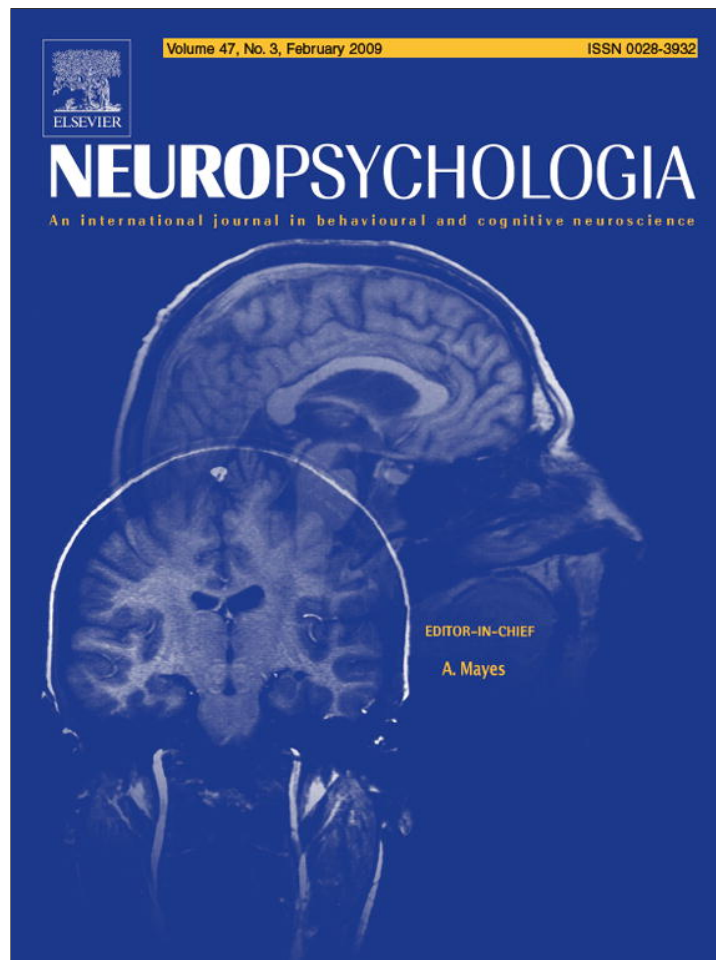


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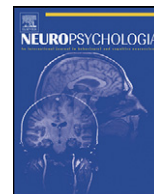
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Hemodynamic changes in the infant cortex during the processing of featural and spatiotemporal information

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ABSTRACT

Over the last 20 years neuroscientists have learned a great deal about the ventral and dorsal object processing pathways in the adult brain, yet little is known about the functional development of these pathways. The present research assessed the extent to which different patterns of neural activation, as measured by changes in blood volume and oxygenation, are observed in infant visual and temporal cortex in response to events that involve processing of featural differences or spatiotemporal discontinuities. Infants aged 6.5 months were tested. Increased neural activation was observed in visual cortex in response to a featural-difference and a spatiotemporal-discontinuity event. In addition, increased neural activation was observed in temporal cortex in response to the featural-difference but not the spatiotemporal-discontinuity event. The outcome of this experiment reveals early functional specialization of temporal cortex and lays the foundation for future investigation of the maturation of object processing pathways in humans.

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1. Introduction

Over the last 20 years a great deal of research has been conducted on the neural basis of object processing. Early studies conducted with non-human primates suggested that there are two main routes for visual object processing (De Yoe & Van Essen, 1988; Goodale & Milner, 1992; Livingstone & Hubel, 1988; Mishkin, Ungerleider, & Macko, 1983; Ungerleider & Mishkin, 1982). The *ventral* route originates from the parvocellular layers of the lateral geniculate nucleus (LGN) and projects from the primary visual cortex to the temporal cortex and mediates processing of visual features important for the recognition and identification of objects. The *dorsal* route originates from the magnocellular layers of the LGN and projects from the primary visual cortex to the parietal cortex and is important for the analysis of motion, depth, and location. More recent studies with non-human (Orban, Van Essen, & Fanduffel, 2004; Tootell, Tsao, & Vanduffel, 2003; Tsunoda, Yamane, Nishizaki, & Tanifuji, 2001; Wang, Tanifuji, & Tanaka, 1998; Wang, Tanaka, & Tanifuji, 1996) and human (Bly & Kosslyn, 1997; Grill-Spector, Kourtzi, & Kanwisher, 2001; Grill-Spector et al., 1998; Haxby et al., 1991; Kourtzi & Kanwisher, 2001; Kraut, Hart, Soher, & Gordon, 1997) primates, using more sophisticated neuroimaging techniques, provide

converging evidence for the functional distinction between these two pathways.

Although we now have extensive information about the neural correlates of object processing in the adult, little is known about the functional development of these pathways. Research conducted with infant monkeys suggests that the temporal cortex undergoes significant structural and neurophysiological development early in life (Bachevalier, Brickson, Hagger, & Mishkin, 1990; Rodman, Skelly, & Gross, 1991; Webster, Ungerleider, & Bachevalier, 1991, 1995). Metabolic, neurophysiological, and neuroanatomical data obtained with human infants also reveals significant neural maturation during the first year (e.g., Braddick, Atkinson, & Wattam-Bell, 2003; Braddick & Atkinson, 2007; Chugani & Phelps, 1986; Conel, 1939–1967; De Haan & Nelson, 1999; Franceschini, Thaker, Themelis, Krishnamoorthy & Bortfeld, 2007; Gunn et al., 2002; Purpura, 1975). However, because there are a limited number of non-invasive techniques available to measure localized functional brain activation in infants, little is known about the functional consequences of neural maturation. Recent advances in optical imaging, including near-infrared spectroscopy (NIRS), now offer the opportunity to study functional activation in human infants.

In NIRS, near-infrared light is projected through the scalp and skull into the brain and the intensity of the light that is diffusely reflected is recorded. Typically, during cortical activation local concentrations of oxyhemoglobin (HbO₂) increase, whereas concentrations of deoxyhemoglobin (HbR) decrease (Hoshi & Tamura, 1993; Jaszewski et al., 2003; Obrig et al., 1996; Strangman,

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Franceschini, & Boas, 2003; Villringer & Dirnagl, 1995). From the summated change in HbO₂ and HbR, the total change in hemoglobin (HbT) can be computed. Given that changes in HbT signal changes in regional cerebral blood flow (rCBF), having a measure of HbT is an important guide in the interpretation of NIRS data. While an increase in blood volume would result in an increase in HbO₂ and HbR, an increase in blood flow results in an increase in HbO₂ and a “washout” of HbR (i.e., an increase in relative concentration of HbO₂ and a decrease in relative concentration of HbR).

Predicting and interpreting changes in HbO₂ and HbR during cortical activation is not always straightforward, however. For example, an increase in rCBF (as indicated by HbT) produces an increase in HbO₂ and a decrease in HbR. At the same time, an increase in oxygen consumption produces a decrease in HbO₂ and an increase in HbR. Furthermore, the effect of these opposing mechanisms may be different in infants than adults (Hintz et al., 2001; Meek et al., 1998; Sakatani, Chen, Lichty, Zuo, & Wang, 1999). Hence it is important to remember that changes in relative concentrations of HbO₂ and HbR are produced by changes in blood volume, rCBF, and oxygen consumption and that the relation between these can be complex.

To capitalize on changes in HbO₂ and HbR, near-infrared light between approximately 650 and 950 nm is utilized. At these wavelengths, light is differentially absorbed by oxygenated and deoxygenated blood (Gratton, Sarno, Maclin, Corballis, & Fabiani, 2000; Villringer & Chance, 1997). Measuring the light intensity modulation during stimulus presentation, and comparing it to the light intensity during a baseline event in which no stimulus is presented, provides important information about the hemodynamic response to brain activation.

Recently, researchers have successfully applied NIRS technology to human infants in the experimental setting (e.g., Baird et al., 2002; Bortfeld, Wruck, & Boas, 2007; Pena et al., 2003; Taga, Asakawa, Maki, Konishi, & Koizumi, 2003; Wilcox, Bortfeld, Woods, Wruck, & Boas, 2005, 2008). Most of these studies have focused on region specific hemodynamic changes in the neocortex during perceptual and cognitive tasks. For example, Wilcox et al. (2005) assessed hemodynamic changes in the visual and the temporal cortex during a visual object processing task. In this task, 6.5-month-olds saw an event in which a green ball and a red box emerged successively to opposite sides of screen (Fig. 1A). Behavioral studies (Wilcox & Baillargeon, 1998a,b; Wilcox & Chapa, 2002) indicate that 4.5–11.5-month-old infants use the featural differences to interpret the event as involving two distinct objects. Analysis of the NIRS data revealed a significant increase in HbO₂ in visual and temporal cortex during the test event. Follow-up studies replicated and extended these findings to other events involving featurally distinct objects (Wilcox et al., 2008) and demonstrated that activation is observed in visual but not temporal cortex in response to control events (e.g., when the same object is seen to both sides of the screen). These data suggest that object processing is functionally localized: whereas visual cortex responds to all events involving visual objects temporal cortex responds only when the objects differ in their featural properties.

What these findings leave open to speculation, however, is the extent to which temporal cortex mediates the processing of other types of object information. For example, in adults the ventral pathway mediates processing of object features but does not typically mediate processing of the spatiotemporal properties of objects. If the ventral pathway in the infant is organized in a way similar to

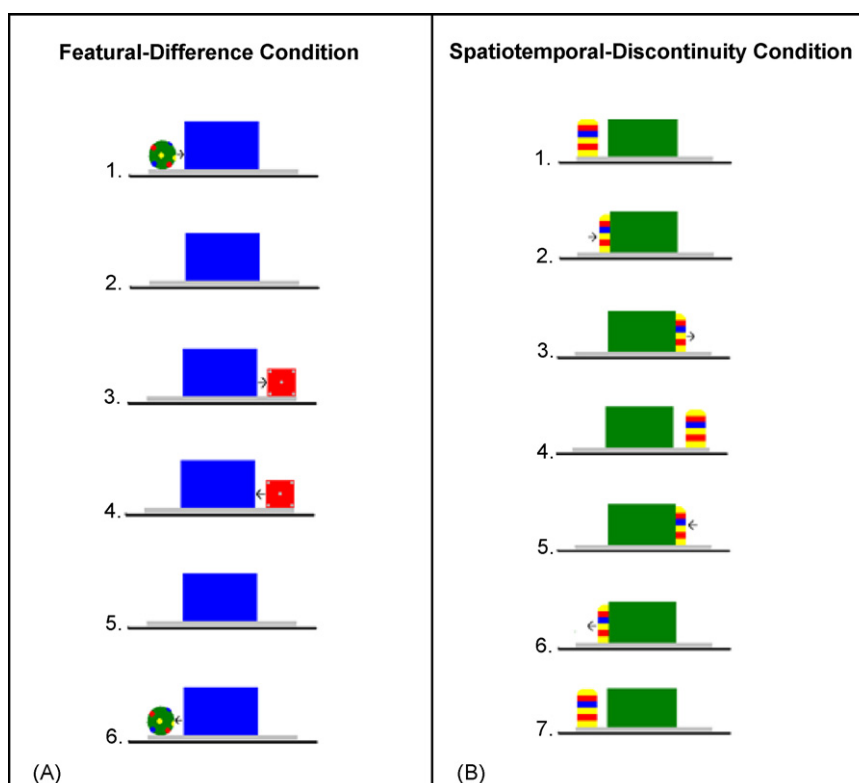


Fig. 1. The test events seen in the featural-difference (A) and spatiotemporal-discontinuity (B) condition. Although not pictured, a hand moved the objects. In the *featural-difference* condition, the ball moved right until it was fully hidden behind the occluder (2 s); the box then emerged and moved to the right edge of the platform (2 s). The box paused (1 s) and the 5 s sequence was seen in reverse. The entire 10 s ball-box cycle then repeated twice to conclude the 30 s trial. When in motion the objects moved at a rate of 12 cm/s and the occlusion interval was 1.8 s. In the *spatiotemporal-discontinuity* condition, the column moved right until it was fully occluded (2.5 s) and then the second column appeared immediately at the right edge of the occluder (the event was produced by two experimenters who had similar sized hands covered by identical white gloves) and moved right until it reached the right end of the platform (2.5 s). The column paused (1 s) and the 6 s sequence was reversed. The entire event was then repeated 1.5 times to conclude the 30 s trial. When visible, the object moved at a rate of 3 cm/s.

that of the adult, then a different pattern of activation should be observed in temporal cortex in response to events involving analysis of object features than to events involving analysis of spatiotemporal information. The present research tests this hypothesis. Infants aged 6.5 months were presented with the featural-difference event of Wilcox et al. (2005; Fig. 1A) and a spatiotemporal-discontinuity event (Fig. 1B) and NIRS data were collected. Behavioral studies have demonstrated that infants 3.5 months and older interpret the spatiotemporal-discontinuity event as involving two objects (Schweinle & Wilcox, 2004; Wilcox & Schweinle, 2003).

2. Materials and methods

2.1. Participants

Twelve 6.5-month-olds, 8 M (M age = 6 months, 15 days, range = 5 months, 12 days to 7 months, 11 days). Twelve additional infants were tested but eliminated from analysis because they failed to contribute usable NIRS data (e.g., large motion artifacts and/or poor signal-to-noise ratio). Seven infants saw the featural-difference event first.

2.2. Apparatus, stimuli, and procedure

Infants sat on a parent's lap facing a puppet-stage apparatus. The green ball used in the featural-difference event was 10.25 cm in diameter with colored dots.

The red box was 10.25 cm square and decorated with silver thumbtacks. The occluder was 21.5 cm × 30 cm and made of blue cardboard. The columns used in the spatiotemporal-discontinuity event were 12 cm × 6 cm × 3 cm and made of colored Duplos. The occluder was 24 cm × 35 cm and dark green. The objects were moved by a gloved hand, which entered the apparatus through a slit in the back wall. The test events are depicted in Fig. 1.

Prior to each 30 s test trial infants were presented with a 10 s baseline (silent pause) in which a muslin-covered shade covered the front opening of the apparatus and hid the stage. The shade was raised at the beginning of each test trial and lowered at the end of each trial. Cloth-covered barriers isolated the infant from the testing room. The stage was illuminated with 20-Watt fluorescent bulbs affixed to each inside wall of the apparatus. No other lighting was used.

The time infants spent looking during each test trial was recorded by two trained observers and looking time data were time-locked to the NIRS data. Inter-observer agreement averaged 96%. A blocked design (four trials of one event followed by four trials of the other) was used because the events were produced live in the puppet-stage apparatus and each event involved different display characteristics (e.g., objects and screen). Frequent changing of display characteristics was difficult for the experimenters and distracting to the infants.

2.3. Instrumentation

The instrumentation was identical to that of Wilcox et al. (2005). Briefly, the imaging equipment contained three major components: (1) two fiber optic cables (1 mm in diameter) that delivered near-infrared light to the scalp of the participant; (2) four fiber optic cables (2.5 mm in diameter) that detected the diffusely reflected light at the scalp; and (3) an electronic control box. The electronic control box pro-

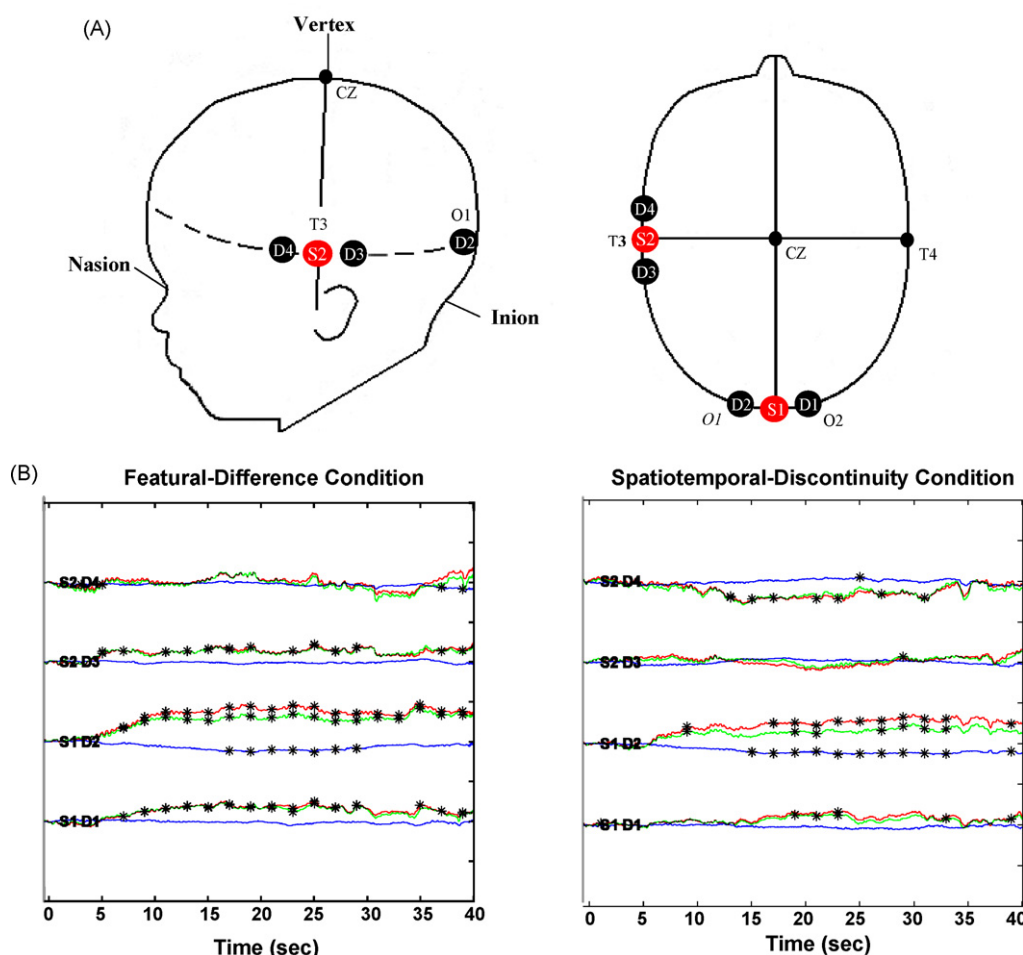


Fig. 2. Location of probe placement (A) and hemodynamic response curves (B) for each detector. Probes were placed on the skull using the International 10–20 system. One probe was positioned so that the source (S1) lay directly above the inion and the detectors (D1 and D2) lay over the right and left visual cortex, respectively. The other source (S2) was positioned at T3 and the corresponding detectors (D3 and D4) lay anterior and posterior to T3 over temporal cortex. Emitter-detector distances were 2 cm. The hemoglobin response curves, in optical density units presented as $\text{mM} \times \text{cm}$ (x-axis), were averaged across participants and trials for the four detectors. For purposes of interpretation, HbT is displayed along with HbO₂ and HbR (green, red, and blue lines, respectively). The left panel of (B) displays responses obtained during the featural-discontinuity event and the right panel displays responses during the spatiotemporal-discontinuity event. The event began at time 0 and continued for 30 s; 31–40 is post-stimulus (silent pause). The asterisks indicate points along the response curves that differed significantly from 0 (baseline).

duced light at 690 and 830 nm wavelengths with two laser-emitting diodes (Boas, Franceschini, Dunn, & Strangman, 2002; TechEn Inc.). Each emitter delivered both wavelengths and each detector responded to both wavelengths. The signals received by the electronic control box were processed and relayed to a DELL Inspiron 7000™ laptop computer.

Prior to test, infants were fitted with custom-made headgear that secured the fiber optics to the scalp. The ends of the fiber optic cables were arranged in two triads, each containing one emitter and two detectors. Triad placement is shown in Fig. 2A.

2.4. Processing of the NIRS data

The NIRS data were processed, for each detector separately, using a procedure similar to that of Wilcox et al. (2005). The raw signals were acquired at the rate of 200 samples per second, digitally low-pass-filtered at 10.0 Hz, a principal components analysis was used to design a filter for systemic physiology and motion artifacts, and the data were converted to relative concentrations of oxygenated (HbO₂) and deoxygenated (HbR) blood using the modified Beer–Lambert law (Strangman, Boas, & Sutton, 2002). Changes in HbO₂ and HbR were examined using 43 s time epochs: the 3 s prior to the onset of the test event, the 30 s test event, and the 10 s following the test event. The mean optical signal from –3 to 0 s (baseline) was subtracted from the signals and other segments of the 43 s time epoch were interpreted relative to this zeroed baseline. Optical signals were averaged across trials and then infants for each condition. Trials objectively categorized as containing motion artifacts (a change in the filtered intensity greater than 5% in 1/20 s during the 3 s baseline and 30 s test event) were eliminated from the mean. A total of 11 trials (of the 96 remaining) were eliminated. Two additional trials were eliminated because the infant failed to watch the event or procedural error.

3. Results

3.1. Looking time data

The infants looked almost continuously throughout the test trials (featural-difference, *M* = 27.62, *S.D.* = 1.85 and speed-discontinuity, *M* = 26.74, *S.D.* = 1.46) suggesting that they found the test events engaging.

3.2. NIRS data

The hemoglobin concentration response curves are shown in Fig. 2. In the featural-difference condition, relative changes in

HbO₂ and HbR concentration from 5 to 30 s following initiation of the event were compared to baseline. The first emergence of the box occurred at 3 s and, allowing 2 s for the hemodynamic response to become initiated, changes in HbO₂ and HbR should be detectable by 5 s. In the spatiotemporal-discontinuity condition, relative changes in HbO₂ and HbR concentration were assessed starting at 6 s (the first immediate emergence occurred at 4 s).

Three sets of analyses were conducted. First, mean HbO₂ and HbR responses for each condition and detector were compared to 0 (Table 1). In visual cortex, a significant increase in HbO₂ was obtained in response to the featural-difference event (at D1 and D2) and the spatiotemporal-discontinuity event (at D2). A significant decrease in HbR was observed at D2 in response to the spatiotemporal-discontinuity event. In temporal cortex, a significant increase in HbO₂ was observed in response to the featural-difference event at D3 but not D4. In contrast, a decrease in HbO₂, which approached significance (*P* = .058), was observed in response to the spatiotemporal-discontinuity event at D4 but not D3.

Second, to test the extent to which the responses observed at each detector varied by the event seen, paired sample *t*-tests were conducted for each detector with condition as the within-subjects factor (Table 1). At D1 and D2 the mean HbO₂ and HbR responses did not vary significantly by condition. However, at D3 the HbO₂ response observed in the featural-difference condition (which was positive in direction) differed from that observed in the spatiotemporal-discontinuity condition. Although the *t*-test was not statistically significant (*P* = .095), the effect size was large, Cohen's *d* = .77 (see Cohen, 1988 for evaluating effect sizes), indicating that the two groups differed in their responses. At D4, the HbO₂ response observed in the spatiotemporal-discontinuity condition (which was negative in direction) differed reliably from that observed in the featural-difference condition, and the effect size was also large, Cohen's *d* = .77.

Finally, to assess the extent to which the response observed at one detector within a neural region differed reliably from that observed at the other detector within that same region, paired-sample *t*-tests were conducted for each neural region and condition

Table 1
Relative change in HbO₂ and HbR concentration displayed by detector and condition.

Neural Region	Detector	Chromophore	Condition		Paired <i>t</i> -tests
			Featural-difference <i>M</i> (<i>S.D.</i>)	Spatiotemporal-discontinuity <i>M</i> (<i>S.D.</i>)	
Visual	D1	HbO ₂	0.0066 (.006)** <i>t</i> = 3.96, d.f. = 11	0.0035 (.011) <i>t</i> = 1.10, d.f. = 11	<i>t</i> = –1.01, d.f. = 11
		HbR	–0.0003 (.003) <i>t</i> = –0.35, d.f. = 11	–0.0009 (.005) <i>t</i> = –0.61, d.f. = 11	<i>t</i> = –0.40, d.f. = 11
	D2	HbO ₂	0.0143 (.018)** <i>t</i> = 2.79, d.f. = 11	0.0092 (.014)* <i>t</i> = 2.21, d.f. = 11	<i>t</i> = –1.11, d.f. = 11
		HbR	–0.0035 (.006) <i>t</i> = –2.03, d.f. = 11	–0.0038 (.006)* <i>t</i> = –2.23, d.f. = 11	<i>t</i> = –0.26, d.f. = 11
Temporal	D3	HbO ₂	0.0058 (.009)** <i>t</i> = 2.34, d.f. = 11	–0.0001 (.006) <i>t</i> = –0.08, d.f. = 10	<i>t</i> = –1.84, d.f. = 10+
		HbR	–0.0004 (.003) <i>t</i> = –0.35, d.f. = 11	0.0007 (.004) <i>t</i> = 0.51, d.f. = 10	<i>t</i> = 0.74, d.f. = 10
	D4	HbO ₂	0.0010 (.009) <i>t</i> = 0.39, d.f. = 10	–0.0063 (.010) <i>t</i> = –2.14, d.f. = 10	<i>t</i> = –2.47, d.f. = 10*
		HbR	–0.0008 (.003) <i>t</i> = –0.94, d.f. = 10	0.0001 (.003) <i>t</i> = 0.18, d.f. = 10	<i>t</i> = –0.69, d.f. = 10

Note: Cells contain mean relative optical density units in mM x cm averaged from 5 to 30 s (featural-difference condition) and 6 to 30 s (spatiotemporal-discontinuity condition). Mean responses were compared to 0 using *t*-tests (two-tailed) and the asterisks indicate those responses that differed significantly from 0 (**P* < .05 and ***P* < .025). The last column contains paired *t*-tests (two-tailed) comparing responses across condition (**P* < .05 and +*P* < .1 with a large effect size). Effect sizes are reported in the text.

Table 2
Comparison of hemodynamic responses across detectors within each neural region.

Condition	Chromophore	Visual cortex D1 vs. D2	Temporal cortex D3 vs. D4
Featural-difference	HbO ₂	<i>t</i> = –1.41, d.f. = 11	<i>t</i> = 2.32, d.f. = 10*
	HbR	<i>t</i> = 1.45, d.f. = 11	<i>t</i> = 0.28, d.f. = 10
Spatiotemporal-discontinuity	HbO ₂	<i>t</i> = –1.30, d.f. = 11	<i>t</i> = 2.19, d.f. = 9+
	HbR	<i>t</i> = 1.69, d.f. = 11	<i>t</i> = 0.55, d.f. = 9

Note: Paired *t*-tests compared hemodynamic responses, for each condition and chromophore, across detectors within each neural region. The symbols indicate the comparison in which the responses observed at the two detectors differed reliably from each other using two-tailed statistics (**P* < .05 and +*P* < .1 with a large effect size). Effect sizes are reported in the text.

with detector as the within-subjects factor (Table 2). In visual cortex, hemodynamic responses did not vary significantly by detector, for infants in either condition. In temporal cortex, some differences emerged. In the featural-difference condition, the HbO₂ response observed at D3 differed significantly from that observed at D4, Cohen's $d = .53$. In the spatiotemporal-discontinuity condition, the HbO₂ response observed at D3 and D4 also differed. Although the t -test was not statistically significant ($P = .056$) the effect size was large, Cohen's $d = .75$.

4. Discussion

The present research used near-infrared spectroscopy to assess neural activation, as measured by changes in blood volume and oxygenation, in visual and temporal cortex in 6.5-month-olds during two object processing tasks, one that involved analysis of object features and the other that required analysis of spatiotemporal information.

As predicted, a hemodynamic response was observed in visual cortex (D1 and D2) in both conditions and the responses did not vary significantly by condition. At the same time, the hemodynamic response obtained at D1 differed qualitatively from that obtained at D2. For example, a significant increase in HbO₂ was observed at D2 in response to both events and a significant decrease in HbR was observed at D2 in response to the featural-difference event. In comparison, a less robust response was observed at D1: a significant increase in HbO₂ was observed in response to the featural-difference event only and no significant decreases in HbR were observed. Why were qualitatively different hemodynamic response patterns observed at D1 and D2? Recall that in the present study the light source was placed directly above theinion, so that D1 lay over the left visual cortex whereas D2 lay over the right visual cortex. One possible explanation is that in the infant, the left and right visual cortices are organized slightly differently, so that structurally analogous areas in the two hemispheres respond differently to the same visual stimuli. Different functional responses could lead to different hemodynamic responses. Alternatively, it is possible that the two hemispheres are not structurally identical, so that measuring from skull locations equidistant from the midline, but in different hemispheres, does not guarantee measurements from structurally (let alone functionally) analogous neural areas. A final possibility is that these results reflect an asymmetry in the degree of neurovascular regulation in visual areas of the two hemispheres. That is, in right visual cortex changes in rCBF may not be as well matched to energy demands (oxygen consumption) as in left visual cortex. Additional research will be needed to (a) establish the extent to which reliable hemispheric differences in hemodynamic responses to visual stimuli exist in V and other areas and (b) identify the basis of these differences. In the meantime, the patterns observed in the present experiment are best interpreted with caution.

As predicted, the hemodynamic response observed in temporal cortex differed by condition, although the way in which this was manifested was unexpected. A significant *increase* in HbO₂ was observed at D3 in the featural-difference condition and a *decrease* in HbO₂ was observed at D4 in the spatiotemporal-discontinuity condition. In addition, the responses observed at each detector differed by condition, and within condition the responses differed by detector. These dissociations suggest that the cortical regions directly anterior to and posterior to T3 are functionally distinct.

On the basis of neuroanatomical data obtained with human infants (Conel, 1939–1967; Purpura, 1975) and on data demonstrating the relation between 10 and 20 skull coordinates and underlying neural structure in adults (Okamoto et al., 2004), we suspect that the area posterior to T3 lies near the temporal–occipital border and may include part of the lateral occipital cortex (LOC). Neuroimag-

ing data obtained with adults (Grill-Spector et al., 1998, 1999, 2001; Haxby et al., 1991; Kourtzi & Kanwisher, 2001; Kraut et al., 1997; Malach et al., 1995) indicate that the LOC is involved in the processing of object features, but does not respond to the spatiotemporal properties of objects. In contrast, we suspect that the area anterior to T3 lies close to the medial or superior temporal gyrus. This area does not appear to mediate processing of featural differences or spatiotemporal discontinuities, at least in the infant.

Unexpectedly, a *decrease* in HbO₂ was observed at D4 in the spatiotemporal-discontinuity condition, and this response differed significantly from the response obtained in the featural-difference condition. In addition, there was a decrease in HbT indicating a decrease in rCBF. One possible explanation for this unusual pattern of results is that viewing the spatiotemporal-discontinuity event led to neural deactivation and a corresponding decrease in rCBF and blood volume. Either of these could have produced an increase in the local concentration of HbR relative to HbO₂. Another, more likely explanation, assumes no activation at D4. The decrease in HbO₂ reflects, instead, the blood supply being diverted to nearby areas that are active. These areas could lie either deeper in the cortex or adjacent on the surface (e.g., areas associated with the dorsal system). Regardless of how to best conceptualize the hemodynamic response obtained at D4 during the spatiotemporal-discontinuity event, it is important to remember that this response differed from that observed at D3 during the spatiotemporal-discontinuity event, suggesting that the neural regions from which these two detectors were measuring differ in their functional response to the event and/or in the nature of their neurovascular regulatory mechanisms.

The data also suggest that HbO₂ is a more robust measure of the hemodynamic response than HbR. A significant increase in HbO₂ was observed at a number of detectors, in both V and T, and in response to both the featural-difference and spatiotemporal-discontinuity events. In contrast, the only detector at which we saw a significant decrease in HbR and a corresponding increase in HbO₂ was at D2 in V (spatiotemporal-discontinuity condition). These results are consistent with those of other investigators who have reported that, generally speaking, HbO₂ is a more robust and reliable measure of neural activation than HbR (Bartocci et al., 2000; Chen et al., 2002; Hoshi & Tamura, 1993; Jaszewski et al., 2003; Kato, Kamei, Takashima, & Ozaki, 1993; Sakatani et al., 1999; Strangman et al., 2002, 2003).

Finally, the hemodynamic responses observed in temporal cortex appeared smaller in magnitude and less robust than those observed in visual cortex. For example, the magnitude of the response observed at D3 in response to the featural-difference event was less than that observed at D2. Although we can only speculate, the less robust responses observed at D3 and D4 may reflect greater immaturity of the temporal as compared to visual cortex. This would be consistent with neuroanatomical and metabolic data (Chugani & Phelps, 1986; Conel, 1939–1967; Franceschini et al., 2007; Purpura, 1975) and might reflect less efficient neural processing and/or energy demands associated with maturational events.

In summary, the present results demonstrate that there are region specific differences in visual object processing in human infants and that NIRS is sufficiently sensitive to detect these differences. The ability to study functional brain activation in awake, processing infants represents a significant advancement in the field of developmental neuroscience and will allow investigators to study localized functional maturation of the human brain.

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