctional
 43 profiles described in relatives of ASD individuals reflect an intermediate
 44 neurobiological pattern between the clinical population and healthy controls.
 45 This suggests that quantitative measures detecting autistic traits in the
 46 general population represent potential candidates for the development of
 47 biomarkers identifying early pathophysiological processes associated with
 48 ASD. Functional near-infrared spectroscopy (fNIRS) has been extensively
 49 employed to investigate neural development and function. In contrast, the
 50 potential of fNIRS to define reliable biomarkers of brain activity has been
 51 barely explored. Features of non-invasiveness, portability, ease of
 52 administration and low-operating costs make fNIRS a suitable instrument to
 53 assess brain function for differential diagnosis, follow-up, analysis of
 54 treatment outcomes and personalized medicine in several neurological
 55 conditions. Here, we introduce a novel standardized procedure with high
 56 entertaining value to measure hemodynamic responses (HDR) in the occipital
 57 cortex of adult subjects and children. We found that the variability of evoked
 58 HDR correlates with the autistic traits of children, assessed by the Autism-
 59 Spectrum Quotient. Interestingly, HDR amplitude was especially linked to
 60 social and communication features, representing the core symptoms of ASD.
 61 These findings establish a quick and easy strategy for measuring visually-
 62 evoked cortical activity with fNIRS that optimize the compliance of young
 63 subjects, setting the background for testing the diagnostic value of fNIRS
 64 visual measurements in the ASD clinical population.

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73 Introduction

74 Autism spectrum disorder (ASD) is a heterogeneous developmental condition that
 75 involves persistent challenges in social interactions, restricted/repetitive behaviors,
 76 and the lack of behavioral and cognitive flexibility¹. Since the pioneering work by
 77 Lorna Wing², increasing epidemiological evidence indicates that autistic traits are
 78 continuously distributed across the general population^{3,4}. This is due to the complex
 79 genetic and epigenetic inheritance pattern of ASD, where multiple candidate loci
 80 contribute to the pathogenesis of the disease^{5,6}. Milder autistic traits have been
 81 termed the extended or broader autism phenotype (BAP), with BAP features being
 82 particularly prevalent in first- and second-degree relatives of individuals with ASD⁶⁻⁹.
 83 Since ASD and broader autistic manifestations share common genetic variants and
 84 neurobiological susceptibility factors¹⁰, the general population emerges as a suitable
 85 testing bed for the development of quantitative measures detecting neurostructural
 86 and neurofunctional hallmarks of autism.

87 Over the last decade, the biological dimension of ASD has been largely
 88 explored, thanks to the growing availability of advanced tools to explore brain
 89 correlates of neurological disorders, including high-density EEG,
 90 magnetoencephalography, positron emission tomography, magnetic resonance
 91 imaging (MRI), and functional near-infrared spectroscopy (fNIRS)^{8,11,12}. A number of
 92 studies reported defective neuroanatomical and neurofunctional features in
 93 individuals with ASD, suggesting that a dysfunction of specific brain areas might
 94 underlie the core symptoms of ASD¹³⁻¹⁷. Interestingly, similar neurological profiles
 95 describe the relatives of autistic probands⁸.

96 fNIRS is an optical imaging technique that allows quantifying oxygen
 97 consumption in different regions of the cerebral cortex, providing an indirect measure
 98 of neuronal activity^{18,19}. This blood-oxygen-level-dependent (BOLD) signal is similar
 99 to that detected with functional MRI (fMRI)²⁰. However, fNIRS is more tolerant to
 100 motion artifacts than fMRI, and the development of robust methods for motion
 101 detection and correction allowed to avoid sedation in children^{21,22}. Furthermore,
 102 fNIRS has the advantage of being totally non-invasive, low-cost, portable, noiseless,
 103 and endowed with high experimental flexibility and no setting constraints. This
 104 methodological strength provides the fNIRS with a high ecological value for
 105 investigating neural circuit maturation either in typically developing children or
 106 clinically relevant populations^{21,23}.

Although the use of fNIRS in autism research is still an emerging area, a number of studies aiming to decipher the neuronal mechanisms and circuits underlying ASD evaluated different aspects of brain function and organization, including resting-state and task-evoked responses^{24,25}. Coherence analyses of resting-state hemodynamic activity showed weaker local and interhemispheric functional connectivity in different cortical regions^{26–31}. Moreover, individuals on the autism spectrum present patterns of atypical activity, including reduced hemodynamic responses within specific brain regions, bilateral differences in neuronal activation and the lack of cortical specialization, in tasks ranging from sensory perception³² to executive functions³³, social perception^{34–37}, joint attention^{38–40}, imitation^{41,42}, facial and emotional processing^{43–46}, speech perception and language^{47–50}. The majority of studies targeting evoked brain activity was focused on the prefrontal and the temporal cortex²⁵, where symptom severity seems to be inversely correlated with the degree of cortical activation^{42,43}.

Growing evidence suggests that fNIRS might be a candidate biomarker for several neuropsychiatric disorders, including ASD^{29,51–55}. In particular, functional network efficiency²⁹, weighted separability of NIRS signals⁵⁶, multi-layer neural networks and sample entropy of spontaneous hemodynamic fluctuations^{54,57} have been proposed as auxiliary diagnosis indexes for ASD. However, all these approaches require complex algorithms to extract high-level features from the fNIRS raw data, while fitness, applicability and translational value of biomarkers greatly depend on their ease of use. In this framework, the analysis of visual phenotype has become an important model to evaluate cortical processing in different neurodevelopmental conditions^{58–63}. Indeed, electrophysiological measurement of visually evoked responses has been introduced as a quantitative method to assess brain function in Rett syndrome^{58,64}, and hemodynamic responses (HDR) emerged as a potential longitudinal biomarker for CDKL5 Deficiency Disorder and Creatine Transporter Deficiency in murine models^{62,63}.

Since clinical studies suggested a dysregulation of sensory processing and functional connectivity in the visual cortex of ASD subjects^{65–67}, we hypothesized that fNIRS visual measures could be related to broad autism dimensions in the general population. To address this issue, we developed a novel standardized procedure to assess HDR changes in the occipital cortex testing 40 randomly selected neuro-

typical adults and 19 children, and measuring inter-individual differences in autistic traits, using the Autism-Spectrum Quotient (AQ^{68,69}).

Results

An animated cartoon-based stimulus is able to evoke visual responses in the adult cortex

We measured the cortical HDR function⁷⁰ elicited by a reversing checkerboard pattern in the adult population. In agreement with the previous literature^{71,72}, we obtained a significant activation of the occipital cortex in response to different conditions of visual stimulation (Fig. 1 and Fig. S1). Grand averages across adult participants (see Table 1 for demographics) of Total Hb (THb), OxyHb (OHb) and DeoxyHb (DHb) concentration changes are plotted in Fig. 2. Using a classic mean-luminance grey screen as baseline, statistical analysis revealed a significant main effect of the checkerboard stimulus (S) with respect to the blank presentation for all HDR metrics (Radial Stimulus condition, RS, Fig. 2A).

To increase the entertaining quality of our experimental paradigm, we devised an innovative visual stimulation protocol blending the checkerboard pattern with an isoluminant commercial cartoon, thus serving as a reference baseline (Fig. 1 and Fig. S1). We found a significant increase of THb and OHb, with a parallel reduction of DHb concentration, in response to S appearance, reflecting the functional activation of visual areas in this condition as well. The cortical response was independent from the cartoon employed as baseline: a comparable HDR, indeed, was clearly elicited both when the baseline movie was fixed a priori by the experimenter (Cartoon Fixed condition, CF; Fig. 2B) and when the cartoon was freely selected by the tested subject (Cartoon Chosen condition, CC; Fig. 2C). Interestingly, a significant pattern of correlations emerged among HDR metrics recorded with different stimulating conditions (Fig. S2), indicating that the quality of visual input does not quantitatively impact HDR. The amplitude of cortical activation was only slightly smaller in response to CF and CC, with the range of OHb and DHb fluctuations being significantly lower with respect to that evoked by RS (Fig. 2D).

Within the CF condition, we also established that the baseline cartoon does not affect the degree of visual activation: indeed, a comparable modification of THb, OHb and DHb concentrations was recorded using “The Lion King”, “The Powerpuff Girls”, “Peppa Pig” or “Kung Fu Panda” (Fig. S3A). Furthermore, no differences of

visually evoked responses were detected modulating the contrast level of the baseline cartoon: THb, OHb and DHb fluctuations, indeed, were comparable when a fixed baseline cartoon was presented at 20%, 40%, or 80% of contrast (Fig. 2E; Fig. S3B-D). Finally, the response latency was homogenous in RS, CF, and CC conditions (Fig. S4A).

Altogether, these results demonstrate the validity of this innovative stimulation procedure to evoke a significant and reliable response in the occipital cortex preserving inter-subject variability.

The cartoon paradigm was reliable in eliciting cortical responses in children

We measured cortical responses in typically developing children (see Table 2 for demographics) viewing the radial checkerboard blended with the animated cartoon. We compared three different conditions: each subject, indeed, was asked to select two cartoons of their preference for the baseline and the first choice was employed for the low-contrast (cartoon 1 low contrast, 20%, L1) and the high-contrast (cartoon 1 high contrast, 80%, H1) stimulation, while the second cartoon was presented only at low-contrast (cartoon 2 low contrast, L2; Fig. S1).

Our data showed a significant activation of the visual cortex, with a prominent change of THb, OHb and DHb concentration in response to the S with respect to the blank for all conditions tested (Fig. 3A-C). The amplitude of elicited cortical responses was comparable following L1, H1 and L2 (Fig. 3D), proving that the HDR is independent from the cartoon narrative selected for the baseline and the contrast level of baseline presentation in children as well. Small differences were observed for response latency among L1, H1 and L2 conditions (Fig. S4B). A highly significant pattern of correlations among different HDR indexes recorded in the diverse conditions was detected (Fig. S5).

In agreement with previous literature⁷², a maturational trend of cortical responsivity was recognized, with children showing significantly higher HDR amplitude with respect to adult subjects (Fig. 3E). On the contrary, age-dependent effects were not identified for response latency (Fig. S4C).

These findings establish a novel method for measuring visually-evoked cortical activity with fNIRS that ensures an elevated compliance of young subjects and high-quality reliability of measurements, suggesting a valuable tool for studying

visual cortical processing in typically developing children, but also in clinically relevant populations.

Negative correlation of HDR amplitude with AQ score

Despite no effects detectable in adults (Fig. 4A-C), the amplitude of visual responses was highly correlated to AQ scores in children (Fig. 4D-E). Consistent with a recent work⁵⁴, the correlation was specific for THb, with higher AQ score being associated with a lower amplitude of THb visually-evoked signals (Fig. 4D-E).

Interestingly, HDR amplitude was especially linked to social and communication autistic traits (Fig. 5, S6): indeed, assessing separately the five AQ subscales⁶⁹ we found a significant correlation of THb and OHb with the Social Skills subscale (AQ_S, Fig. 5A-B), while only THb modulation was related to the Communication subscale (AQ_C, Fig. 5C-D). Given the reliability across different visual tasks (L1 and H1), the strongest interaction was between THb and AQ_S (Fig. 5A-B).

Discussion

We measured hemodynamic responses in the occipital cortex while subjects viewed a reversing checkerboard pattern on a grey isoluminant baseline or the same stimulus blended with a commercial animated cartoon. In all participants, the patterned stimulus elicited a significant change of cortical Hb (upwards for THb and OHb, down for DHb) independently from the reference baseline, while no response was detected following blank presentation. Since HDR consists of an initial increase in oxygen rich blood followed by a smaller depletion of deoxy-haemoglobin, with a tight interrelation among total, oxygenated and reduced Hb levels, reporting all data allows for a more accurate physiological interpretation of the results⁷⁰.

Interestingly, the level of occipital cortex activation did not depend on either the movie selected as reference baseline or the baseline contrast, with only a slight reduction of OHb and DHb change in response to the stimulus blended to animated cartoons with respect to the classic RS condition. These data demonstrate the reliability of this novel procedure with high entertaining and ecological value in eliciting cortical activity. Thus, our approach might be helpful for studying cortical function in children with an atypical trajectory of brain development, commonly showing a reduced compliance in experimental environments.

The magnitude of HDR modulation, and in particular of THb, was inversely correlated with AQ scores in children. Indeed, we found that the higher were the AQ scores of subjects the lower was the amplitude of THb response to visual stimulation, suggesting that visually-evoked fNIRS responses are able to capture the dimension of autistic traits in the general young population. Our findings are consistent with previous studies showing that cortical activation measured with fNIRS and the performance in visual psychophysics negatively reflected ASD symptom severity^{42,43,73–76}. Moreover, these data reinforce the concept that THb changes could provide richer discriminative information for classifying between typically developing children and ASD subjects⁵⁴. A tentative explanation of reduced HDR in children with stronger autistic traits might be found in the difference of perceptual styles in the general population^{77,78}: the preference for focusing on local details vs. the global stimulus configuration, indeed, is a defining feature of ASD⁷⁹ and locally centered perception could be less effective in activating neural circuits of visual cortex. Interestingly, a recent study established a vascular link to ASD, showing early dysfunction of endothelial cells and impaired endothelium-dependent vasodilation in a mouse model of 16p11.2 deletion⁸⁰. Since the HDR measured with fNIRS strongly relies on neurovascular coupling, this suggests that lower neurophysiological activity may stem in part from endothelial-dependent vascular factors. In contrast, we failed to detect any correlation between HDR and AQ scores in adult subjects. This is likely due to the different dynamic range and amplitude of HDR responses measured in children and adult participants. Accordingly, the maturation of neural and vascular networks over the first years of life affects neurovascular coupling with a pattern of hemodynamic responses significantly different in developing and adult brain⁸¹.

In our experiment, the THb index describes about 45% of the variance in AQ scores. This correlation is remarkably high, considering that it is detected across two separate visual stimulating procedures, i.e., the vision of low- and the high-contrast blended RS-animated cartoons. Although the autistic questionnaire for children is well-validated, showing good test-retest reliability and high internal consistency, not all items have the same validity and factor analysis identified five subscales, named ‘Social Skills’, ‘Communication’, ‘Attention to Detail’, ‘Imagination’ and ‘Attention Switching’⁶⁹. We observed variability in the strength of correlation between HDR and AQ subscales, with the highest correlation for the ‘Social Skills’ and ‘Communication’

subscales. Interestingly, 'Social Skills' and 'Communication' are the subscales with higher construct validity performance in differentiating individuals with or without ASD^{82,83}, while 'Attention to Detail' was the poorest classifying domain⁸⁴. It is also worth stressing that among the items composing the 'Attention to Detail' subscale, only four actually relate to sensory perception (e.g. 'My child usually notices details that others do not'), the others being more focused on cognitive functions (e.g. 'My child is fascinated by numbers'). Accordingly, the first pilot report of the AQ questionnaire showed lower internal consistency (measured by Cronbach's alpha coefficient) of 'Attention to Detail' items compared to other subscales⁶⁸.

Although primarily affecting social functioning, there is a growing body of evidence showing that ASD is also associated with abnormalities in multiple sensory domains, fluctuating between hyper- and hypo-sensitivity to sensory stimuli^{73,85}. In addition to a higher incidence of refractive errors and strabismus⁸⁶, anomalies in visual processing, visual attention and visual-motor integration have been described in ASD population^{66,73,87}. Interestingly, sensory symptoms are correlated with the severity of the disorder, at least in children⁸⁸. Moreover, commonly observed alterations in social skills might have a visual component⁷³ and perception deficits could impact with cascading effects on the maturation of cognitive and social domains⁸⁷.

It has been recently suggested that an early assessment of pupil size modulation and visual behavior might improve the diagnostic process of ASD^{66,77,87,89}. Currently, ASD diagnosis and follow-up almost entirely rely on phenotypic information collected via clinical measures and parental input that are highly prone to subjective bias⁹⁰. Moreover, the late appearance of some behavioral autistic traits often delays the diagnosis until mid-childhood^{91,92}. Thus, the identification of solid brain biomarkers early predicting ASD pathophysiology is a critical step to anticipate tailored interventions, leading to better outcomes for patients and possibly even the prevention of certain behaviors typically associated with ASD. Objective biomarkers have also the potential to be helpful in the management of patients, allowing the classification of disease severity and monitoring response to treatments¹². A recent systematic review highlighted that both functional and structural neuroimaging features might predict ASD diagnosis in the early pre-symptomatic period^{93,94}, but further studies are needed to validate the promising performance of such biomarkers¹². Lately, resting-state fNIRS

measurements have been suggested as candidate biomarkers for ASD^{29,54,56,57}. As stressed above, fNIRS offers significant advantages with respect to other neuroimaging tools, including non-invasiveness, ease of use, no need of sedation, tolerance to movements and portability, making it a child-friendly approach. However, the extraction of metrics with diagnostic value from resting-state recordings involves complex algorithms.

In contrast, our analysis of visually evoked responses is quick, easy and requires only that children pay attention to a short movie of their choice. Since our stimulating strategy has been studied to optimize the compliance of young subjects, we believe that our results might set the background for testing the predictive value of fNIRS visual measurements to empower early detection of autistic traits. Moreover, screening for autistic traits in the general population may be helpful in epidemiological research because it may provide a large sample size to investigate the correlation between autism phenotype severity and other pathophysiological processes⁸³.

Materials and methods

Subjects

We recruited a total of 40 adult subjects (20 women, age: 31.05 ± 3.94 (SD) years) and 19 children (5 girls, age: 7.20 ± 3.01 (SD) years). All participants reported normal or corrected-to-normal vision and had no diagnosed neuropsychiatric condition. Experimental procedures on children were authorized by the Regional Pediatrics Ethics Board (Comitato Etico Pediatrico Regionale-Azienda Ospedaliero-Universitaria Meyer-Firenze, Italy; authorization number 201/2019) and were performed according to the declaration of Helsinki. Written informed consent was obtained from all adult participants and from the parents of each child, authorizing the use of anonymized data for research purposes. Assent was also obtained from the children involved in the study before participation.

AQ score

Adult participants filled in the Autistic-traits Quotient (AQ) questionnaire, a 50-items self-administered report validated for the Italian version^{68,95}. The items consist of descriptive statements assessing personal preferences and typical behavior. For each item, participants respond on a 4-point Likert scale: “strongly agree”, “slightly

agree", "slightly disagree", and "strongly disagree". The items are grouped in five subscales: Social Skills, Communication, Attention to Details, Imagination and Attention Switching. All the questionnaires were scored by a neuropsychiatrist blinded to subject data: 1 point was assigned when the participant's response was characteristic of ASD (slightly or strongly), 0 points were attributed otherwise. Total scores range between 0 and 50, with 32 being the clinical threshold for autism risk⁶⁸. No subjects scoring above 32 points were recorded. The mean (min-max) of the scores was 15.0 (3-32) with SD of 6.5. Since child self-report might be affected by reading and comprehension difficulties, the children's version of Autism Spectrum Quotient (Italian version of AQ-child) was completed by parents⁶⁹. This version of the AQ questionnaire includes 50 items as well, grouped in the same subscales described above, and parents were required to report for each statement the degree of consistency with their child's behavior. Scores range from 0 to 150, since the response scale is treated as a 4-point Likert scale with 0 representing definitely agree; 1 slightly agree; 2 slightly disagree; and 3 definitely disagree. Items were reverse scored as needed. The threshold score is 76⁶⁹. All subjects scored below 76 points. The mean (min-max) of the scores was 32.1 (17-49) with SD of 10.7.

Apparatus and montages

To measure changes in total Hb (THb) concentration and relative oxygenation levels (OHb and DHb) in the occipital cortex during the task, we used a continuous-wave NIRS system (NIRSport 8x8, NIRx Medical Technologies LLC, Berlin, Germany). Our NIRSport system consists of 8 red light-sources operating at 760 nm and 850 nm, and 7 detectors which can be placed into a textile EEG cap (EASYCAP, Herrsching, Germany), forming an array of 22 multi-distant channels⁹⁶. Textile EEG caps of different sizes were used. The probe arrangement was fixed in each of the caps using grommets, optode stabilizers, colored labels and holders in order to assure comparable probe mapping over all subjects. For data recording, the Aurora Software 1.4.1.1 (NIRx Medical Technologies LLC) was employed. The sampling rate was 10.2 Hz. Visual areas were identified according to the craniocerebral topography within the international 10-20 system and the placement of the optodes was done using fOLD v2.2⁹⁷ and NIRSite 2.0 (NIRx Medical Technologies LLC) softwares. Sources and detectors were symmetrically distributed to define 22 channels around the region of interest, each adjacent pair of sources and detectors

defining one channel (min-max source-detector separation: 20-44 mm for adults, 22-30 mm for children; Fig. S2).

Experimental Design and Visual Stimulation

Prior to the experiment, adult participants (or parents for children) filled in the AQ questionnaire. Then, subjects were asked to sit on a comfortable chair and the fNIRS cap was positioned. Optodes were placed into the cap and the calibration of light coupling between sensors and detectors was performed. All experimental sessions lasted 30 minutes. Visual stimuli were generated using Python 3 and Psychopy3⁹⁸ and displayed with gamma correction on a monitor (Sharp LC-32LE352EWH, 60Hz refresh rate, 45 cd/m² mean luminance, resolution of 800×600 pixels) placed 70 cm from the subject. Cortical hemodynamics in response to full-field, reversing, square wave, radial checkerboard, with abrupt phase inversion (spatial frequency: 0.33 cycles per degree, temporal frequency: 4 Hz; Fig. 1A) was evaluated in the time domain by measuring the peak-to-baseline amplitude and latency. To have an internal control with blank stimulation, we used an event-related design consisting of: i) 20 cycles of 5 seconds stimulus 'on' (reversing checkerboard, 90% of contrast) followed by 10 seconds stimulus 'off' and ii) 20 cycles of 5 seconds mock stimulus 'on' (reversing checkerboard, 0% of contrast) followed by 10 seconds stimulus 'off'. The two stimulating conditions were pseudo randomly interleaved for each subject during the recording. Blocks lasted 10 minutes and participants were permitted to take rest between recordings. Fig. 1D shows a schematic representation of the experimental procedure. Visual events were synchronized with NIRSport over wireless LAN communication through the Python version of LabStreamingLayer (<https://github.com/sccn/labstreaminglayer>).

Recordings in adult participants- Experiment 1 for adults (exp1) aimed to understand whether a reliable hemodynamic signal could be recorded in response to the radial checkerboard merged with an animated cartoon. Thus, exp1 started with a 10-minutes recording using the reversing checkerboard as stimulus 'on' and the grey screen as stimulus 'off' (RS condition), and continued with the vision of two different blended animated cartoons, where the stimulus 'on' was a merge between the reversing checkerboard and the movie, whereas the stimulus 'off' was the grey-scale isoluminant cartoon (CF and CC conditions; Fig. S1). The merging procedure was achieved using Python3 OpenCV⁹⁹. During the appearance of the stimulus each

frame was filtered using an automatic Canny edge detection algorithm (<https://www.pyimagesearch.com/2015/04/06/zero-parameter-automatic-canny-edge-detection-with-python-and-opencv/>), then the filtered cartoon was blended with the radial checkerboard. Each pixel of the animated cartoon with the same color of the corresponding pixel of the radial checkerboard was inverted, to obtain a fully visible image. The result was a RS with an overlaid cartoon frame (Fig 1B). The first cartoon was randomly selected by the operator within a group of 4 (“The Lion King”, “The Powerpuff Girls”, “Peppa Pig” or “Kung Fu Panda”; CF), whereas the latter was a free choice of the subject (CC). Exp2, aiming to dissect the contribution of baseline contrast to visual responses, was performed in a subset of adult participants (n = 15). Exp2 consisted of 3 consecutive recordings of a CF (“Peppa Pig”; “Hide-and-seek”, “Fly the kite”, “Polly parrot” episodes) with the modulation of the baseline contrast (20%, 40%, 80%; Fig. S1). The presentation order of different contrast levels was randomly shuffled.

Recordings in children- To confirm that the baseline movie and its contrast do not affect the emergence of visual responses to the radial checkerboard in children, we measured hemodynamic signals in response to 2 different blended RS-animated cartoons freely decided by the subject: cartoon 1 was presented at both low (20%, L1) and high (80%, H1) contrast, while only low contrast was recorded for cartoon 2 (L2; Fig. S1). In this case, the presentation order was decided by the child, in order to maximize subject compliance.

During the experimental sessions, data were quickly analyzed and visualized using nirsLAB software (NIRx Medical Technologies LLC, v2019.4).

Signal Processing and Statistical analysis

Data preprocessing was completed using the Homer3 package (v1.29.8) in MATLAB (R2020a). We created a processing stream tailored on recent guidelines for analysis of fNIRS data²². First, the raw intensity data were converted to optical density (OD) changes (*hmR_Intensity2OD*). Then, channels showing very high or low optical intensity were excluded from further analyses using the function *hmR_PruneChannels* (dRange: 5e-04-1e+00, SNRthresh: 2; SDrange: 0.0-45.0). Motion artifacts were then removed by a multistep rejection protocol. After a step of motion artifact detection using the *hmR_MotionArtifactByChannel* function (tMotion: 1.0, tMask: 1.0; STDEVthresh 13.0; AMPthresh: 0.40), motion correction was

performed with a combination of Spline interpolation (*hmR_MotionCorrectSpline*, p: 0.99) and Wavelet filtering (*hmR_MotionCorrectWavelet*, iqr: 0.80) functions²². The remaining uncorrected motion artifacts were identified using the *hmR_MotionArtifactByChannel*. A band-pass filter (*hmR_BandpassFilt: Bandpass_Filter_OpticalDensity*, hpf: 0.01, lpf: 0.50) was applied to decrease slow drifts and high-frequency noise, and the OD data were converted to Hb concentration changes using the modified Beer–Lambert law (*hmR_OD2Conc*, ppf: 1.0 1.0 1.0). Finally, trials of each subject were block-averaged for every stimulating condition and channel (*hmR_BlockAvg: Block_Average_on_Concentration_Data*, trange: -2.0 20.0)²². The resulting txt file was imported in Python as a Pandas DataFrame. For each subject, only the channel with the highest response amplitude was analyzed. The peak response was identified as the maximal value for THb and OHb and the minimum value for DHb. A grand average was taken of the 20 trials of data per stimulating condition and differences between visual stimulation ‘on’ (reversing checkerboard) and ‘off’ (blank) were compared. All data were normalised with respect to the blank-evoked response using a subtraction method. Statistical analysis was carried out using *pingouin* Python library¹⁰⁰ and the following functions: *pingouin.ttest* (paired and two-sided t-test), *pingouin.rm_anova* (one-way repeated measures ANOVA), *pingouin.pairwise_ttests* (post-hoc analysis), *pingouin.pairwise_corr* (Spearman correlation), *pingouin.regplot* (Linear regression). T-test, ANOVA and post-hoc analysis were used to assess differences in fNIRS peak responses following different stimulating conditions, whereas we tested the interaction between the amplitude of fNIRS measures and AQ scores with Spearman correlation and we employed the Linear regression to plot such correlation. Adjustments for multiple comparisons were performed using the Benjamini/Hochberg false discovery rate (BH-FDR) correction. The effect size calculated for the ANOVA was the generalized eta-squared. All the plots have been generated using *Matplotlib* Python library¹⁰¹. All statistical metrics and details are reported in Table S1.

Data availability

The datasets generated during the current study and scripts used for visual stimulation are available, respectively on Zenodo (<http://doi.org/10.5281/zenodo.5101912>) and GitHub website (https://github.com/raffaelemazziotti/FNIRS_code).

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480 **References**

- 481 1. Maenner, M. J. *et al.* Potential impact of DSM-5 criteria on autism spectrum
482 disorder prevalence estimates. *JAMA Psychiatry* **71**, 292–300 (2014).
- 483 2. Wing, L. The Continuum of Autistic Characteristics. *Diagnosis and Assessment*
484 *in Autism* 91–110 (1988) doi:10.1007/978-1-4899-0792-9_7.
- 485 3. Posserud, M.-B., Lundervold, A. J. & Gillberg, C. Autistic features in a total
486 population of 7-9-year-old children assessed by the ASSQ (Autism Spectrum
487 Screening Questionnaire). *J. Child Psychol. Psychiatry* **47**, 167–175 (2006).
- 488 4. Ruzich, E. *et al.* Measuring autistic traits in the general population: a systematic
489 review of the Autism-Spectrum Quotient (AQ) in a nonclinical population sample
490 of 6,900 typical adult males and females. *Mol. Autism* **6**, 2 (2015).
- 491 5. Persico, A. M. & Bourgeron, T. Searching for ways out of the autism maze:
492 genetic, epigenetic and environmental clues. *Trends Neurosci.* **29**, 349–358
493 (2006).
- 494 6. Ruzich, E. *et al.* Subgrouping siblings of people with autism: Identifying the
495 broader autism phenotype. *Autism Res.* **9**, 658–665 (2016).
- 496 7. Piven, J., Palmer, P., Jacobi, D., Childress, D. & Arndt, S. Broader autism
497 phenotype: evidence from a family history study of multiple-incidence autism
498 families. *Am. J. Psychiatry* **154**, 185–190 (1997).
- 499 8. Billeci, L. *et al.* The Broad Autism (Endo)Phenotype: Neurostructural and
500 Neurofunctional Correlates in Parents of Individuals with Autism Spectrum
501 Disorders. *Front. Neurosci.* **10**, 346 (2016).
- 502 9. Carpita, B. *et al.* The broad autism phenotype in real-life: clinical and functional
503 correlates of autism spectrum symptoms and rumination among parents of
504 patients with autism spectrum disorder. *CNS Spectr.* **25**, 765–773 (2020).
- 505 10. Bralten, J. *et al.* Autism spectrum disorders and autistic traits share genetics and
506 biology. *Mol. Psychiatry* **23**, 1205–1212 (2018).
- 507 11. Pagnozzi, A. M., Conti, E., Calderoni, S., Fripp, J. & Rose, S. E. A systematic
508 review of structural MRI biomarkers in autism spectrum disorder: A machine
509 learning perspective. *Int. J. Dev. Neurosci.* **71**, 68–82 (2018).
- 510 12. Frye, R. E. *et al.* Emerging biomarkers in autism spectrum disorder: a systematic
511 review. *Ann Transl Med* **7**, 792 (2019).

- 512 13. Amaral, D. G., Schumann, C. M. & Nordahl, C. W. Neuroanatomy of autism.
513 *Trends Neurosci.* **31**, 137–145 (2008).
- 514 14. Bellani, M., Calderoni, S., Muratori, F. & Brambilla, P. Brain anatomy of autism
515 spectrum disorders I. Focus on corpus callosum. *Epidemiol. Psychiatr. Sci.* **22**,
516 217–221 (2013).
- 517 15. Bellani, M., Calderoni, S., Muratori, F. & Brambilla, P. Brain anatomy of autism
518 spectrum disorders II. Focus on amygdala. *Epidemiol. Psychiatr. Sci.* **22**, 309–
519 312 (2013).
- 520 16. Billeci, L. *et al.* On the application of quantitative EEG for characterizing autistic
521 brain: a systematic review. *Front. Hum. Neurosci.* **7**, 442 (2013).
- 522 17. Calderoni, S., Bellani, M., Hardan, A. Y., Muratori, F. & Brambilla, P. Basal
523 ganglia and restricted and repetitive behaviours in Autism Spectrum Disorders:
524 current status and future perspectives. *Epidemiol. Psychiatr. Sci.* **23**, 235–238
525 (2014).
- 526 18. Lloyd-Fox, S., Blasi, A. & Elwell, C. E. Illuminating the developing brain: the past,
527 present and future of functional near infrared spectroscopy. *Neurosci. Biobehav.*
528 *Rev.* **34**, 269–284 (2010).
- 529 19. Gervain, J. *et al.* Near-infrared spectroscopy: a report from the McDonnell infant
530 methodology consortium. *Dev. Cogn. Neurosci.* **1**, 22–46 (2011).
- 531 20. Raichle, M. E. & Mintun, M. A. Brain work and brain imaging. *Annu. Rev.*
532 *Neurosci.* **29**, 449–476 (2006).
- 533 21. Vanderwert, R. E. & Nelson, C. A. The use of near-infrared spectroscopy in the
534 study of typical and atypical development. *Neuroimage* **85 Pt 1**, 264–271 (2014).
- 535 22. Di Lorenzo, R. *et al.* Recommendations for motion correction of infant fNIRS
536 data applicable to multiple data sets and acquisition systems. *Neuroimage* **200**,
537 511–527 (2019).
- 538 23. Yamasaki, T. *et al.* Rapid maturation of voice and linguistic processing systems
539 in preschool children: a near-infrared spectroscopic study. *Exp. Neurol.* **250**,
540 313–320 (2013).
- 541 24. Mazzoni, A., Grove, R., Eapen, V., Lenroot, R. K. & Bruggemann, J. The
542 promise of functional near-infrared spectroscopy in autism research: What do we
543 know and where do we go? *Soc. Neurosci.* **14**, 505–518 (2019).
- 544 25. Zhang, F. & Roeyers, H. Exploring brain functions in autism spectrum disorder:
545 A systematic review on functional near-infrared spectroscopy (fNIRS) studies.

- 546 *Int. J. Psychophysiol.* **137**, 41–53 (2019).
- 547 26. Keehn, B., Wagner, J. B., Tager-Flusberg, H. & Nelson, C. A. Functional
548 connectivity in the first year of life in infants at-risk for autism: a preliminary near-
549 infrared spectroscopy study. *Front. Hum. Neurosci.* **7**, 444 (2013).
- 550 27. Zhu, H., Fan, Y., Guo, H., Huang, D. & He, S. Reduced interhemispheric
551 functional connectivity of children with autism spectrum disorder: evidence from
552 functional near infrared spectroscopy studies. *Biomed. Opt. Express* **5**, 1262–
553 1274 (2014).
- 554 28. Li, J. *et al.* Characterization of autism spectrum disorder with spontaneous
555 hemodynamic activity. *Biomed. Opt. Express* **7**, 3871–3881 (2016).
- 556 29. Li, Y. & Yu, D. Weak network efficiency in young children with Autism Spectrum
557 Disorder: Evidence from a functional near-infrared spectroscopy study. *Brain*
558 *and Cognition* vol. 108 47–55 (2016).
- 559 30. Jia, H., Li, Y. & Yu, D. Attenuation of long-range temporal correlations of
560 neuronal oscillations in young children with autism spectrum disorder.
561 *Neuroimage Clin* **20**, 424–432 (2018).
- 562 31. Cao, W. *et al.* The Development of Brain Network in Males with Autism
563 Spectrum Disorders from Childhood to Adolescence: Evidence from fNIRS
564 Study. *Brain Sci* **11**, (2021).
- 565 32. Xu, M., Minagawa, Y., Kumazaki, H., Okada, K.-I. & Naoi, N. Prefrontal
566 Responses to Odors in Individuals With Autism Spectrum Disorders: Functional
567 NIRS Measurement Combined With a Fragrance Pulse Ejection System. *Front.*
568 *Hum. Neurosci.* **14**, 523456 (2020).
- 569 33. Xiao, T. *et al.* Response inhibition impairment in high functioning autism and
570 attention deficit hyperactivity disorder: evidence from near-infrared spectroscopy
571 data. *PLoS One* **7**, e46569 (2012).
- 572 34. Lloyd-Fox, S. *et al.* Reduced neural sensitivity to social stimuli in infants at risk
573 for autism. *Proc. Biol. Sci.* **280**, 20123026 (2013).
- 574 35. Braukmann, R. *et al.* Diminished socially selective neural processing in 5-month-
575 old infants at high familial risk of autism. *Eur. J. Neurosci.* **47**, 720–728 (2018).
- 576 36. Lloyd-Fox, S. *et al.* Cortical responses before 6 months of life associate with
577 later autism. *Eur. J. Neurosci.* **47**, 736–749 (2018).
- 578 37. Bhat, A. N., McDonald, N. M., Eilbott, J. E. & Pelphrey, K. A. Exploring cortical
579 activation and connectivity in infants with and without familial risk for autism

- during naturalistic social interactions: A preliminary study. *Infant Behav. Dev.* **57**, 101337 (2019).
38. Iwanaga, R. *et al.* Usefulness of near-infrared spectroscopy to detect brain dysfunction in children with autism spectrum disorder when inferring the mental state of others. *Psychiatry Clin. Neurosci.* **67**, 203–209 (2013).
39. Zhu, B. & Godavarty, A. Functional connectivity in the brain in joint attention skills using near infrared spectroscopy and imaging. *Behav. Brain Res.* **250**, 28–31 (2013).
40. Zhu, H. *et al.* Atypical prefrontal cortical responses to joint/non-joint attention in children with autism spectrum disorder (ASD): A functional near-infrared spectroscopy study. *Biomed. Opt. Express* **6**, 690–701 (2015).
41. Tamura, R., Kitamura, H., Endo, T., Abe, R. & Someya, T. Decreased leftward bias of prefrontal activity in autism spectrum disorder revealed by functional near-infrared spectroscopy. *Psychiatry Res.* **203**, 237–240 (2012).
42. Su, W.-C. *et al.* Differences in cortical activation patterns during action observation, action execution, and interpersonal synchrony between children with or without autism spectrum disorder (ASD): An fNIRS pilot study. *PLoS One* **15**, e0240301 (2020).
43. Kita, Y. *et al.* Self-face recognition in children with autism spectrum disorders: a near-infrared spectroscopy study. *Brain Dev.* **33**, 494–503 (2011).
44. Nakadoi, Y. *et al.* Multi-channel near-infrared spectroscopy shows reduced activation in the prefrontal cortex during facial expression processing in pervasive developmental disorder. *Psychiatry Clin. Neurosci.* **66**, 26–33 (2012).
45. Fox, S. E., Wagner, J. B., Shrock, C. L., Tager-Flusberg, H. & Nelson, C. A. Neural processing of facial identity and emotion in infants at high-risk for autism spectrum disorders. *Front. Hum. Neurosci.* **7**, 89 (2013).
46. Mori, K. *et al.* Neuroimaging in autism spectrum disorders: 1H-MRS and NIRS study. *J. Med. Invest.* **62**, 29–36 (2015).
47. Minagawa-Kawai, Y. *et al.* Cerebral laterality for phonemic and prosodic cue decoding in children with autism. *Neuroreport* **20**, 1219–1224 (2009).
48. Funabiki, Y., Murai, T. & Toichi, M. Cortical activation during attention to sound in autism spectrum disorders. *Res. Dev. Disabil.* **33**, 518–524 (2012).
49. Edwards, L. A., Wagner, J. B., Tager-Flusberg, H. & Nelson, C. A. Differences in Neural Correlates of Speech Perception in 3 Month Olds at High and Low Risk

- for Autism Spectrum Disorder. *J. Autism Dev. Disord.* **47**, 3125–3138 (2017).
50. Pecukonis, M., Perdue, K. L., Wong, J., Tager-Flusberg, H. & Nelson, C. A. Exploring the relation between brain response to speech at 6-months and language outcomes at 24-months in infants at high and low risk for autism spectrum disorder: A preliminary functional near-infrared spectroscopy study. *Dev. Cogn. Neurosci.* **47**, 100897 (2021).
51. Yasumura, A. *et al.* Age-related differences in frontal lobe function in children with ADHD. *Brain Dev.* **41**, 577–586 (2019).
52. Chou, P.-H., Huang, C.-J. & Sun, C.-W. The Potential Role of Functional Near-Infrared Spectroscopy as Clinical Biomarkers in Schizophrenia. *Curr. Pharm. Des.* **26**, 201–217 (2020).
53. Husain, S. F. *et al.* Validating a functional near-infrared spectroscopy diagnostic paradigm for Major Depressive Disorder. *Sci. Rep.* **10**, 9740 (2020).
54. Xu, L. *et al.* Characterizing autism spectrum disorder by deep learning spontaneous brain activity from functional near-infrared spectroscopy. *J. Neurosci. Methods* **331**, 108538 (2020).
55. Yang, D., Hong, K.-S., Yoo, S.-H. & Kim, C.-S. Evaluation of Neural Degeneration Biomarkers in the Prefrontal Cortex for Early Identification of Patients With Mild Cognitive Impairment: An fNIRS Study. *Front. Hum. Neurosci.* **13**, 317 (2019).
56. Yanagisawa, K., Nakamura, N., Tsunashima, H. & Narita, N. Proposal of auxiliary diagnosis index for autism spectrum disorder using near-infrared spectroscopy. *Neurophotronics* **3**, 031413 (2016).
57. Xu, L. *et al.* Identification of autism spectrum disorder based on short-term spontaneous hemodynamic fluctuations using deep learning in a multi-layer neural network. *Clin. Neurophysiol.* **132**, 457–468 (2021).
58. Durand, S. *et al.* NMDA receptor regulation prevents regression of visual cortical function in the absence of Mecp2. *Neuron* **76**, 1078–1090 (2012).
59. de Freitas Dotto, P. *et al.* Sweep visually evoked potentials and visual findings in children with West syndrome. *Eur. J. Paediatr. Neurol.* **18**, 201–210 (2014).
60. Begenisic, T., Sansevero, G., Baroncelli, L., Cioni, G. & Sale, A. Early environmental therapy rescues brain development in a mouse model of Down syndrome. *Neurobiol. Dis.* **82**, 409–419 (2015).
61. Boggio, E. M. *et al.* Visual impairment in FOXP1-mutated individuals and mice.

- 648 *Neuroscience* **324**, 496–508 (2016).
- 649 62. Mazziotti, R. *et al.* Searching for biomarkers of CDKL5 disorder: early-onset
650 visual impairment in CDKL5 mutant mice. *Hum. Mol. Genet.* **26**, 2290–2298
651 (2017).
- 652 63. Mazziotti, R. *et al.* Novel translational phenotypes and biomarkers for creatine
653 transporter deficiency. *Brain Commun* **2**, fcaa089 (2020).
- 654 64. LeBlanc, J. J. *et al.* Visual evoked potentials detect cortical processing deficits in
655 Rett syndrome. *Ann. Neurol.* **78**, 775–786 (2015).
- 656 65. Keehn, B., Westerfield, M. & Townsend, J. Brief Report: Cross-Modal Capture:
657 Preliminary Evidence of Inefficient Filtering in Children with Autism Spectrum
658 Disorder. *J. Autism Dev. Disord.* **49**, 385–390 (2019).
- 659 66. Little, J.-A. Vision in children with autism spectrum disorder: a critical review.
660 *Clin. Exp. Optom.* **101**, 504–513 (2018).
- 661 67. Seymour, R. A., Rippon, G., Gooding-Williams, G., Schoffelen, J. M. & Kessler,
662 K. Dysregulated oscillatory connectivity in the visual system in autism spectrum
663 disorder. *Brain* **142**, 3294–3305 (2019).
- 664 68. Baron-Cohen, S., Wheelwright, S., Skinner, R., Martin, J. & Clubley, E. The
665 autism-spectrum quotient (AQ): evidence from Asperger syndrome/high-
666 functioning autism, males and females, scientists and mathematicians. *J. Autism*
667 *Dev. Disord.* **31**, 5–17 (2001).
- 668 69. Auyeung, B., Baron-Cohen, S., Wheelwright, S. & Allison, C. The Autism
669 Spectrum Quotient: Children's Version (AQ-Child). *J. Autism Dev. Disord.* **38**,
670 1230–1240 (2008).
- 671 70. Tachtsidis, I. & Scholkmann, F. False positives and false negatives in functional
672 near-infrared spectroscopy: issues, challenges, and the way forward.
673 *Neurophotonics* **3**, 031405 (2016).
- 674 71. Chen, L.-C., Sandmann, P., Thorne, J. D., Herrmann, C. S. & Debener, S.
675 Association of Concurrent fNIRS and EEG Signatures in Response to Auditory
676 and Visual Stimuli. *Brain Topogr.* **28**, 710–725 (2015).
- 677 72. Ward, L. M., Aitchison, R. T., Tawse, M., Simmers, A. J. & Shahani, U. Reduced
678 Haemodynamic Response in the Ageing Visual Cortex Measured by Absolute
679 fNIRS. *PLoS One* **10**, e0125012 (2015).
- 680 73. Simmons, D. R. *et al.* Vision in autism spectrum disorders. *Vision Res.* **49**,
681 2705–2739 (2009).

74. Park, W. J., Schauder, K. B., Zhang, R., Bennetto, L. & Tadin, D. High internal noise and poor external noise filtering characterize perception in autism spectrum disorder. *Sci. Rep.* **7**, 17584 (2017).
75. Noel, J.-P., Lakshminarasimhan, K. J., Park, H. & Angelaki, D. E. Increased variability but intact integration during visual navigation in Autism Spectrum Disorder. *Proc. Natl. Acad. Sci. U. S. A.* **117**, 11158–11166 (2020).
76. Noel, J.-P., Zhang, L.-Q., Stocker, A. A. & Angelaki, D. E. Individuals with autism spectrum disorder have altered visual encoding capacity. *PLoS Biol.* **19**, e3001215 (2021).
77. Turi, M., Burr, D. C. & Binda, P. Pupillometry reveals perceptual differences that are tightly linked to autistic traits in typical adults. *Elife* **7**, (2018).
78. Tortelli, C., Turi, M., Burr, D. C. & Binda, P. Objective pupillometry shows that perceptual styles covary with autistic-like personality traits. *Elife* **10**, (2021).
79. Hallen, R. V. der *et al.* Global processing takes time: A meta-analysis on local–global visual processing in ASD. *Psychological Bulletin* vol. 141 549–573 (2015).
80. Ouellette, J. *et al.* Vascular contributions to 16p11.2 deletion autism syndrome modeled in mice. *Nat. Neurosci.* **23**, 1090–1101 (2020).
81. Kozberg, M. & Hillman, E. Neurovascular coupling and energy metabolism in the developing brain. *Prog. Brain Res.* **225**, 213–242 (2016).
82. Broadbent, J., Galic, I. & Stokes, M. A. Validation of Autism Spectrum Quotient Adult Version in an Australian Sample. *Autism Research and Treatment* vol. 2013 1–7 (2013).
83. Lundqvist, L.-O. & Lindner, H. Is the Autism-Spectrum Quotient a Valid Measure of Traits Associated with the Autism Spectrum? A Rasch Validation in Adults with and Without Autism Spectrum Disorders. *J. Autism Dev. Disord.* **47**, 2080–2091 (2017).
84. Wouters, S. G. M. & Spek, A. A. The use of the Autism-spectrum Quotient in differentiating high-functioning adults with autism, adults with schizophrenia and a neurotypical adult control group. *Research in Autism Spectrum Disorders* vol. 5 1169–1175 (2011).
85. Leekam, S. R., Nieto, C., Libby, S. J., Wing, L. & Gould, J. Describing the sensory abnormalities of children and adults with autism. *J. Autism Dev. Disord.* **37**, 894–910 (2007).
86. Scharre, J. E. & Creedon, M. P. Assessment of visual function in autistic

- children. *Optom. Vis. Sci.* **69**, 433–439 (1992).
87. Apicella, F., Costanzo, V. & Purpura, G. Are early visual behavior impairments involved in the onset of autism spectrum disorders? Insights for early diagnosis and intervention. *European Journal of Pediatrics* vol. 179 225–234 (2020).
88. Kern, J. K. *et al.* Sensory correlations in autism. *Autism* vol. 11 123–134 (2007).
89. Jones, W. & Klin, A. Attention to eyes is present but in decline in 2-6-month-old infants later diagnosed with autism. *Nature* **504**, 427–431 (2013).
90. McPartland, J. C. *et al.* Looking Back at the Next 40 Years of ASD Neuroscience Research. *J. Autism Dev. Disord.* (2021) doi:10.1007/s10803-021-05095-5.
91. Baird, G., Cass, H. & Slonims, V. Diagnosis of autism. *BMJ* **327**, 488–493 (2003).
92. Zwaigenbaum, L. *et al.* Stability of diagnostic assessment for autism spectrum disorder between 18 and 36 months in a high-risk cohort. *Autism Res.* **9**, 790–800 (2016).
93. Emerson, R. W. *et al.* Functional neuroimaging of high-risk 6-month-old infants predicts a diagnosis of autism at 24 months of age. *Sci. Transl. Med.* **9**, (2017).
94. Hazlett, H. C. *et al.* Early brain development in infants at high risk for autism spectrum disorder. *Nature* **542**, 348–351 (2017).
95. Ruta, L., Mazzone, D., Mazzone, L., Wheelwright, S. & Baron-Cohen, S. The Autism-Spectrum Quotient--Italian version: a cross-cultural confirmation of the broader autism phenotype. *J. Autism Dev. Disord.* **42**, 625–633 (2012).
96. Vrana, A., Meier, M. L., Hotz-Boendermaker, S., Humphreys, B. K. & Scholkman, F. Cortical Sensorimotor Processing of Painful Pressure in Patients with Chronic Lower Back Pain-An Optical Neuroimaging Study using fNIRS. *Front. Hum. Neurosci.* **10**, 578 (2016).
97. Zimeo Morais, G. A., Balardin, J. B. & Sato, J. R. fNIRS Optodes' Location Decider (fOLD): a toolbox for probe arrangement guided by brain regions-of-interest. *Sci. Rep.* **8**, 3341 (2018).
98. Peirce, J. *et al.* PsychoPy2: Experiments in behavior made easy. *Behav. Res. Methods* **51**, 195–203 (2019).
99. Gollapudi, S. OpenCV with Python. *Learn Computer Vision Using OpenCV* 31–50 (2019) doi:10.1007/978-1-4842-4261-2_2.
100. Vallat, R. Pinguin: statistics in Python. *Journal of Open Source Software* vol. 3 1026 (2018).

101. Hunter, J. D. Matplotlib: A 2D Graphics Environment. *Computing in Science & Engineering* vol. 9 90–95 (2007).

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Figure legends

Fig.1: Visual stimulation and experimental paradigm. A: Representative frame of baseline grey screen (upper row, stimulus ‘off’) and reversing checkerboard (lower row, stimulus ‘on’) for RS condition. The small black square indicates the fixation point. **B:** Representative frame of low-contrast (20%) grey-scale baseline animated cartoon (upper row, stimulus ‘off’) and blended checkerboard-cartoon (lower row, stimulus ‘on’) for CF and CC conditions. **C:** Representative HDR in the occipital cortex during the stimulus ‘off’ (upper row) and stimulus ‘on’ activation phase (lower row) according to the output of nirsLAB software. The Look Up table is reported under the images. **D:** Experimental protocol showing that the cycles of visual stimulation were structured in blocks of 40 trials (20 trials with the reversing checkerboard and 20 trials with the ‘mock’ stimulus) for a total duration of 10 minutes.

Fig. 2: HDR was reliably detected in adults using both RS and blended RS-animated cartoons. For all panels, values in the y-axis are multiplied for 10^4 . **A:** On the left, the average time course for THb (green line), OHb (red line) and DHb (blue line) in response to the RS are shown. The three plots on the right depict the

average peak response to the stimulus (S) vs. the blank (B) across all the adult subjects. The stimulus-driven signal was significantly different from the blank for all the conditions (t-test, $p < 0.001$ for all comparisons). **B:** Same plots as above for the CF condition. On the left, the average time course of the evoked HDR is depicted. On the right, the graphs showed that the HDR amplitude was significantly higher in response to S with respect to B for THb, OHb and DHb (t-test, $p < 0.001$ for all comparisons). **C:** CC condition. Also in this case the S elicited significantly higher responses for THb, OHb and DHb with respect to the B (t-test, $p < 0.001$ for all comparisons). **D:** Comparison among different visual stimulations (RS: Radial Stimulus, CF: fixed cartoon, CC: chosen cartoon) shows no differences in evoked amplitudes for THb, whereas a significant difference was detected between RS and CC for OHb (One-way RM ANOVA, $p < 0.01$, post hoc BH-FDR, RS vs. CC $p < 0.01$) and a more complex pattern of differences emerged for DHb (One-way RM ANOVA, $p < 0.001$, post hoc BH-FDR, RS vs. CF $p < 0.01$, RS vs. CC $p < 0.001$, CF vs CC $p < 0.05$). **E:** No differences of evoked responses were detected with different contrast levels of the baseline movie (L: low, M: medium, H: high). For statistical metrics and details, refer to table S1. Data are shown as average $\pm$ s.e.m. * $p < 0.05$; ** $p < 0.01$; *** $p < 0.001$.

Fig. 3: HDR signal was reliably detected in children using blended RS-animated cartoons with low and high contrast. For all panels, values in the y-axis are multiplied for 10^4 . **A:** On the left, the average time course for THb (green line), OHb (red line) and DHb (blue line) in response to low-contrast (20%) blended RS-animated cartoons is shown. On the right, the graphs represent the average amplitude of the evoked HDR following the stimulus (S) and the blank (B). A significantly different response to S with respect to the B was detectable for all metrics (t-test, $p < 0.001$ for all comparisons). **B:** The average time course of the HDR to high-contrast (80%) blended RS-animated cartoons is shown. Here, the baseline cartoon is the same as the experiment described in panel A, but a different part of the movie was used. THb, OHb and DHb showed a significantly higher deflection to the S with respect to the B in this condition as well (t-test, $p < 0.001$ for all comparisons). **C:** On the left, the average time course of HDR following the second low-contrast blended RS-animated cartoon selected by the subject. On the right, the analysis of peak amplitudes revealed significantly higher responses during

S compared to B for THb, OHb and DHb (t-test, $p < 0.001$ for all comparisons). **D:** Comparison among different contrast levels of the baseline cartoon revealed no differences in the amplitude of HDR. **E:** Response amplitudes for low-contrast blended RS-animated cartoons in adults and children. More specifically, we compared the response to CF condition of adults with L1 condition for children. The average amplitude of HDR was significantly higher in children (t-test, $p < 0.001$ for THb, $p < 0.05$ for OHb, $p < 0.01$ for DHb). For statistical metrics and details, refer to table S1. Data are shown as average $\pm$ s.e.m. * $p < 0.05$; ** $p < 0.01$; *** $p < 0.001$.

Fig. 4: Correlation between HDR and AQ scores. For all panels, values in the x-axis are multiplied for 10^4 . The ρ (rho) index in each plot indicates the Spearman correlation value. **A-C:** Correlation between HDR and AQ scores in adults, for amplitudes obtained using RS (**A**), CF (**B**), and CC (**C**). No significant correlations were detected for adult participants. **D-E:** Correlation between HDR and AQ scores in children, for amplitudes obtained using high (**D**), and low (**E**) contrast baseline cartoons. A significant correlation was found between THb and AQ scores for both high- and low-contrast blended stimuli ($p < 0.05$ for both cases). Circles are individual values, lines represent the linear regression model fit and shaded regions are the 95% CI.

Fig. 5: Correlation between HDR and AQ subscales in children. For all panels, values in the x-axis are multiplied for $\times 10^4$. The ρ (rho) index in each plot indicates the Spearman correlation value. **A-B:** Correlations between HDR and AQ Social Skills (AQ_S) subscale. A significant correlation between THb and AQ_S was detected using both high- (A) and low-contrast blended stimuli (B; $p < 0.05$ for both cases). In addition, OHb recorded in response to the low-contrast blended RS-cartoon was significantly correlated with AQ_S (B; $p < 0.05$). **C-D:** Correlations between HDR and AQ Communication (AQ_C) subscale. THb amplitude in response to the high-contrast blended RS-cartoon was significantly correlated with AQ_C ($p < 0.05$). Circles are individual values, lines represent the linear regression model fit and shaded regions are the 95% CI.

Table 1: Demographic characteristics of adult subjects. Age (years), gender, head circumference (head, cm), cap size (cm), total AQ score (AQ), AQ subscale scores

(AQ_S, AQ_C, AQ_A, AQ_D, AQ_I) and the movies used for visual stimulation (CF and CC according to the experimental protocol) are listed for each participant. For movies, production company, release date and episode title are indicated as well.

Table 2: Demographic characteristics of children. Age (years), gender, head circumference (head, cm), cap size (cm), total AQ score (AQ), AQ subscale scores (AQ_S, AQ_C, AQ_A, AQ_D, AQ_I) and the movies used for visual stimulation (C1 and C2 according to the experimental protocol) are listed for each participant. For movies, production company, release date and episode title are indicated as well.

Fig 1

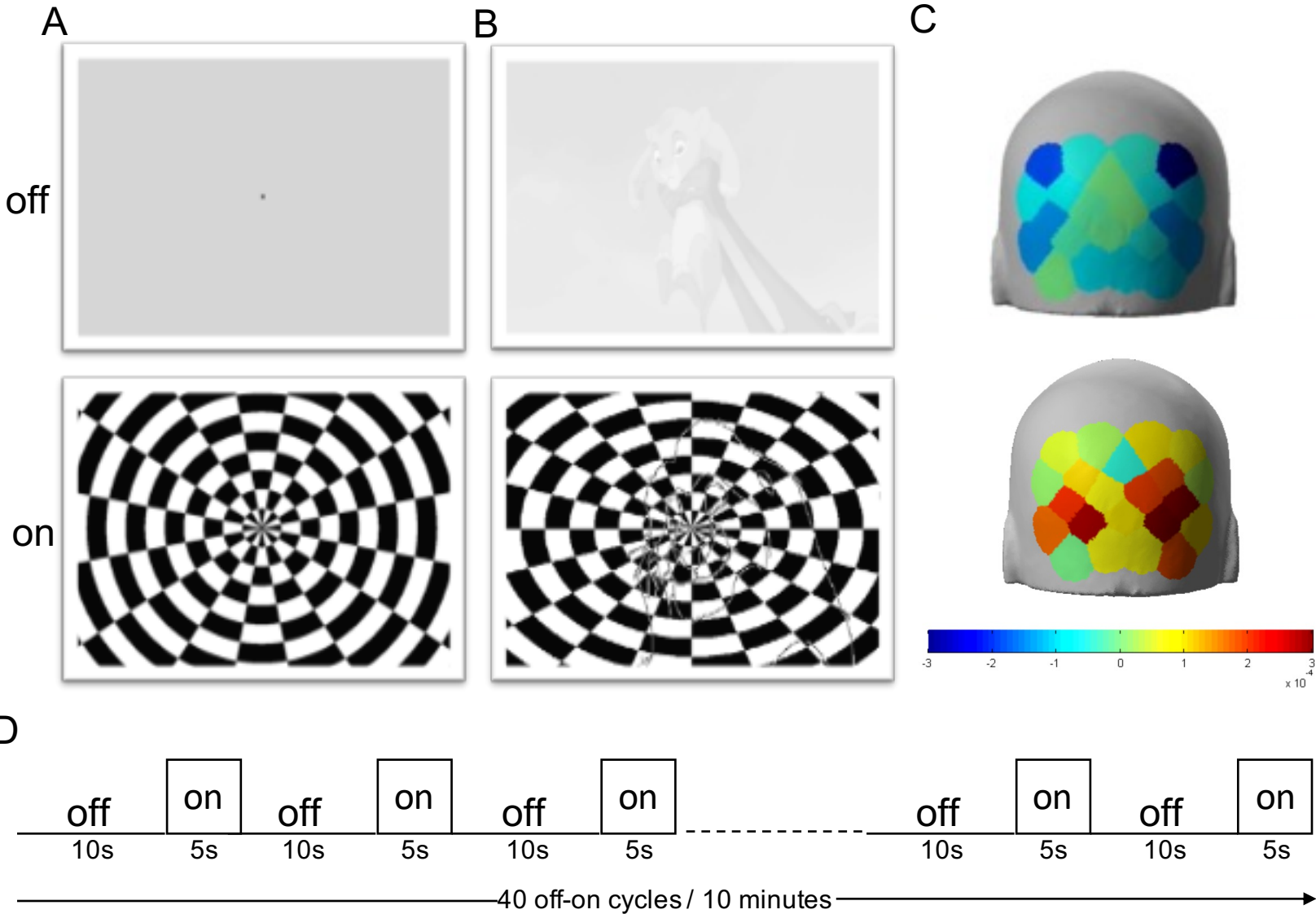


Fig 2

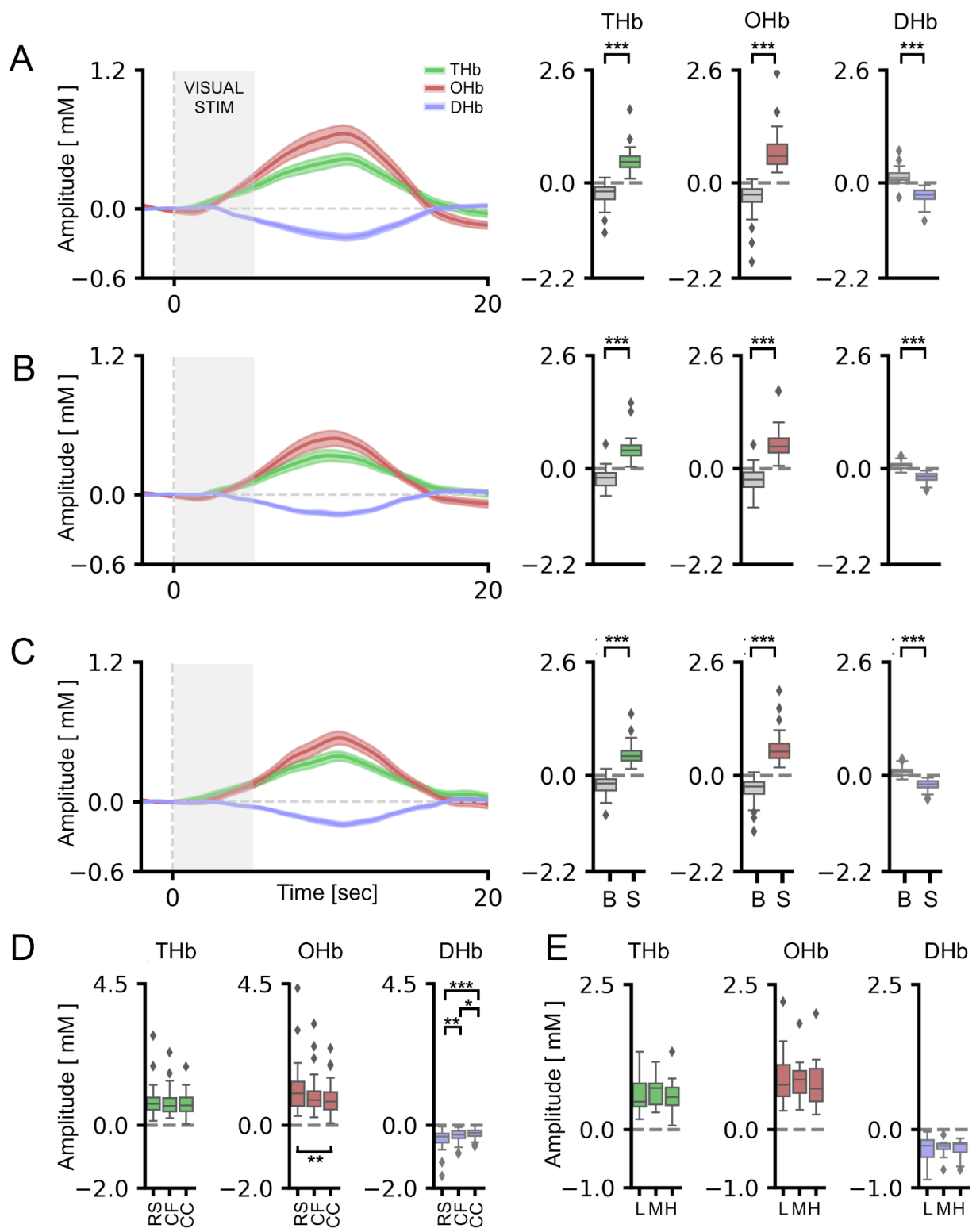


Fig 3

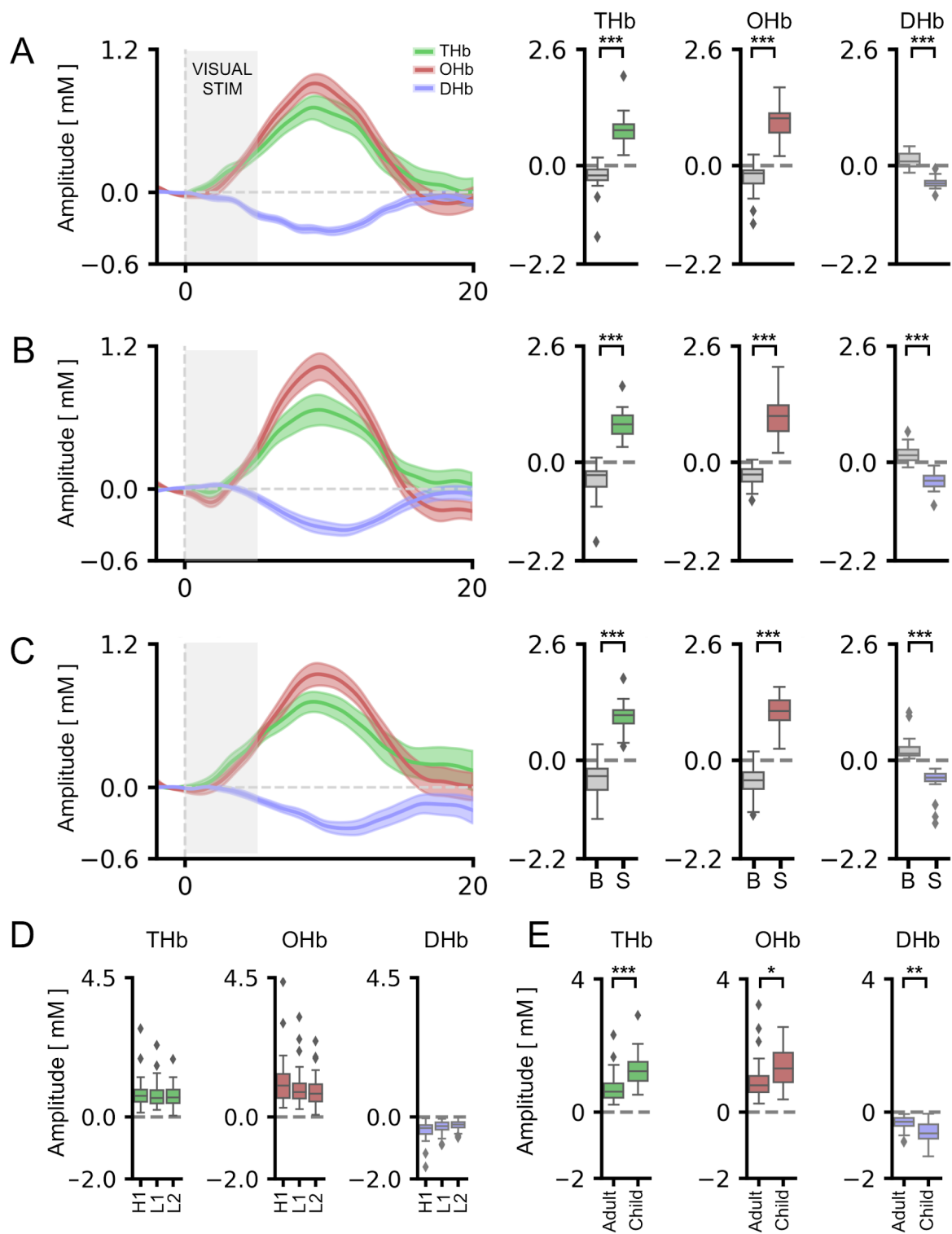


Fig 4

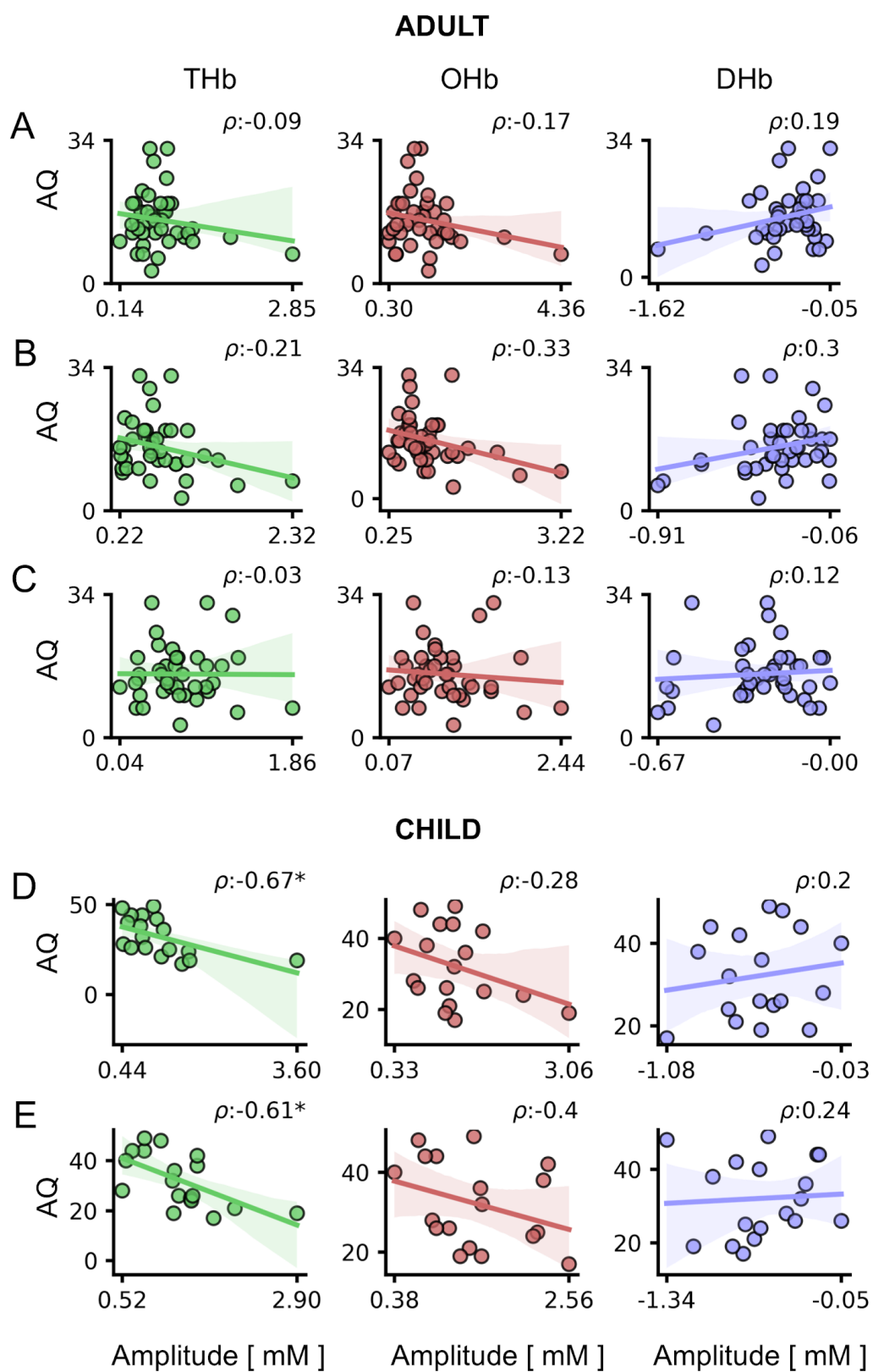
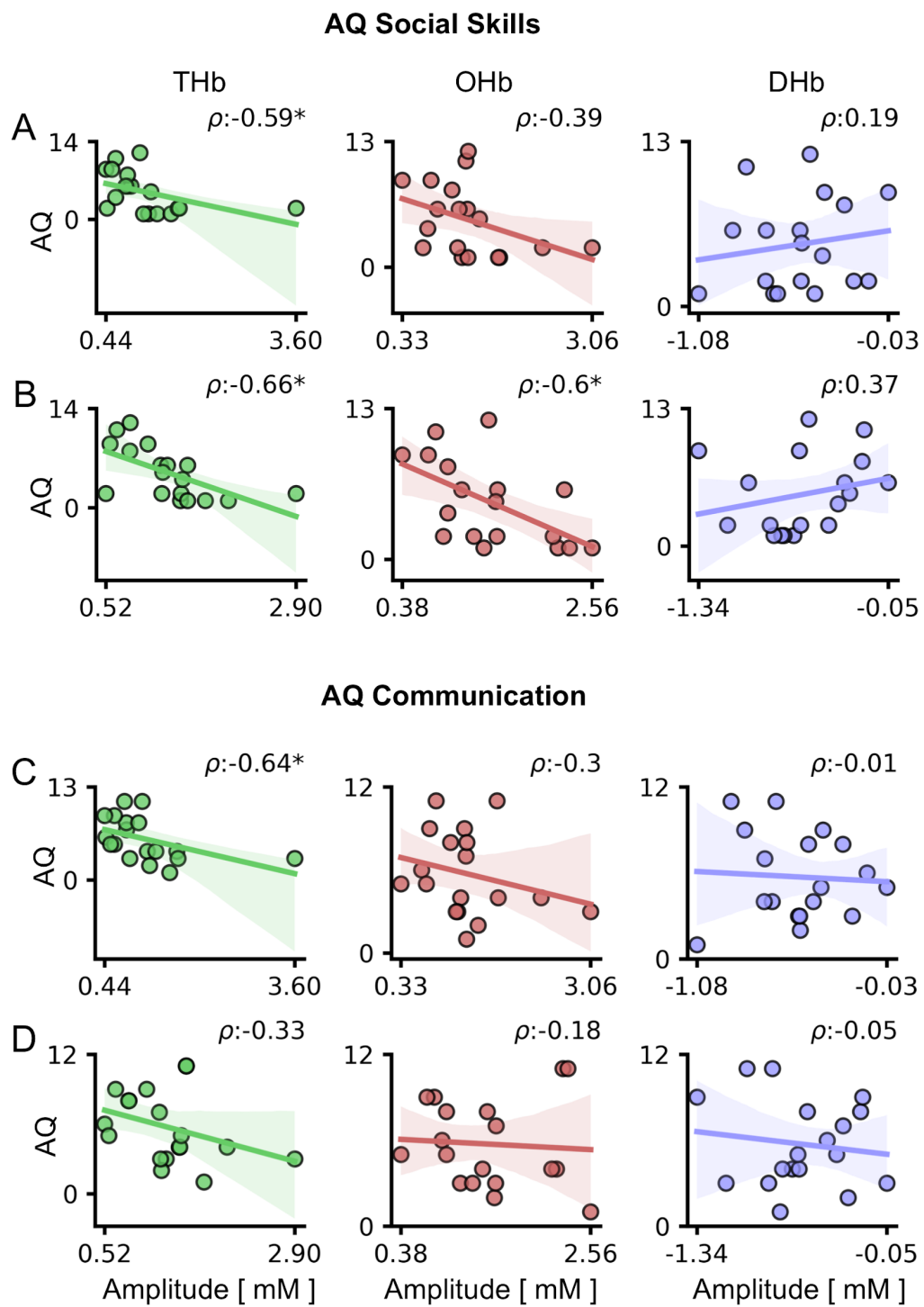


Fig 5



ID	Age	Gender	Head	Cap size	AQ	AQ_S	AQ_C	AQ_A	AQ_D	AQ_I	CF	CC
A1	29	F	57	56	11	0	1	7	1	2	The Lion King Walt Disney Pictures,1994	The Sword in the Stone Walt Disney Productions,1963
A2	36	F	56	56	17	0	0	6	9	2	The Powerpuff Girls special episode "Twas the Fight Before Christmas", Cartoon Network Studios, 2003	The Aristocats Walt Disney Productions,1970
A3	35	M	57	56	17	3	3	5	0	6	Kung Fu Panda DreamWorks Animation, 2008	Aladdin Walt Disney Pictures,1992
A4	28	F	57	56	3	0	0	1	1	1	Peppa Pig "Hide-and-seek", "Fly the kite", "Polly parrot" episodes, Entertainment One, 2004	The Simpsons S29E02 "Springfield Splendor", 20th Television, 2017
A5	25	F	57	56	19	4	1	7	6	1	The Lion King Walt Disney Pictures,1994	101 Dalmatians Walt Disney Productions,1961
A6	35	F	57	56	15	4	2	4	5	0	The Powerpuff Girls special episode "Twas the Fight Before Christmas", Cartoon Network Studios, 2003	The Simpsons S29E02 "Springfield Splendor", 20th Television, 2017
A7	29	F	57	56	11	0	2	5	3	1	Peppa Pig "Hide-and-seek", "Fly the kite", "Polly parrot" episodes, Entertainment One, 2004	101 Dalmatians Walt Disney Productions,1961
A8	30	F	56	56	13	0	2	4	5	2	Kung Fu Panda DreamWorks Animation, 2008	Wall-E Disney-Pixar, 2008
A9	29	F	59	56	7	1	1	3	2	0	The Lion King Walt Disney Pictures,1994	The Sword in the Stone Walt Disney Productions,1963
A10	29	M	58	56	12	1	1	5	4	1	The Powerpuff Girls special episode "Twas the Fight Before Christmas", Cartoon Network Studios, 2003	Aladdin Walt Disney Pictures,1992
A11	27	M	56.5	56	17	2	3	5	4	3	Peppa Pig "Hide-and-seek", "Fly the kite", "Polly parrot" episodes, Entertainment One, 2004	101 Dalmatians Walt Disney Productions,1961
A12	30	F	56	56	9	1	2	2	3	1	Kung Fu Panda DreamWorks Animation, 2008	Inside out Disney-Pixar, 2015
A13	29	M	57	56	14	1	3	6	2	2	The Lion King Walt Disney Pictures,1994	Wall-E Disney-Pixar, 2008
A14	36	M	58.5	56	19	2	2	5	7	3	The Powerpuff Girls special episode "Twas the Fight Before Christmas", Cartoon Network Studios, 2003	101 Dalmatians Walt Disney Productions,1961
A15	36	M	57	56	16	4	1	4	5	2	Peppa Pig "Hide-and-seek", "Fly the kite", "Polly parrot" episodes, Entertainment One, 2004	Aladdin Walt Disney Pictures,1992
A16	29	F	54	56	22	1	3	9	7	2	Kung Fu Panda DreamWorks Animation, 2008	The Rescuers Walt Disney Productions,1977
A17	28	F	58.5	56	7	1	0	2	1	3	The Lion King Walt Disney Pictures,1994	101 Dalmatians Walt Disney Productions,1961
A18	29	F	55	56	12	0	1	5	5	1	The Powerpuff Girls special episode "Twas the Fight Before Christmas", Cartoon Network Studios, 2003	Peter Pan Walt Disney Productions,1953
A19	34	M	60	56	19	6	2	5	5	1	Peppa Pig "Hide-and-seek", "Fly the kite", "Polly parrot" episodes, Entertainment One, 2004	SpongeBob S12E01 "FarmerBob", Nickelodeon Animation Studio, 2018
A20	26	M	58	56	10	3	0	5	2	0	Kung Fu Panda DreamWorks Animation, 2008	Inside out Disney-Pixar, 2015

A21	36	F	56	56	15	1	2	2	7	3	The Lion King Walt Disney Pictures,1994	Inside out Disney-Pixar, 2015
A22	30	M	60.5	56	19	1	2	6	4	6	The Lion King Walt Disney Pictures,1994	The Sword in the Stone Walt Disney Productions,1963
A23	26	F	56	56	12	0	1	5	6	0	The Powerpuff Girls special episode "Twas the Fight Before Christmas", Cartoon Network Studios, 2003	Wall-E Disney-Pixar, 2008
A24	32	M	60.5	56	7	0	1	5	0	1	Kung Fu Panda DreamWorks Animation, 2008	Aladdin Walt Disney Pictures,1992
A25	34	F	56	56	10	1	1	4	2	2	The Lion King Walt Disney Pictures1994	Beauty and the Beast Walt Disney Pictures,1994
A26	29	F	57	56	32	6	5	8	9	4	The Powerpuff Girls special episode "Twas the Fight Before Christmas", Cartoon Network Studios, 2003	The Aristocats Walt Disney Productions,1970
A27	31	M	59	56	32	10	6	6	2	8	Kung Fu Panda DreamWorks Animation, 2008	The Simpsons S29E02 "Springfield Splendor", 20th Television, 2017
A28	39	F	57	56	21	2	4	4	8	3	Peppa Pig "Hide-and-seek", "Fly the kite", "Polly parrot" episodes, Entertainment One, 2004	Lady Oscar, "A Funeral Bell Tolls in the Twilight" episode, Discotek Media, 1980
A29	29	M	57	56	6	0	0	2	2	2	The Lion King Walt Disney Pictures,1994	The Simpsons S29E02 "Springfield Splendor", 20th Television, 2017
A30	29	M	57	56	17	3	2	5	6	1	The Powerpuff Girls special episode "Twas the Fight Before Christmas", Cartoon Network Studios, 2003	The Sword in the Stone Walt Disney Productions,1963
A31	36	M	59.5	56	12	0	0	6	4	2	Peppa Pig "Hide-and-seek", "Fly the kite", "Polly parrot" episodes, Entertainment One, 2004	The Sword in the Stone Walt Disney Productions,1963
A32	27	M	59	56	12	1	2	5	2	2	The Powerpuff Girls special episode "Twas the Fight Before Christmas", Cartoon Network Studios, 2003	The Aristocats Walt Disney Productions,1970
A33	40	M	57	56	13	2	3	1	6	1	Peppa Pig "Hide-and-seek", "Fly the kite", "Polly parrot" episodes, Entertainment One, 2004	The Simpsons S29E02 "Springfield Splendor", 20th Television, 2017
A34	31	F	55	56	15	1	1	4	4	5	Kung Fu Panda DreamWorks Animation, 2008	The Sword in the Stone Walt Disney Productions,1963
A35	27	M	56.5	56	19	6	2	4	7	0	The Lion King Walt Disney Pictures,1994	Frozen Walt Disney Pictures, 2013
A36	30	M	58	56	14	5	1	3	4	1	The Powerpuff Girls special episode "Twas the Fight Before Christmas", Cartoon Network Studios, 2003	Wall-E Disney-Pixar, 2008
A37	25	M	58	56	29	6	8	6	7	2	Peppa Pig "Hide-and-seek", "Fly the kite", "Polly parrot" episodes, Entertainment One, 2004	Wall-E Disney-Pixar, 2008
A38	31	M	57.5	56	10	0	1	5	3	1	Kung Fu Panda DreamWorks Animation, 2008	Inside out Disney-Pixar, 2015
A39	38	M	57.5	56	25	2	4	6	8	5	The Lion King Walt Disney Pictures,1994	The Sword in the Stone Walt Disney Productions,1963
A40	33	F	57.5	56	13	0	0	3	10	0	The Powerpuff Girls special episode "Twas the Fight Before Christmas", Cartoon Network Studios, 2003	Beauty and the Beast Walt Disney Pictures,1994

ID	Age	Gender	Head	Cap size	AQ	AQ-S	AQ_C	AQ_A	AQ_D	AQ_I	C1 (L1 and H1)	C2 (L2)
B1	5	M	51.5	52	21	1	4	7	5	4	Uncle Grandpa S3E04 "Uncle Easter", Cartoon Network Studios, 2016	Teen Titans Go! To the Movies Warner Bros. Animation, 2018
B2	5	M	51	52	32	6	7	3	10	6	Wile E. Coyote & Road Runner "Coyote falls", ""Fur of flying" and "Rabid rider" episodes, Acme Corporation, 2010	ABCs song Little Baby Bum - Nursery Rhymes & Kids Songs youtube channel, 2014
B3	13	M	57	56	26	6	3	4	7	6	The Simpsons S29E02 "Springfield Splendor", 20th Television, 2017	Futurama, S7E01 "The Bots and the Bees", Comedy Central, 2012
B4	12	M	56	56	44	8	8	11	13	4	The Incredibles Disney-Pixar, 2004	Wile E. Coyote & Road Runner "Coyote falls", ""Fur of flying" and "Rabid rider" episodes, Acme Corporation, 2010
B5	12	M	56	56	25	1	4	10	0	10	Big Hero 6 Walt Disney Pictures, 2014	The Amazing World of Gumball S01E01 "The DVD", Cartoon Network Development Studio Europe, 2011
B6	6	F	53	52	17	1	1	4	8	3	Inside out Disney-Pixar, 2015	Floopaloo S1E22 "Squirrel for a Day", Marc du Pontavice, 2012
B7	7	M	53	52	24	2	4	1	12	5	The Chipmunks "Bye, George" episode, DIC Entertainment, 1989	Floopaloo S1E22 "Squirrel for a Day", Marc du Pontavice, 2012
B8	4	F	50.5	52	38	6	11	7	11	3	Frozen Walt Disney Pictures, 2013	Bolt Walt Disney Pictures, 2008
B9	4	M	52	52	42	1	11	11	13	6	Spider-Man: Into the Spider-Verse Columbia Pictures, 2018	Bolt Walt Disney Pictures, 2008
B10	4	M	51	52	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.	Cars Walt Disney Pictures, 2006	Thomas & Friends "Diesel and the duckling" episode, Mattel, 2019
B11	9	F	54	56	28	2	6	3	13	4	The Simpsons S29E02 "Springfield Splendor", 20th Television, 2017	Zig & Sharko S01E28 "Moby Zig", Xilam Animation, 2010
B12	8	M	54.5	56	26	4	5	5	4	8	Beauty and the Beast Walt Disney Pictures, 1994	Ice Age Blue Sky Studios, 2002
B13	4	M	53	56	44	11	9	8	9	7	Finding Nemo Disney-Pixar 2003	101 Dalmatians Walt Disney Productions, 1961
B14	6	M	53	52	36	5	2	3	19	7	Trolls DreamWorks Animation, 2016	Curious George S1E03 "Zeros to Donuts", Universal Animation Studios, 2006
B15	7	M	51.5	52	49	12	8	10	9	10	Cars Walt Disney Pictures, 2006	Pup Academy S1E02 "Tell Us About Your Human Day", Air Bud Entertainment, 2019
B16	10	F	55	56	48	9	9	10	10	10	Frozen Walt Disney Pictures, 2013	Descendants Disney Channel Original Productions, 2015
B17	10	M	54	56	40	9	5	7	13	6	The Simpsons S29E02 "Springfield Splendor", 20th Television, 2017	Uncle Grandpa S3E04 "Uncle Easter", Cartoon Network Studios, 2016
B18	4	F	49.5	52	19	2	3	5	6	3	Frozen Walt Disney Pictures, 2013	Bing S1E06 "Smoothie", Tandem Films e Digitales Studios, 2014
B19	7	M	54	56	19	2	3	5	6	3	Ranger Rob S1E12 "Big Stink in Big Sky Park" Nelvana Enterprises Inc, 2016	Arex&Vastatore "Casket of fear" episode Arex&Vastatore YouTube Channel, 2021