Network inefficiencies in autism spectrum disorder at 24 months

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ORIGINAL ARTICLE

Network inefficiencies in autism spectrum disorder at 24 months

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Autism spectrum disorder (ASD) is a developmental disorder defined by behavioral symptoms that emerge during the first years of life. Associated with these symptoms are differences in the structure of a wide array of brain regions, and in the connectivity between these regions. However, the use of cohorts with large age variability and participants past the generally recognized age of onset of the defining behaviors means that many of the reported abnormalities may be a result of cascade effects of developmentally earlier deviations. This study assessed differences in connectivity in ASD at the age at which the defining behaviors first become clear. There were 113 24-month-old participants at high risk for ASD, 31 of whom were classified as ASD, and 23 typically developing 24-month-old participants at low risk for ASD. Utilizing diffusion data to obtain measures of the length and strength of connections between anatomical regions, we performed an analysis of network efficiency. Our results showed significantly decreased local and global efficiency over temporal, parietal and occipital lobes in high-risk infants classified as ASD, relative to both low- and high-risk infants not classified as ASD. The frontal lobes showed only a reduction in global efficiency in Broca’s area. In addition, these same regions showed an inverse relation between efficiency and symptom severity across the high-risk infants. The results suggest delay or deficits in infants with ASD in the optimization of both local and global aspects of network structure in regions involved in processing auditory and visual stimuli, language and nonlinguistic social stimuli.

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INTRODUCTION

Autism spectrum disorder (ASD) is a developmental disorder defined by impairments in social communication and social interaction and a restricted repertoire of activities and interests. Recent research has indicated that abnormalities in connectivity may underlie many of the behavioral symptoms. This research, however, is equivocal with regard to both the nature of connectivity abnormalities and the regions in which they occur, and currently provides little insight into their early developmental origins. Numerous studies have reported long-range under-connectivity in ASD, some have reported short-range over-connectivity accompanying long-range under-connectivity, and some have reported a less clearly divided mixture of over- and under-connectivity. Such connectivity abnormalities have been variously reported for virtually all pairs of brain regions.

This disparate set of findings reflects a variety of important methodological differences between studies, as well as small sample sizes and substantial etiological and clinical heterogeneity. Prominent among the sources of variability is the wide variety of ages of the participants in these various studies. ASD is a developmental disorder involving complex networks comprising multiple spatially distributed brain regions which are developing in the context of a clearly altered growth trajectory. Children with ASD show abnormally rapid brain growth during the first years of life and abnormalities in the development of connectivity. These brain-based abnormalities are likely to contribute to a developmental cascade of behavioral and neural abnormalities. Thus, differences in findings of connectivity abnormalities in ASD may be owing to differences in the ages of the participants in various studies. The multitude of abnormalities in connectivity reported in adults with ASD, or even children with ASD, may tell us little about where in the brain these abnormalities first emerge. Identifying where connectivity abnormalities originate and how they develop has the potential to inform as to their cause, and to suggest more etiologically relevant paths to successful intervention. However, if we are to avoid confusing downstream developmental effects with source effects, this identification must be on the basis of data from individuals with ASD as early in development as possible, and with as little variation in age as possible.

Behavioral abnormalities may appear during the first year of life or early in the second year, but typically the characteristic symptoms of ASD necessary for classification only become stable by the end of the second year. If connectivity abnormalities underlie these symptoms of ASD, then differences in connectivity should also be present by the end of the second year of life, but downstream developmental effects should be minimal. Thus, to localize the early neural abnormalities in ASD, we assess connectivity at 24 months of age.

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We utilize a whole-brain network analysis to assess white matter connectivity abnormalities in three groups of 24-month-olds: those with symptoms of ASD who also have an older sibling with ASD; those having an older sibling with ASD, but who are not themselves developing symptoms of ASD; and typically developing 24-month-olds with an older sibling and no first-degree relative with ASD or intellectual disability. Children with a sibling who has been diagnosed with ASD, but who do not receive a diagnosis themselves, often exhibit milder ASD symptoms. Thus this group might be expected to show a milder form of any connectivity abnormalities found in infants with significant symptoms of ASD.

A network analysis assesses connectivity on a whole-brain basis, rather than in terms of individual connections, thus providing measures which are relevant to overall brain organization. Brain areas are treated as nodes in a graph; the connections between the brain areas constitute the edges of the graph. Properties of the overall network, or of subnetworks, associated with each participant can then be measured, and group differences assessed. This analytic approach has proved effective in characterizing differences in brain networks in a variety of clinical populations, including ASD.

Network analysis methods have evolved over the past 15 years, from straightforward applications of graph theory, which assess only network topology, to more sophisticated approaches which take into account the spatial aspects of connectivity to assess the efficiency of information transfer within the network. Such approaches utilize measures of the length and strength of connections between all pairs of anatomical regions to estimate how efficiently information can be transferred between regions. This provides a way to assess the impact of abnormalities in spatial organization on communication, both across the brain and within local subnetworks. We use probabilistic tractography to estimate the strength of connectivity between all pairs of regions, and the length of the connections between regions. We then assess the group differences in local and global efficiency for each brain region, as well as the relation of efficiency to symptom severity, to determine where communication efficiency might be impaired in the brains of 24-month-olds with symptoms of ASD.

MATERIALS AND METHODS

Participants

The participants for this study were drawn from an ongoing multisite study of brain and behavioral development in infants at high risk for autism due to having an older sibling with ASD, known as the IBIS (Infant Brain Imaging Study), a study funded by the National Institute of Health via an Autism Center of Excellence grant. IBIS has enrolled and collected neuroimaging and behavioral data from these high-risk infants at 6, 12 and 24 months of age, with confirmation upon enrollment of the diagnosis of their older sibling by medical record review and the Autism Diagnostic Interview. Data are also collected for infants deemed at relatively low risk for autism by virtue of having no first-degree relative with ASD or intellectual disability (as assessed by the Social Communication Questionnaire and the Family Interview for Genetic Studies), and an older sibling. A parent or legal guardian gave informed consent. Exclusionary criteria include the following conditions: (1) diagnosis or physical signs of genetic conditions/syndromes (for example, fragile X syndrome), (2) significant medical or neurological conditions affecting growth, development or cognition (for example, CNS infection, seizure disorder, congenital heart disease), (3) sensory impairments such as vision or hearing loss, (4) low birth weight (< 2000) or prematurity (< 36 weeks gestation), (5) evidence of significant perinatal brain injury or exposure to utero neurotoxins (for example, alcohol, selected prescription medications), (6) non-English speaking families, (7) contraindication for MRI (magnetic resonance imaging; for example, metal implants), (8) adopted children and (9) family history of first-degree relative with intellectual disability (not associated with ASD), psychosis, schizophrenia or bipolar disorder. The data were collected at four sites (University of North Carolina at Chapel Hill, University of Washington, Children’s Hospital of Philadelphia, Washington University in St Louis); quality control and pre-processing were done at the University of Utah, the University of North Carolina at Chapel Hill and the Montreal Neurological Institute (McGill University), the Data Coordinating Center. The study was approved by the institutional review boards at all sites.

The subset of data used in the current study includes all 24-month data that passed quality control as of June 2012. These data divide according to whether the infants are high-risk or low-risk, and according to whether they showed ASD symptoms, as indexed by the Autism Diagnostic Observation Schedule (ADOS), above the clinical cutoff for ASD. Participants were administered the ADOS at 24 months of age by trained researchers. Descriptives for the participants are shown in Table 1, with ADOS scores transformed to calibrated severity scores, as per Gotham et al. Note that Table 1 includes high-risk infants above the clinical cutoff for ASD (HRPOS), high-risk infants below the clinical cutoff for ASD (HRNEG) and low-risk infants below the clinical cutoff for ASD (LRNEG). There were three low-risk infants who received ADOS scores above the cutoff for ASD; their data were excluded. All other infants (a total of 136) were included in the analysis.

Imaging and image processing

MRI scans were done at each site on Siemens 3T TIM Trio scanners (Siemens Medical Systems, Erlangen, Germany) with 12-channel head coils. Intra- and inter-site reliability was verified. Data were collected while infants were naturally sleeping. Three types of images were acquired: (1) T1-weighted images using a 3D MPGR sequence (resolution = 1.0 × 1.0 × 1.0 mm; TE = 3.16 ms; TR = 2400 ms; matrix = 224 × 256); (2) T2-weighted images using a 3D FSE sequence (resolution = 1.0 × 1.0 × 1.0 mm; TE = 80 ms; TR = 3200 ms; matrix = 224 × 256); and (3) diffusion-weighted images using a 2D echo planar sequence (resolution = 2.0 × 2.0 × 2.0; TE = 102 ms; TR = 12 800 ms; directions = 25; b-values: unique values evenly distributed between 0 and 1000 s/mm²). Data were quality controlled, starting at the MRI console; where motion artifacts were observed, the scan sequence was repeated to possibly acquire artifact free data.

The T1- and T2-weighted images were subjected to manual quality control, and corrected for geometric distortion. The diffusion-weighted images were cleaned of motion and other artifacts using DTIPrep, which corrects artifacts where possible, and excludes directions from the data when correction is not possible. Only data sets with acceptable T1-, T2- and diffusion-weighted images were included in the analysis.

The distortion-corrected T1- and T2-volumes were then processed with CIVET, a fully automated structural image analysis pipeline developed at the Montreal Neurological Institute. CIVET corrects intensity nonuniformities using N3, aligns the input volumes to the Talairach-like ICBM-152-nl template, classifies the image into white matter, gray matter, cerebrospinal fluid and background; extracts the white matter and pial surfaces, and warps these to a common surface template. The CIVET results were used to construct the seed, stop and target masks for use with FSL’s probtrackx. Seed masks specify the voxels from which tracts propagate; seed masks were the entire white matter. Stop masks specify where tract propagation is halted; stop masks were the voxels on the boundary of the white matter. Target masks determine the interpretation of tracts; target masks were the cortical labels of the automatic anatomical

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Table 1. Sample characteristics

<table>
<thead>
<tr>
<th>Age (months)</th>
<th>HRPOS</th>
<th>HRNEG</th>
<th>LRNEG</th>
</tr>
</thead>
<tbody>
<tr>
<td>N</td>
<td>Mean ± s.d.</td>
<td>N</td>
<td>Mean ± s.d.</td>
</tr>
<tr>
<td>Male</td>
<td>24</td>
<td>24.65 ± 0.64</td>
<td>51</td>
</tr>
<tr>
<td>Female</td>
<td>7</td>
<td>25.05 ± 0.82</td>
<td>31</td>
</tr>
<tr>
<td>ADOS severity</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>—</td>
<td>6.13 ± 1.60</td>
<td>—</td>
</tr>
<tr>
<td>Female</td>
<td>—</td>
<td>5.14 ± 0.86</td>
<td>—</td>
</tr>
</tbody>
</table>

Abbreviations: HRPOS, high-risk infants with ADOS scores above the clinical cutoff for ASD; HRNEG, high-risk infants with ADOS scores below the clinical cutoff for ASD; LRNEG, low-risk infants with ADOS scores below the clinical cutoff for ASD.
labelling (AAL) atlas, a parcelation based on anatomical landmarks, predominately sulci, defined on the common surface template. The diffusion-weighted images were unwarped via nonlinear registration to the distortion-corrected T1-volume, and then preprocessed for probabilistic tractography with FSL’s bedpostX. The default parameters for bedpostX were used: a maximum of two fibers per voxel, a multivariate factor analysis, on the prior for the second fiber and a burn-in of 1000. The unwarped diffusion volumes were also affine registered to the T1-volumes using FSL’s flirt to provide the mapping from the seed, stop and target masks to the diffusion space. Probabilistic tractography, utilizing FSL’s probtrackX was then seeded twice from 10 000 random locations within each voxel of the seed masks: with distance-bias correction to generate the number of tracts connecting voxels in the target mask, and without distance-bias correction to generate the physical lengths of those tracts. These results were then compiled for each AAL region, generating undirected matrices of the total number of connections between each pair of AAL regions, and the mean physical length of those connections. The total number of connections between each pair of AAL regions divided by the average surface area of the two AAL regions is referred to as ‘connection efficiency’. 

Analysis

Network analyses investigate the organization of sets of connections, rather than considering the connections independently. The metric utilized here is efficiency, as defined by Latora and Marchiori. Network efficiency is the capacity to exchange information across a network. Latora and Marchiori defined the efficiency of the communication between two AAL regions as the total number of connections between each pair of AAL regions divided by the shortest path length of those tracts. These results were then compiled for each AAL region, generating undirected matrices of the total number of connections between each pair of AAL regions, and the mean physical length of those connections. The total number of connections between each pair of AAL regions divided by the average surface area of the two AAL regions is referred to as ‘connection efficiency’.

RESULTS

Relative to LRNEG infants, the HRPOS infants showed reductions in nodal local efficiency bilaterally in the temporal and occipital lobes in inferior and medial regions, and predominately in the right hemisphere for lateral regions, extending to the supramarginal gyrus (Figure 1, top left). Relative to LRNEG infants, the HRPOS infants showed reductions in nodal global efficiency in bilateral temporal and occipital regions, extending in the right hemisphere to the angular and supramarginal gyri; the left hemisphere reductions include the pars triangularis and orbital gyri. (Figure 1, bottom left). There were no significant increases in either nodal local or nodal global efficiency in the HRPOS infants relative to the LRNEG infants. The group differences in nodal local efficiency for HRPOS compared with LRNEG infants are similar to those in the comparison of the HRPOS and LRNEG infants, but left lateralized and less extensive. Relative to the HRNEG infants, the HRPOS infants showed reductions in nodal local efficiency over the left hemisphere temporal and occipital lobes, and the precuneus; the right hemisphere differences are mostly nonsignificant.
Figure 1. The comparison of nodal local and nodal global efficiency for LR\textsuperscript{NEG} versus HR\textsuperscript{POS} infants, and for HR\textsuperscript{NEG} versus HR\textsuperscript{POS} infants, in a model with all three groups, controlling for age, sex and site. The top half of the figure shows the $t$- and $p$-statistic for the comparison of nodal local efficiency in each region of the AAL atlas; the bottom half of the figure shows the $t$- and $p$-statistic for the comparison of nodal global efficiency. The left column shows the comparisons for LR\textsuperscript{NEG} versus HR\textsuperscript{POS} infants; the right column for HR\textsuperscript{NEG} versus HR\textsuperscript{POS} infants. A positive $t$-statistic represents a relative reduction in efficiency in HR\textsuperscript{POS} infants. No regions show significantly increased nodal local or global efficiency in HR\textsuperscript{POS} infants. Significant reductions in nodal local efficiency in HR\textsuperscript{POS} compared with LR\textsuperscript{NEG} infants are seen bilaterally in the temporal and occipital lobes in inferior and medial regions, and predominately in the right hemisphere for lateral regions, extending to the supramarginal gyrus. Significant reductions in nodal local efficiency in HR\textsuperscript{POS} compared with HR\textsuperscript{NEG} infants are seen in the left temporal and occipital lobes, extending into the precuneus, and in several posterior regions in the right hemisphere. Significant reductions in nodal global efficiency in HR\textsuperscript{POS} infants compared with LR\textsuperscript{NEG} infants are seen bilaterally in the temporal and occipital lobes, extending in the right hemisphere to the angular and supramarginal gyri. The left pars triangularis and medial orbital gyrus also show reduced nodal global efficiency. Significant reductions in nodal global efficiency in HR\textsuperscript{POS} infants compared with HR\textsuperscript{NEG} infants are seen in bilateral temporal lobes, extending in the left hemisphere to the occipital cortex, and in the right hemisphere to the angular and supramarginal gyri. The left hemisphere reductions include the medial temporal and occipital regions, and the pars triangularis. The right hemisphere reductions include the precuneus. Note that in all cases the reductions are predominately posterior. HR\textsuperscript{NEG}, high-risk infants with ADOS scores below the clinical cutoff for ASD; HR\textsuperscript{POS}, high-risk infants with ADOS scores above the clinical cutoff for ASD; LR\textsuperscript{NEG}, low-risk infants with ADOS scores below the clinical cutoff for ASD.
Figure 1. The group differences in nodal global efficiency for HRPOS compared with HRNEG infants are also similar to those in the comparison of the HRPOS and LRNEG infants, but with greater reductions in the left hemisphere. Significant reductions in HRPOS infants compared with HRNEG infants are seen in bilateral temporal lobes, extending in the left hemisphere to the occipital cortex, and in the right hemisphere to the angular and supramarginal gyri. The left hemisphere reductions include the pars triangularis. The right hemisphere reductions include the pars triangularis. The top half of the figure shows the t- and p-statistic for the relation of severity and nodal local efficiency in each region of the AAL atlas; the bottom half of the figure shows the t- and p-statistic for the relation of severity and nodal global efficiency. A negative t-statistic represents an inverse relation between severity and efficiency, that is, greater symptom severity corresponds to lesser efficiency. AAL, automatic anatomical labelling; ADOS, Autism Diagnostic Observation Schedule.

Figure 2. The relationship between the measures of network efficiency and the ADOS-based calibrated severity scores within the high-risk infants are similar to the group differences in efficiency for the two subgroups of the high-risk infants, that is, the HRPOS compared with HRNEG infants. There is a significant inverse relation of severity to nodal local efficiency in the left temporal and occipital lobes (Figure 2, top). This inverse relation is significant in the lingual gyrus, and throughout the temporal lobe, with the exception of the middle temporal gyrus. There is a significant inverse relation of severity to nodal global efficiency in the left temporal lobe and in Broca’s area (Figure 2, bottom). This inverse relation is significant throughout the temporal lobe, with the exception of the inferior temporal gyrus. Note that where there is a significant relation between efficiency and severity, in all cases this is an inverse relation, thus greater symptom severity corresponds to lesser efficiency. Also note that, as with the group differences, this inverse relation is only present in posterior regions, and in Broca’s area for nodal global efficiency.

The post hoc analyses of the group differences in the strengths of the connections comprising the paths contributing to the efficiency measures, the tract-based lengths of those paths, and the number of edges in those paths indicates that these results stem from both weaker connections and abnormalities in network structure. The reductions in nodal local and nodal global efficiency in HRPOS infants are partially paralleled by reductions in connection strength in the paths involved. Relative to LRNEG infants, the HRPOS infants showed reductions in nodal local connection strength bilaterally in the temporal, parietal and occipital lobes, and in the left medial orbital gyrus (Figure 3, top left). Relative to LRNEG infants, the HRPOS infants showed reductions in nodal global connection strength in the right inferior temporal and occipital lobes (Figure 3, bottom left). There were no significant increases in either nodal local or nodal global connection strength in the HRPOS infants relative to the LRNEG infants. Relative to HRNEG infants, the HRPOS infants showed reductions in nodal local connection strength in bilateral temporal and parietal lobes, the right precuneus and gyrus rectus, and the left pars triangularis and medial orbital gyrus (Figure 3, top right). Relative to HRNEG infants, the HRPOS infants showed reductions in
Figure 3. The comparison of the mean connection strength in the paths comprising the measures of nodal local and nodal global efficiency for LR\textsuperscript{NEG} versus HR\textsuperscript{POS} infants, and for HR\textsuperscript{NEG} versus HR\textsuperscript{POS} infants, in a model with all three groups, controlling for age, sex and site. The top half of the figure shows the $t$- and $p$-statistic for the comparison of nodal local connection strength in each region of the AAL atlas; the bottom half of the figure shows the $t$- and $p$-statistic for the comparison of nodal global connection strength. The left column shows the comparisons for LR\textsuperscript{NEG} versus HR\textsuperscript{POS} infants; the right column for HR\textsuperscript{NEG} versus HR\textsuperscript{POS} infants. A positive $t$-statistic represents a relative reduction in connection strength in HR\textsuperscript{POS} infants. No regions show significantly increased connection strength in HR\textsuperscript{POS} infants. Significant reductions in nodal local connection strength in HR\textsuperscript{POS} compared with LR\textsuperscript{NEG} infants are seen bilaterally in the temporal, parietal and occipital lobes, and in the left medial orbital gyrus. Significant reductions in nodal local connection strength in HR\textsuperscript{POS} compared with HR\textsuperscript{NEG} infants are seen in bilateral temporal and parietal lobes, the right precuneus and gyrus rectus, and the left pars triangularis and medial orbital gyrus. Significant reductions in nodal global connection strength in HR\textsuperscript{POS} infants compared with LR\textsuperscript{NEG} infants are seen in the right fusiform gyrus and inferior occipital lobe. Significant reductions in nodal global connection strength in HR\textsuperscript{POS} infants compared with HR\textsuperscript{NEG} infants are seen in the right temporal and occipital lobes. Note that, as with efficiency, in all cases the reductions are predominately posterior. HR\textsuperscript{NEG}, high-risk infants with ADOS scores below the clinical cutoff for ASD; HR\textsuperscript{POS}, high-risk infants with ADOS scores above the clinical cutoff for ASD; LR\textsuperscript{NEG}, low-risk infants with ADOS scores below the clinical cutoff for ASD.
Figure 4. The comparison of the mean length of the paths comprising the measures of nodal local and nodal global efficiency for LR$^\text{NEG}$ versus HR$^\text{POS}$ infants, and for HR$^\text{NEG}$ versus HR$^\text{POS}$ infants, in a model with all three groups, controlling for age, sex and site. The top half of the figure shows the $t$- and $p$-statistic for the comparison of nodal local mean path length in each region of the AAL atlas; the bottom half of the figure shows the $t$- and $p$-statistic for the comparison of nodal global mean path length. The left column shows the comparisons for LR$^\text{NEG}$ versus HR$^\text{POS}$ infants; the right column for HR$^\text{NEG}$ versus HR$^\text{POS}$ infants. A positive $t$-statistic represents a relative reduction in mean path length in HR$^\text{POS}$ infants. Significant reductions in nodal local mean path length in HR$^\text{POS}$ infants are seen bilaterally in the temporal, parietal and occipital lobes, and in the left medial orbital gyrus. Significant reductions in nodal local mean path length in HR$^\text{POS}$ infants compared with HR$^\text{NEG}$ infants are seen in bilateral temporal, parietal and occipital lobes, the right precuneus, paracentral lobule and gyrus rectus, and the left pars triangularis and medial orbital gyrus. No regions show significantly increased nodal local mean path length in HR$^\text{POS}$ infants relative to either LR$^\text{NEG}$ or HR$^\text{NEG}$ infants. Significant increases in nodal global mean path length in HR$^\text{POS}$ infants compared with LR$^\text{NEG}$ infants are seen in the right parietal lobe, the right pars triangularis and the left medial orbital gyrus. Significant increases in nodal global mean path length in HR$^\text{POS}$ infants compared with HR$^\text{NEG}$ infants are seen bilaterally in the parietal lobe, and in the right temporal lobe and left medial orbital gyrus. No regions show significantly decreased nodal global mean path length in HR$^\text{POS}$ infants relative to either LR$^\text{NEG}$ or HR$^\text{NEG}$ infants. Note that, as with efficiency, in all cases the group differences are predominately posterior. AAL, automatic anatomical labelling; HR$^\text{NEG}$, high-risk infants with ADOS scores below the clinical cutoff for ASD; HR$^\text{POS}$, high-risk infants with ADOS scores above the clinical cutoff for ASD; LR$^\text{NEG}$, low-risk infants with ADOS scores below the clinical cutoff for ASD.
Figure 5. The comparison of the mean number of edges in the paths comprising the measures of nodal local and nodal global efficiency for LR\textsuperscript{NEG} versus HR\textsuperscript{POS} infants, and for HR\textsuperscript{NEG} versus HR\textsuperscript{POS} infants, in a model with all three groups, controlling for age, sex and site. The top half of the figure shows the $t$- and $p$-statistic for the comparison of nodal local mean number of edges in each region of the AAL atlas; the bottom half of the figure shows the $t$- and $p$-statistic for the comparison of nodal global mean number of edges. The left column shows the comparisons for LR\textsuperscript{NEG} versus HR\textsuperscript{POS} infants; the right column for HR\textsuperscript{NEG} versus HR\textsuperscript{POS} infants. A positive $t$-statistic represents a relative reduction in mean number of edges in HR\textsuperscript{POS} infants compared with LR\textsuperscript{NEG} infants. Significant reductions in nodal local mean number of edges in HR\textsuperscript{POS} compared with LR\textsuperscript{NEG} infants are seen bilaterally in the temporal, parietal and occipital lobes, and in the left medial orbital gyrus. Significant reductions in nodal local mean number of edges in HR\textsuperscript{POS} compared with HR\textsuperscript{NEG} infants are seen in bilateral temporal and parietal lobes, the right precuneus, paracentral lobule, and gyrus rectus, and the left cuneus, pars triangularis and medial orbital gyrus. No regions show significantly increased nodal local mean number of edges in HR\textsuperscript{POS} compared with HR\textsuperscript{NEG} infants. Significant increases in nodal global mean number of edges in HR\textsuperscript{POS} infants compared with LR\textsuperscript{NEG} infants are seen in bilateral temporal, parietal and occipital lobes, the right superior frontal gyrus, and the left medial orbital gyrus. Significant increases in nodal global mean number of edges in HR\textsuperscript{POS} infants compared with HR\textsuperscript{NEG} infants are seen in bilateral temporal, parietal and occipital lobes, the left superior frontal gyrus, medial orbital gyrus, gyrus rectus and pars opercularis, and the right pars triangularis, paracentral lobule and cingulate cortex. No regions show significantly decreased nodal global mean number of edges in HR\textsuperscript{POS} infants relative to either LR\textsuperscript{NEG} or HR\textsuperscript{NEG} infants. Note that, as with efficiency, in all cases the group differences are predominately posterior. HR\textsuperscript{NEG}, high-risk infants with ADOS scores below the clinical cutoff for ASD; HR\textsuperscript{POS}, high-risk infants with ADOS scores above the clinical cutoff for ASD; LR\textsuperscript{NEG}, low-risk infants with ADOS scores below the clinical cutoff for ASD.
nodal global connection strength in the right temporal and occipital lobes (Figure 3, bottom right). There were no significant increases in either nodal local or nodal global connection strength in the HRPOS infants relative to the HRNEG infants. Note that, as with efficiency, in all cases the reductions are predominately posterior.

The reductions in nodal local and nodal global efficiency in HRPOS infants appear to be not only the result of weaker connections, but also abnormalities in network structure. Relative to LRNEG infants, the HRPOS infants showed significantly reduced nodal local mean path length in bilateral temporal, parietal and occipital lobes, and the left medial orbital gyrus (Figure 4, top left). The fact that this group difference is almost identical to the group difference in nodal local mean number of edges (Figure 4, top left) indicates that these networks have reduced spatial extent. Relative to LRNEG infants, the HRPOS infants showed significantly increased nodal global mean path length in the right parietal lobe, the right pars triangularis, and the left medial orbital gyrus (Figure 4, bottom left). The comparison of nodal global mean number of edges also shows increases in HRPOS infants in posterior regions, and more bilaterally in this case, as well as in the right lateral orbital and superior frontal gyri, and the left medial orbital gyrus (Figure 4, bottom left). Thus, relative to LRNEG infants, HRPOS infants have on average longer, less direct paths connecting a node with other brain regions. Relative to HRNEG infants, the HRPOS infants showed significantly reduced nodal local mean path length in bilateral temporal and parietal lobes, the left occipital lobe, pars triangularis and medial orbital gyrus, and the right precuneus, paracentral lobule, and gyrus rectus (Figure 4, top right). Again, this group difference is exactly mirrored by the group difference in nodal local mean number of edges (Figure 5, top right) indicating that nodal local networks have reduced spatial extent in HRPOS infants. Relative to HRNEG infants, the HRPOS infants showed significantly increased nodal global mean path length in bilateral parietal lobes, the right temporal lobe and pars triangularis, and the left medial orbital gyrus (Figure 4, bottom right). The comparison of nodal global mean number of edges shows increases in HRPOS infants in bilateral temporal, parietal and occipital lobes, as well as in the left medial orbital and superior frontal gyri, the left medial orbital gyrus and pars orbitalis, and the right pars triangularis, paracentral lobule and cingulate gyrus (Figure 5, bottom right). Thus, relative to HRNEG infants, HRPOS infants have on average longer, less direct paths connecting a node with other brain regions.

**DISCUSSION**

Network efficiency measures from diffusion-based probabilistic tractography were significantly reduced, predominately in posterior brain regions, in HRPOS infants relative to both LRNEG and HRNEG infants at 24 months of age, and were inversely related to ASD symptom severity. HRPOS infants showed reduced nodal global efficiency, relative to both LRNEG and HRNEG infants, over the right superior temporal, angular and supramarginal gyri, and the left paracentral lobule, corpus callosum, and lingual gyrus. HRPOS infants showed reduced nodal local efficiency, relative to both LRNEG and HRNEG infants, over bilateral inferior temporal gyri, the left fusiform, parahippocampal and lingual gyrus, temporal pole, and medial occipital cortex. Relative to the HRNEG infants, the HRPOS infants showed more bilateral reductions in nodal global efficiency and more left lateralized reductions in nodal local efficiency. In either case, the abnormalities were concentrated in posterior regions; the frontal lobes showed no differences, with the exception of the anterior portion of Broca’s area. The significant differences in the HRPOS infants were all reductions in efficiency. Further, both nodal local and nodal global efficiency showed an inverse relation to ASD symptoms, as indexed by ADOS-based calibrated severity scores within the high-risk infants, over left temporal and occipital regions, and in Broca’s area for nodal global efficiency. The regions implicated are involved in processing auditory and visual stimuli, language and nonlinguistic social stimuli, for example, faces, and in memory, all of which show abnormalities in ASD.58–65

These results align with a number of findings in infant siblings of children with autism, and older children with ASD. Infant siblings of children with autism have been shown to have atypicalities in both visual and auditory processing,66–68 as well as an inability to integrate audio-visual information;77 in addition, connections between the bilateral occipital lobes have been implicated in relation to visual orienting in infants who go on to develop symptoms of ASD.69 Studies in toddlers with symptoms of ASD have shown reduced correlation in functional data between bilateral superior temporal gyri,71 and reduced activation in response to speech stimuli in the left superior temporal gyrus.72 Slightly older children with ASD show reduced correlation in functional data between bilateral inferior frontal gyri.73 Hypoperfusion in bilateral temporal and temporal-parietal regions has also been reported in older children with ASD,73–76 and a negative relation between cerebral blood flow and autism severity in the left superior temporal gyrus.77 Structural abnormalities in older children with autism include localized gray matter reductions within ventral and superior temporal cortical areas, as well as parietal areas.78

Frontal cortico-cortical connectivity abnormalities, which are well documented in adolescents and adults with ASD,7,79,80 presumably emerge over development together with the behavioral abnormalities with which they are associated. In children, the frontal lobes show few network abnormalities, whereas posterior regions show widespread abnormalities both locally and globally.9,35

The reductions in nodal local and nodal global efficiency shown here for the HRPOS infants correspond to a number of organizational differences. The paths underlying these reductions in efficiency were shown to be comprised of generally weaker connections, that is, fewer, smaller or less well-myelinated fibers. In the case of nodal global efficiency the paths involved were also shown to be on average physically longer and more indirect, that is, to pass through more intermediary nodes. In the case of nodal local efficiency the paths involved were comprised of weaker, but generally physically shorter and more direct paths in the HRPOS infants. But as the degree to which a node is a neighbor of another is determined by the strength of the direct connection between them, and the nodal global connection strength is decreased in the HRPOS infants, the physically shorter and more direct paths in the HRPOS infants represents reduced spatial extent of the nodal local networks. Thus, the reduction in nodal local efficiency in the HRPOS infants also reflects a reduction in long-distance connectivity. The reductions in nodal local and nodal global efficiency in the HRPOS infants are thus not simply indices of under-connectivity, but also reflect a more random configuration with more limited local connectivity and less direct connections to other brain regions.

Networks with a high degree of spatially local connectivity, but with few long-range connections, that is, shortcuts, have high local efficiency and low global efficiency; networks with a high degree of long-range connectivity, but which lack spatially local clustering, have high global efficiency and low local efficiency. Biological systems in general, and neural networks in particular, reliably balance global efficiency with local efficiency, having strong local clustering mixed with sufficient long-range connectivity to allow rapid communication between distant nodes; these have been dubbed ‘small-world’ properties.4,36,37 The results here suggest that cortico-cortical organization is deficient in infants with significant symptoms of ASD in terms of both the long-range connectivity that provides for rapid integration of information between different brain regions, and the spatial clustering that provides for efficient processing for
subtasks. Notably, these abnormalities are predominately posterior, and in regions involved in early processing of auditory and visual stimuli, audio-visual integration, language and nonlinguistic social stimuli.

The economical ‘small-world’ organization of the typical human brain is refined over development. The results here might relate to brain overgrowth and emerging symptoms. The developmental nature of the network inefficient economical organization. Further research is needed to determine the key to understanding development is development itself. Longitudinal data spanning infancy and toddlerhood will be essential in addressing these questions, as will four-dimensional multimodal methods to accurately estimate and analyze changes in connection lengths, connection strengths, network organization and behavior.

CONFLICT OF INTEREST
The authors declare no conflict of interest.

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Network inefficiencies in ASD at 24 months
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