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Enhanced Visual Search in Infancy Predicts Emerging Autism Symptoms

Highlight
- We measured visual search abilities in infants at familial risk for autism
- Enhanced visual search at 9 months predicted a higher level of autism symptoms at 2 years
- Atypical perception is intrinsically linked to the emerging autism phenotype

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In Brief
It remained unknown whether superior perception, a common feature of the autism phenotype, contributes to the emergence of core social interaction and communication symptoms. Gliga et al. show that superior performance in a visual search task in 9-month-old infants predicts a higher level of autism symptoms at 15 months and 2 years.
Enhanced Visual Search in Infancy Predicts Emerging Autism Symptoms

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SUMMARY

In addition to core symptoms, i.e., social interaction and communication difficulties and restricted and repetitive behaviors, autism is also characterized by aspects of superior perception [1]. One well-replicated finding is that of superior performance in visual search tasks, in which participants have to indicate the presence of an odd-one-out element among a number of foils [2–5]. Whether these aspects of superior perception contribute to the emergence of core autism symptoms remains debated [4, 6]. Perceptual and social interaction atypicalities could reflect co-occurring but biologically independent pathologies, as suggested by a “fractionable” phenotype model of autism [7]. A developmental test of this hypothesis is now made possible by longitudinal cohorts of infants at high risk, such as of younger siblings of children with autism spectrum disorder (ASD). Around 20% of younger siblings are diagnosed with autism themselves [8], and up to another 30% manifest elevated levels of autism symptoms [9]. We used eye tracking to measure spontaneous orienting to letter targets (O, S, V, and +) presented among distractors (the letter X; Figure 1). At 9 and 15 months, emerging autism symptoms were assessed using the Autism Observation Scale for Infants (AOSI; [10]), and at 2 years of age, they were assessed using the Autism Diagnostic Observation Schedule (ADOS; [11]). Enhanced visual search performance at 9 months predicted a higher level of autism symptoms at 15 months and at 2 years. Infant perceptual atypicalities are thus intrinsically linked to the emerging autism phenotype.

RESULTS AND DISCUSSION

Eighty-two high-risk infants (37 girls) and 27 low-risk controls (13 girls) took part in this study (Table S1). We analyzed the proportion of trials in which infants made a first look toward one of the targets, after fixating at the center of the screen. Infants with at least four valid trials were included in the analysis. Above-chance performance was measured in the group as a whole, at all ages (9 months: t(103) = 5.62, p < 0.001; 15 months: t(95) = 4.31, p < 0.001; 2 years: t(94) = 7.9, p < 0.001; Table S2). Performance was not related to either the age or IQ of the participant (all p > 0.09) during any of the visits.

A shift from categorical to continuous characterization of psychopathology is encouraged by clinical and genetics research [12, 13]. Thus, to take into account longitudinal relationships between visual search performance and emerging autism symptoms (Figure 1), we entered search performance at 9 months, 15 months, and 2 years in an autoregressive model (Figure 2; model fit: $\chi^2(4) = 6.87, p = 0.14$, comparative fit index [CFI] = 0.95) with continuous measures of symptom severity at 9 and 15 months (Autism Observation Scale for Infants [AOSI] score) and 2 years of age (Autism Diagnostic Observation Schedule [ADOS] score). Nine-month visual search significantly predicted the 15-month AOSI ($\beta = 0.22$, SE = 0.10, p = 0.03) and the 2-year ADOS ($\beta = 0.24$, SE = 0.10, p = 0.02; see also Figure S2) score, with increased visual search accuracy predicting higher symptom severity. Findings were very similar when only the high-risk group was included in the analysis (model fit: $\chi^2(4) = 10.001$, p = 0.04, CFI = 0.88). The relationship with ADOS remained substantively similar: 9-month visual search was significantly related to both 15-month AOSI ($\beta = 0.223$, SE = 0.11, p = 0.049) and 2-year ADOS ($\beta = 0.27$, SE = 0.11, p = 0.02) scores. To test whether visual search at 9 months continued to predict ADOS score after accounting for earlier autism markers, we ran an autoregressive model with regressions, rather than correlations, between AOSI and ADOS (model fit: $\chi^2(4) = 6.87$, p = 0.14, CFI = 0.95; Figure S1). The relationship between 9-month visual search and 15-month AOSI remained significant ($\beta = 0.182$, SE = 0.09, p = 0.046), but the direct relationship with later ADOS (i.e., accounting for 9- and 15-month AOSI) became nonsignificant ($\beta = 0.13$, SE = 0.09, p = 0.13), suggesting a developmental pathway in which infant visual search contributes to autism symptoms at 15 months of age and that in turn contributes to autism severity at 2 years of age. The lack of a concurrent relationship between visual search and symptom severity at 9 months ($\beta = 0.08$, SE = 0.10, p = 0.44) is suggestive of a causal pathway from early perception to later emerging autism symptoms. Moreover, although 9- and 15-month visual search performances are correlated ($\beta = 0.24$, SE = 0.10, p = 0.02), performance at later time points, i.e., at 15 months and 2 years of age, does not relate to symptomatology. This differential relationship points to particular periods in early postnatal development within which atypical perception, in addition to other risk factors, may set development on a pathway to pathology and...
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rather than causing/explaining, deficits in social cognition” [6].
aspect of cognition in autism spectrum disorder (ASD) alongside,
connectivity [17] during the first year of life of those infants that
later develop autism symptoms, by suggesting that perturbations
in oculomotor behavior described in this population [14]. Also,
given a higher incidence of hyperlexia in autism [15], future
studies should address the question of whether the demonstrat-
ated superior visual search also predicts better recognition
of letters later in childhood.
Views on atypical perception have alternated between assign-
ing it a core, causal role in autism [16] and portraying it as “one
aspect of cognition in autism spectrum disorder (ASD) alongside,
rather than causing/explaining, deficits in social cognition” [6].
Importantly, our findings corroborate evidence for atypical oculo-
motor behavior [14] and increased frontal-occipital functional
connectivity [17] during the first year of life of those infants that
later develop autism symptoms, by suggesting that perturbations
in general processes, such as perception or attention, are more
important than previously believed in the developmental pathway
to this disorder [18]. With this shift away from “social brain” the-
ories of autism (e.g., [19]) comes also the challenge of explaining
the mechanisms through which domain-general atypicalities
could contribute to the emergence of specific autism symptoms.
Moreover, the striking predictive association between superior
visual search and autism may also prove useful as one additional
component of early autism identification, given a context in which
most current infant markers are based on impairments common
to multiple neurodevelopmental outcomes (e.g., [20–22]).

EXPERIMENTAL PROCEDURES

Participants
Participants took part in a longitudinal study of children at risk for autism. At the
time of enrollment, none of the infants had been diagnosed with any medical or
developmental condition. Twenty-seven low-risk participants and 82 high-risk
participants took part in this study. High-risk infants had at least one older sibling
(hereafter, proband) with a community clinical diagnosis of ASD. Proband diag-
nosis was confirmed by an expert clinician (T.C.) based on information using the
Development and Well-Being Assessment (DAWBA; [23]) and the parent-report
Social Communication Questionnaire (SCQ; [24]). Parent-reported family med-
ical histories were examined for significant medical conditions in the proband or
extended family members, with no exclusions made on this basis. Infants in the
low-risk control group were recruited from a volunteer database. Inclusion
criteria included full-term birth, normal birth weight, and lack of any ASD within
first-degree family members (as confirmed through parent interview regarding
family medical history). All low-risk participants had at least one older sibling.
Screening for possible ASD in these older siblings was undertaken using the
SCQ, with no child scoring above instrument cut-off for ASD. The data pre-
sented in this paper were collected during three consecutive visits, at around
9 months, 15 months, and 2 years of age. All but two low-risk participants
and all but two high-risk participants contributed data from at least two visits.
General and visit-specific participant characteristics are presented in Table S1.

Stimuli and Procedure
We created arrays of eight letters, situated on an imaginary circle and on a
white background. In each array, all but one stimulus was an “X” letter. The
eightth stimulus was either a “+,” a “V,” an “S,” or an “O” (the targets). For
each target type, eight different arrays were created, varying in the position
of the target, i.e., 32 different stimuli in total. To increase variability, we used
letters in an array that were black, blue, red, or green (25% of arrays for
each color). Because of time constraints, only 50% of the stimuli were pre-
sented in this paper were collected during three consecutive visits, at around
2 years-old visit. For each target type (+, V, S, or O), we chose
four out of the existing eight stimuli, those where targets were in even area
of interest (AOI) positions on the slide (see Figure 1). Infants were seated on
mother’s lap, at approximately 60 cm from a Tobii T120 screen. A five-point
calibration routine was run. The experiment was started only after at least
four points were marked as being properly calibrated for each eye. The infant’s
behavior was monitored by a video camera placed above the Tobii monitor.
Stimuli were presented with TobiiStudio software. Each of the stimuli was pre-
sented once, in a random order, for 1.5 s. Before each stimulus, the child’s
attention was directed to the center of the screen using a short audio-video an-
imation. Only trials in which the center of the screen was fixated within the first
100 ms of stimulus onset were used for subsequent analysis.

Measures of Autism Symptoms
The AOSI [10] is a validated clinical measure of infant risk markers, focusing on
precursors of impairments present in the ASD phenotype, including response
Figure 2. Relationship between Visual Search and Emerging Autism Symptoms

Visual search performance at 9 months predicts later autism symptom severity in an autoregressive model. Standardized coefficient values are presented for significant results (represented as black arrows).

SUPPLEMENTAL INFORMATION

Supplemental Information includes Supplemental Experimental Procedures, two figures, and three tables and can be found with this article online at http://dx.doi.org/10.1016/j.cub.2015.05.011.

CONSORTIA

The members of The BASIS Team are Simon Baron-Cohen, Patrick Bolton, Celeste Cheung, Kim Davies, Michelle Liew, Janice Fernandes, Issy Gammer, Helen Maris, Erica Salomone, Greg Pasco, Andrew Pickles, Helena Ribeiro, and Leslie Tucker. A full list of affiliations for members of The BASIS Team can be found in Table S3.

AUTHOR CONTRIBUTIONS

T.G. designed the eye-tracking study, T.G. and The BASIS Team collected the data. T.G. and R.B. analyzed the study and wrote the paper, with contribution from M.H.J. and T.C. M.H.J. and T.C. led The BASIS Team and designed the overall BASIS study, with other members of The BASIS Team.

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