Synchronous activity of two people's prefrontal cortices during a cooperative task measured by simultaneous near-infrared spectroscopy

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Abstract. The brain activity during cooperation as a form of social process is studied. We investigate the relationship between coinstantaneous brain-activation signals of multiple participants and their cooperative-task performance. A wearable near-infrared spectroscopy (NIRS) system is used for simultaneously measuring the brain activities of two participants. Each pair of participants perform a cooperative task, and their relative changes in cerebral blood are measured with the NIRS system. As for the task, the participants are told to count 10 s in their mind after an auditory cue and press a button. They are also told to adjust the timing of their button presses to make them as synchronized as possible. Certain information, namely, the “intertime interval” between the two button presses of each participant pair and which of the participants was the faster, is fed back to the participants by a beep sound after each trial. When the spatiotemporal covariance between the activation patterns of the prefrontal cortices of each participant is higher, the intertime interval between their button-press times was shorter. This result suggests that the synchronized activation patterns of the two participants’ brains are associated with their performance when they interact in a cooperative task. © 2011 Society of Photo-Optical Instrumentation Engineers (SPIE). [DOI: 10.1117/1.3602853]

Keywords: near-infrared spectroscopy; wearable optical topography; brain activity; hemoglobin; simultaneous measurement; hyperscanning.

Paper 10656R received Dec. 14, 2010; revised manuscript received Apr. 29, 2011; accepted for publication Jun. 2, 2011; published online Jul. 18, 2011.

1 Introduction

One of the common symptoms of autism,^{1,2} Asperger syndrome,^{1,3} schizophrenia, and depression is a lack of social skill. To alleviate these mental disorders on the basis of a neuroscience approach, it is important to investigate the relationship between social skills and brain function.⁴ Social skills, in general, include adaptability of one’s own behavior to a social context, reading the atmosphere or others’ intentions, communication skills, and ethical sense.^{5,6} Most of these social skills are acquired through interactions with others. To study the social-brain function, it is therefore meaningful to investigate interacting brains of multiple participants.

Although the “social brain”^{7–10} has been studied by many researchers, it is mostly measured in only one brain at a time by using various modalities, such as electroencephalography (EEG),^{11,12} positron-emission tomography,^{13–15} single-photon-emission computed tomography,¹⁶ functional magnetic-resonance imaging (fMRI),^{17–21} and functional near-infrared spectroscopy (fNIRS).^{22–26} Consequently, little is known about how social functions are processed in interacting brains of multiple participants.

There are various ways to study two or more brains at once. We used a wearable-optical-topography (WOT) system based on near-infrared spectroscopy (NIRS)^{27–29} because this system can simultaneously measure the brain activities of two or more people who are interacting in a natural manner, namely, dur-

ing face-to-face communication. An NIRS system measures the changes in cerebral blood volume by radiating weak visible or near-infrared light into the head and detecting the transmitted light from another position.^{30–33} On the basis of NIRS, optical topography (OT),^{34,35} which uses multiple light sources and detectors, obtains two-dimensional topographical images of the changes in cerebral blood volume. OT systems have been widely used for research and clinical purposes,^{36–40} especially for measuring the brain activity of infants and children,^{25,41–45} because they have a high level of safety^{46,47} and require few constraints. For simultaneously measuring of multiple participants’ brains (i.e., hyperscanning) and thereby investigating the relationship between brain activity and social interaction, some researchers used EEG,^{48–51} which is less expensive and has a higher temporal resolution than NIRS. The spatial resolution of EEG, however, remains lower than that of NIRS.^{52,53} Others used fMRI,^{54,55} which has a higher spatial resolution than NIRS and EEG, but it is difficult for participants to interact in a natural manner in an fMRI environment. To measure multiple participants’ brain activity during face-to-face interaction and to demonstrate the effectiveness of simultaneous NIRS measurement, we used a wearable NIRS system in this study.

It has been reported that when people interact or cooperate during common social-interaction tasks, a specific brain area activates.^{18,48} For example, in an EEG study, the medial prefrontal region was activated in conflict situations during the “prisoner’s dilemma” task (one common social-interaction task),⁴⁸ and in an fMRI study, during the “trust and reciprocity” game, the medial prefrontal region of highly cooperative

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participants was significantly more activated during human–human interactions than during human–computer interactions.¹⁸ It is therefore reasonable to suppose that the relationship between spatial brain-activity patterns of two people is effective in the investigation of brain functions related to social interactions.

In the present study, considering that the medial prefrontal region has been reported to be activated during social interaction, we focused our attention on activity in the prefrontal-cortex region during a cooperative task as a form of social process. To investigate this region, we proposed an analytical method using spatiotemporal activation patterns of multiple participants. If the relationship between the analyzed brain-activity data of two participants in a natural attitude and the “quality of cooperation” were obtained by using simultaneous NIRS measurement and data analysis of multiple participants, this methodology would become an important tool in research on social cognition. The aim of this study was to investigate the relationship between multiparticipants’ coinstantaneous brain-activation signals and cooperative-task performance by simultaneous NIRS measurement and to show the significance of the simultaneous measurement of two or more brains. A WOT system was used to investigate interactive brain activities during the cooperative task (namely, a cooperative button-pressing) because it is suitable for measuring the prefrontal cortex of multiple participants interacting in a natural manner.^{27–29}

2 Materials and Methods

2.1 Participants

Twelve healthy adults (six pairs), nine males and three females, between 25 and 50 years old (mean $\pm$ SD: 37.5 ± 8.0) participated as volunteer subjects in this study. All participants provided written informed consent after a complete explanation of the study. The experiments were approved by the Ethics Committee of Hitachi, Ltd.

2.2 Near-Infrared-Spectroscopy Measurement

A WOT system, the same one used in our previous study,²⁹ was used for measuring cerebral blood-volume change of the 12 participants. The system generally measures the changes in the product of hemoglobin (Hb) concentration (C) and effective optical path length (L), called simply “Hb change (ΔCL)” here, in human-brain tissue, such as the cerebral cortex. The unit of Hb change (in mM times millimeter) is molar concentration (in mM = mmol/l) multiplied by optical path length (in millimeters). Two kinds of Hb change, oxygenated (oxy-) and deoxygenated (deoxy-), are obtained by using two different absorption-coefficient spectra as well as absorption data for at least two wavelengths according to the modified Beer–Lambert law.^{34,56} The WOT system does not have optical fibers, which may limit the participant’s range of movement, and enables simultaneous measurement of multiple participants by using a personal computer through a wireless local area network (LAN). The details of the WOT system are described previously.^{27–29} The WOT system is small, light, and covers the entire forehead. The probe positions of the light sources and detectors and the channel positions of the WOT system are shown in Fig. 1. The channel position is defined as the center point between each light source and detector. The WOT probes were placed on each

Fig. 1 Probe positions of light sources and detectors and channel positions of wearable optical topography system. Each channel position is defined as the center point between each light source and detector.

participant’s head so that the bottom line of the probes was on the participant’s eyebrow line.

2.3 Interparticipant Variability of Channel Positions

To evaluate the variability of the measurement positions on all the participants, the probe positions were measured with a three-dimensional digitizer on the basis of the international 10–20 system,⁵⁷ namely, the standard method for electrode placement in electroencephalography, and the measurement positions on the scalp were converted to those on the brain in the Montreal Neurological Institute (MNI) coordinate system by the probabilistic-determination method.^{58–60} Figure 2 shows the measurement positions averaged among the 12 participants. For the 12 participants, Table 1 lists the means and standard deviations of the channel positions in regard to each axis. Channel 5 had the maximal square sum of the standard deviations in the three directions. The standard deviations of the channel-5 positions in regard to the three axes were 4.7 mm on the x -axis, 8.5 mm on the y -axis, and 7.0 mm on the z -axis. Because these deviations were much smaller than the 30-mm probe intervals, the difference and variability of the positions of the measurement channels for different participants could be ignored and the data of a channel for one participant was compared to those of the same channel for the other participants.

2.4 Measurement of Activation in mPFC

It has been reported that the depth of light penetration from the surface of the brain in adult humans is several centimeters,⁶¹ depending on the thickness of superficial tissue.⁶² Some researchers reported to have measured parts of relatively deep areas by noninvasive NIRS imaging. For example, NIRS imaging was used to measure the anterior-dorsal region of the medial prefrontal cortex (mPFC) regions of adults,⁶³ the prefrontal region (which is a similar region to the mPFC region in adults) of four-month-old infants,⁶⁴ and anterior orbitofrontal cortex

Fig. 2 Measurement positions on a standard brain in the MNI coordinate system.

Table 1 Mean and standard deviation (SD) of channel positions for 12 participants.

Channel	Mean $\pm$ SD		
	X (mm)	Y (mm)	Z (mm)
1	64.1 $\pm$ 3.7	10.5 $\pm$ 11.0	7.3 $\pm$ 6.7
2	61.4 $\pm$ 3.1	20.4 $\pm$ 10.5	21.1 $\pm$ 5.6
3	58.2 $\pm$ 2.5	29.4 $\pm$ 10.9	-2.3 $\pm$ 7.4
4	55.4 $\pm$ 3.5	38.4 $\pm$ 8.8	11.8 $\pm$ 6.9
5	47.4 $\pm$ 4.7	44.6 $\pm$ 8.5	26.4 $\pm$ 7.0
6	47.3 $\pm$ 4.6	54.4 $\pm$ 7.4	1.6 $\pm$ 7.6
7	38.3 $\pm$ 6.0	59.5 $\pm$ 6.2	16.5 $\pm$ 8.1
8	26.9 $\pm$ 6.7	59.7 $\pm$ 5.9	29.9 $\pm$ 7.8
9	27.4 $\pm$ 6.4	69.1 $\pm$ 3.3	5.6 $\pm$ 8.5
10	16.9 $\pm$ 5.6	69.5 $\pm$ 3.2	19.0 $\pm$ 8.4
11	4.3 $\pm$ 6.4	63.6 $\pm$ 5.0	30.4 $\pm$ 7.7
12	3.8 $\pm$ 7.4	71.6 $\pm$ 1.6	5.9 $\pm$ 8.6
13	-11.0 $\pm$ 4.6	69.4 $\pm$ 3.2	18.1 $\pm$ 8.5
14	-21.5 $\pm$ 4.2	61.6 $\pm$ 6.4	29.0 $\pm$ 7.1
15	-22.1 $\pm$ 5.0	69.9 $\pm$ 2.4	4.1 $\pm$ 7.9
16	-34.6 $\pm$ 4.2	61.4 $\pm$ 4.6	14.4 $\pm$ 7.2
17	-44.1 $\pm$ 3.3	47.2 $\pm$ 7.2	24.4 $\pm$ 5.6
18	-43.8 $\pm$ 3.8	55.7 $\pm$ 4.9	-1.0 $\pm$ 6.2
19	-52.4 $\pm$ 3.1	40.6 $\pm$ 7.1	8.5 $\pm$ 5.6
20	-58.1 $\pm$ 2.6	23.5 $\pm$ 8.7	18.4 $\pm$ 4.0
21	-54.4 $\pm$ 1.6	33.0 $\pm$ 7.0	-4.2 $\pm$ 4.2
22	-60.3 $\pm$ 2.6	13.8 $\pm$ 9.3	5.2 $\pm$ 3.5

regions of mothers and infants.²⁶ Mitchell et al.¹⁹ reported the MNI coordinates of a part of the mPFC region that is related to social cognition as ($X = 9$ mm, $Y = 54$ mm, $Z = 36$ mm). This position is near the superficial layer of the brain and close to channel 11 in the WOT system (see Table 1). The superficial part of the mPFC region is therefore assumed to be within the optical paths of an NIRS imaging system with 30-mm source-detector spacing.

2.5 Experimental Setup

Figure 3 illustrates the experimental setup used to simultaneously measure the brain activity two participants by using the WOT system. Two participants sat face-to-face across a table. Each pair of participants wore WOT probes connected to portable boxes for measurement control, signal processing, and

Fig. 3 Experimental setup. (a) Face-to-face arrangement of two participants across a table. (b) System diagram of simultaneous measurement of two participants' brain activity by using the WOT system. The two WOTs are controlled via wireless LAN by a PC, and the timing of auditory stimulation and marker input are controlled via a stimulation-presentation PC (SP-PC). The button-press time of each participant is recorded by the SP-PC.

wireless communication. The portable boxes were controlled via a wireless-LAN access point by a personal computer (PC) for a WOT controller.

The stimulation-presentation PC (SP-PC) presented auditory cues and feedback sounds to the participants. The feedback sounds were presented a few seconds after he or she pressed the button (connected to the SP-PC) with different tones for each participant. When the auditory cue to start counting is given and when the first button is pressed by one of the participants in each pair (hereafter, first "button-press time"), the SP-PC sends a marker signal to the WOT controller and records the button-press times of each pair of participants.

2.6 Cooperative Button-Press Task

Each pair of participants performed the following cooperative button-press task while the changes in their cerebral blood volume were measured by the WOT system. Figure 4 shows the time sequence of this task. At the start of the task, the two participants closed their eyes and put their index finger on a button on a PC keyboard. In each trial, they were told to press their button after counting 10 s in their mind and were also told to adjust their button-press times to make them as synchronized as possible. The auditory cue to start the time counting was given by a 1000-Hz beep sound (duration 200 ms) emitted from a speaker.

In the cooperative task, the "intertime interval" (namely, the time between the button-press timings of the two

Fig. 6 Button-press time during a cooperative task.

(including both the used and interpolated channels) were therefore analyzed. The effect of using the interpolated channels is to enhance the effective activated area of the prefrontal cortex and, consequently, make the covariance between AV maps higher even when the activated areas of two participants' brains were located at different (but nearby) channel positions.

AVs for the used and interpolated channels were calculated by using the data set composed of the baseline and activation periods shown in Fig. 4; that is, 1 s before the start of counting was taken as the baseline period, and the time period from the 6-s point after the start of counting until the first button press was taken as the activation period. The AVs of the two participants in each trial are called $AV_1(i, c)$ ($i = 1, 2, \dots, 10; c = 1, 2, \dots, 33$) and $AV_2(i, c)$ ($i = 1, 2, \dots, 10; c = 1, 2, \dots, 33$), where i indicates trial number, and c indicates the analyzed channel number including interpolated channels. Covariance $r(i)$ ($i = 1, 2, \dots, 10$) between AV_1 and AV_2 in each trial was calculated from Eqs. (2), (3a), and (3b), and $r(i)$ of each pair was normalized to zero-average and unit standard deviation among trials,

$$r(i) = \sum_{c=1}^{33} \phi_1(i, c) \phi_2(i, c), \quad (2)$$

$$\phi_1(i, c) = AV_1(i, c) - \frac{1}{33} \sum_{c=1}^{33} AV_1(i, c), \quad (3a)$$

$$\phi_2(i, c) = AV_2(i, c) - \frac{1}{33} \sum_{c=1}^{33} AV_2(i, c). \quad (3b)$$

A high covariance of the AV maps [$r(i) > 0$] means that the activated areas of the two brains are located at similar positions. On the other hand, a low covariance [$r(i) < 0$] means that the activated areas of the two brains have different patterns. The intertime intervals between the two button presses of each pair were also normalized to zero-average and unit standard deviation because the baselines of both the covariances and intertime intervals depend on each participant pair.

The ten-trial data of all the pairs were divided into two groups: one included the trials in which the normalized covariances of the AV maps were positive (referred to hereafter as “high-covariance trials”), and the other included those in which the normalized covariances were negative (referred to hereafter as “low-covariance trials”). To investigate the relationship between the normalized covariance of the brain activations of two participants and the normalized intertime intervals between the two button presses of each pair, a Wilcoxon rank-sum tests using the intertime intervals of the two groups (i.e., the high- and low-covariance trials) was conducted. We confirmed in advance that at least one of the two groups did not have a normal distribution using both coefficient of skewness and coefficient of kurtosis. The covariances and intertime intervals for the first trial in the cooperative task were not used because it was assumed that neither participant would be able to anticipate the button-press time of the other in the first trial; that is, the participants would not be able to cooperate with one another before getting feedback. Data processing and statistical analyses of the NIRS signals

Fig. 7 Mean intertime intervals between button presses among all the pairs for each trial.

were performed using Matlab Version 7.4 (The MathWorks, Inc., Natick, MA).

3 Results

3.1 Behavioral Results

Figure 6 plots the behavioral results, namely, button-press time, for each trial in the cooperative task performed in Exps. 1–6. The horizontal axis represents trial number, and the vertical axis represents button-press time. In the eighth trial of Exp. 4, a behavioral error occurs in the results for participant 8, who said after the experiment that she missed pressing the button in the instructed way by mistake. The result for the eighth trial of Exp. 4 was therefore not used in the following analysis. Figure 7 plots the mean intertime intervals between button presses among all the pairs for each trial. The error bars represent standard deviations. The intertime interval of the first trial is the longest of all the trials and greatly decreases at the second trial. From the second to tenth trials, the mean intertime intervals do not greatly change.

Table 2 lists the means and standard deviations of the button-press times of all the participants participating in the control task. In the control task, because there was no explicit interaction between any of the paired participants and no feedback in each trial, it was expected that the participants would behave in the same way in all trials. Only the means and standard deviations of the button-press times were therefore calculated.

In Exps. 1, 3, and 6, the cooperative task was performed first, after which the control task was performed. In contrast, in Exps. 2, 4, and 5, the order of the tasks was opposite. The

Table 2 Mean and standard deviation (SD) of button-press time during a control task.

Exps.	Participants	Mean $\pm$ SD(s)	Participants	Mean $\pm$ SD(s)
1	1	8.6 $\pm$ 0.4	2	11.8 $\pm$ 0.9
2	3	12.6 $\pm$ 2.5	4	12.9 $\pm$ 3.1
3	5	15.6 $\pm$ 0.7	6	16.5 $\pm$ 1.9
4	7	11.1 $\pm$ 0.4	8	10.1 $\pm$ 1.2
5	9	10.2 $\pm$ 0.6	10	12.1 $\pm$ 1.3
6	11	10.5 $\pm$ 0.3	12	9.7 $\pm$ 0.3

intertime intervals between the button presses of each pair performing the control task under both task-order conditions were not significantly different (Student's *t*-test, $p > 0.2$). It is thus considered that the behavioral results for the control task were not influenced by the order of the two tasks.

3.2 NIRS Data

Figure 8 shows the trial-averaged oxy-Hb time-course data for participant 1 (Exp. 1), measured with a WOT system, as representative data. The horizontal axis represents time, and the vertical axis represents oxy-Hb change (ΔCL). The start time of counting is $t = 1$ s. The trial-averaged oxy-Hb change obtained in the control task was subtracted from the data obtained at each time point for oxy-Hb in the cooperative task. The cooperative-minus-control data was thus obtained. The trial-averaged control-task data were calculated by using data from all the trials because there was no feedback sound in the control task. The cooperative-minus-control data of trials 2–10 were averaged to provide the data plotted in Fig. 8.

According to Fig. 8, during the counting period, oxy-Hb change in the middle and right-side prefrontal area increases, whereas that at both ends of the prefrontal area decreases. The spatial patterns of the changes in oxy-Hb change vary according to the participant. In the case of many of the participants, however, it is very common for the middle frontal area to be activated. Figure 9 shows a gray-scale map of AVs (at each analyzed channel) averaged from data of all the participants in trials 2–10. In Fig. 9, the color density of each element in the map represents the AV in the gray-scale bar. It is clear that the AVs in the middle frontal area are higher on average.

3.3 Relationship between Covariance of Two AV Maps and Intertime Intervals between Button Presses

The trials, except for the first trials, were divided into two groups: one including the trials in which the normalized covariance of the AV maps had a positive value (high-covariance trials), and one including those with a negative value (low-covariance trials). To compare the performance of the two groups, a Wilcoxon rank-sum test using the inter-time intervals of the two groups was conducted.

Figure 10 plots the mean and standard error of the inter-time intervals of the low- and high-covariance groups; the numbers of data of those groups are 21 and 32, respectively. The error bars represent the standard errors of normalized intertime interval data. It was found that the intertime intervals of the high-covariance trials are significantly shorter than those of the low-covariance trials ($p < 0.01$). When only the control-task data were used for calculating covariances, no relationship between normalized covariances and intertime intervals was obtained (Wilcoxon rank-sum test, $p > 0.4$). The number of combinations for selecting two people out of the 12 participants is ${}_{12}C_2 = 66$ patterns in total. The number of combinations of paired participants in this study is six patterns; thus, the number of combinations of unpaired participants is 60 patterns in total. To confirm the pair specificity, the spatiotemporal covariances of the 60 unpaired participants were analyzed in the same way. This analysis found that there was not a significant difference between the intertime intervals of the higher- and lower-covariance

Fig. 8 Trial-averaged oxy-Hb time-course data for participant 1 (Exp. 1) measured with a WOT system. The data of all channels (including both the used and interpolated channels) are shown. The horizontal axis represents time, and the vertical axis represents the oxy-Hb change. The start time of counting is $t = 1$ s.

groups (Wilcoxon rank-sum test, $p > 0.6$). These results show that when the spatiotemporal brain-activity patterns of the two participants were more synchronous, the time interval between their button-press times became shorter.

4 Discussion

4.1 Adjusted Button-Press Time in Cooperative Task

According to the behavioral results (Fig. 6), during the cooperative task, each participant seems to have adjusted his or her button-press time in each trial. In the ninth trial of Exp. 1, the fourth and ninth trials of Exp. 2, fifth and tenth trials of Exp. 3, seventh trial of Exp. 4, and eighth trial of Exp. 6, however, the intertime interval increased, possibly because of overadjusting. The effect of overadjusting is that the order of the button presses of the participants is inverted and the intertime interval becomes larger than the last one. The overadjusting trials are in the second half of the ten trials, except for the fourth trial of Exp. 2. That trend might reflect a break in concentration or the limited stability of the 10-s time-production task.

4.2 Relationship between Brain-Activation Patterns and Task Performance

The relationship between the synchronized pattern of the activated area during the counting period and cooperative-task

Fig. 9 Gray-scale map of activation values at each analyzed channel. Activation values are averaged from data of 12 participants in trials 2–10.

performance was investigated. To investigate the brain activities under a cooperative condition, the data obtained during counting period before button presses was used. It was found that when the spatial brain-activity patterns during the counting period of the two participants were more synchronous, the intertime interval between their button-press times became shorter. This result cannot be contaminated by a motion artifact because the data recorded after button presses was not used in the analysis. The possible reasons to explain this result are presented in the following.

It has been reported that when a human performs a branching task, which involves the process of integrating working memory with attention-resource allocation, the frontopolar prefrontal cortex is activated.⁶⁶ The cooperative task used in this study possibly requires two simultaneous mental processes: time counting and thinking about the timing of the button press in order to adjust it to that of the other person. It is therefore possible that the

Fig. 10 Mean and standard error of intertime intervals for low- and high-covariance groups. Black squares represent the mean value, and error bars represent the standard errors of the normalized inter-time intervals.

frontopolar prefrontal cortices of the two participants' brains during this cooperative task might be synchronously activated because they perform a kind of branching task at the same time.

The difference between the cooperative task and control task consisted of whether or not feedback was provided to the participants. It is possible that providing the participants with feedback made them think about each other. During the cooperative task, each participant might consider the adjusting precision and feedback gain (slight or drastic adjustment, which may be dependent on individuals) of another participant in order to prevent overadjusting. Even if these processes might not be actual cooperation or communication, they are important in inferring thinking or future activities of others. It has been reported that when people "mentalize," namely, think about other people's intentions, the mPFC region is activated.^{15,67–69} If adjusting the timing to other people during the cooperative task can be considered a kind of mentalizing process, then the AV map shown in Fig. 9 might reflect the activation of the superficial mPFC region.

4.3 Effects of Motor Planning and Intrinsic Fluctuation

It has been reported that the lateral prefrontal cortex plays a key role in preparing or planning an intended movement or action.^{70–72} The cooperative and control tasks used in this study include a common motor-planning process, that is, planning to press a button at certain time. The effect of the motor-planning function in regard to the difference between the spatiotemporal activity patterns during the cooperative and control tasks was therefore considered negligible.

Another possibility accounting for the high cooperative-task performances (i.e., short intertime intervals) with high covariances between the two participants' brains might be the influence of intrinsic fluctuations within cortical systems.⁷³ However, because the covariances of the AV maps obtained in the cooperative and control tasks are significantly different in terms of the relationship with the task performance, it is reasonable to assume that the results reflect the influence of the cooperative behavior of the two participants.

4.4 Limitations

In each experiment, neither the degree of interhuman relations between any two participants nor their personality characteristics regarding adaptability or cooperativeness was controlled. Moreover, all of the participants recruited in this study already knew each other. The cooperative-task performance might change in accordance with factors that are affected by interhuman relationships, such as whether the participants know each other and the personalities of the participants. These factors should be investigated further.

In addition, the following four limitations exist, in principle, when the oxy- and deoxy-Hb signals obtained from a participant are compared to those from another participant. The first one is the difference in the structure (thickness of scalp, skull, and gray/white matter) of the head,⁷⁴ which causes a difference in penetration depth and path length of the near-infrared light.⁷⁵ If the structural details of the head were measured by x-ray computed tomography or MRI scanner, the Hb-change signal could

be more precisely quantified by using the simulated optical path length.⁷⁶ The second limitation is the variation of probe positions. Figure 2 shows that the probe positions for each participant vary slightly. To reduce the effects of difference in the activated positions when the AV maps of two participants are compared, interpolated channels (in addition to the measurement channels) were used. It was thus possible to robustly calculate the covariance of the AV maps and analyze the relation between the two brains of a participant pair. The standard deviations of the measurement positions on the participants who participated in this study are within 10 mm, which is regarded as being sufficiently small for an OT study using a 30-mm source-detector distance. If the optodes were placed on the scalp on the basis of the probabilistic registration⁵⁹ using three-dimensional position data, the accuracy of interparticipant comparison would be better. The third limitation is a diverse brain-activation response induced by a participant-dependent hemodynamic response. An analytical method that takes account of the phase difference between NIRS signals by using a Hilbert transform⁷⁷ could improve the results presented here.

5 Conclusion

To investigate the relationship between mutually interactive brain-activation signals and cooperative-task performance, six pairs of participants took part in trials in which each pair of participants were instructed to count 10 s in their mind after hearing an auditory cue and then simultaneously press a button. The obtained brain-activity data of two participants were analyzed by calculating the covariance of AV maps. It was found that the synchronized activation patterns of the brains of the two participants of each pair during counting before the button presses is associated with the intertime interval between the two button presses of each pair when the participants tried to adjust their button-press timings to each other. This result cannot be explained in terms of a motion artifact; instead, it suggests that there is a relationship between synchronization of the brain activities and cooperative-task performance.

It is concluded that the spatiotemporal coherence of the NIRS signals between the paired participants is associated with their performance of a cooperative activity. The effect of interhuman relationships and personality on the NIRS signals should be further investigated. It was demonstrated that NIRS hyperscanning can be used to evaluate human interaction. Human-brain functions related to social skills should thus be progressively studied by taking an approach using coinstantaneous spatiotemporal activation patterns of two or more participants. A WOT system, which measures multiple brains in a shared space in natural manners, will contribute to a new research field of interbrain interaction in everyday life.

Acknowledgments

We thank Dr. Atsushi Maki, Dr. Takusige Katura, Dr. Akiko Obata, Dr. Hirokazu Tanaka, and Ryuta Aoki for their helpful comments on our research. We also thank Dr. Kazuo Saitoh for contributing to our fruitful discussions and Dr. Haruo Takeda for his overall support.

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