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**Short-term motor training, but not observational training,
alters neurocognitive mechanisms of action processing in
infancy**

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Running head: UNIQUE MOTOR ACTIVITY FOR ACTIVE LEARNING

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18 Short-term motor training, but not observational training, alters neurocognitive
19 mechanisms of action processing in infancy
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25 Gerson, S. A.,¹ Bekkering, H.,¹ & Hunnius, S.¹
26

27 Radboud University Nijmegen, Donders Institute for Brain, Cognition, and Behaviour
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43 **Corresponding author:**
44

45 Sarah Gerson
46

47 Montessorilaan 3
48

49 6522CC Nijmegen
50

51 The Netherlands
52

53 +31 (06) 29 77 81 35
54

55 s.gerson@donders.ru.nl
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Research Highlights

- We examined the effect of active versus observational experience on the neural correlates of action perception in 10-month-olds
- We found more motor activity to the effects of actively learned actions than to actions only observed
- We can conclude that active experience is critical to action perception on a neural level
- The findings shed light on the developmental origins of shared neural representations

For Review Only

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Abstract:

The role of motor experience in the processing of perceived actions is hotly debated on both behavioral (e.g., action understanding) and neural (e.g., activation of the motor system) levels of interpretation. Whereas some researchers focus on the role of motor experience in the understanding of and motor activity associated with perceived actions, others emphasize the role of visual experience with the perceived actions. The question of whether prior firsthand motor experience is critical to motor system activation during perception of actions performed by others is best addressed through studies with infants who have a limited repertoire of motor actions. In this way, infants can receive motor or visual training with novel actions that are not mere recombinations of previously acquired actions. In this study, 10-month-old infants received active training with a motorically unfamiliar action that resulted in a distinct sound effect. They received observational experience with a second, similarly unfamiliar action. Following training, we assessed infants' neural motor activity via electroencephalography (EEG) while they listened to the sounds associated with the actions relative to a novel sound. We found a greater decrease in mu power to sounds associated with the motorically learned action than to those associated with the observed action that the infants had never produced. This effect was directly related to individual differences in the degree of motor learning via motor training. These findings indicate a unique effect of active experience on neural correlates of action perception.

Keywords: action production; action perception; cognitive development; infancy; learning

Short-term motor training, but not observational training, alters neurocognitive mechanisms of action processing

Links between action and perception have been established across several species, throughout development, and via a variety of measures. Across these findings, the general notion is that action and perception of goal-directed actions are connected via shared neural representations and that this connection allows us to use our action experience to recognize and predict the goals of others' actions (Prinz, 1997). The identifying feature of these shared representations, the so-called *mirror system*, is that similar brain regions or neurons (as measured via fMRI, EEG, single cell recording, or other neuroimaging measures) are active when performing an action and perceiving the same action (or the sound or effect of that action; Kohler et al., 2002).

The Controversial Role of Expertise in Mirror System Activation

To understand the intricate relation between perception and action and differentiate the roles of motoric and visual experience, researchers are investigating the links between motor expertise, action perception, and activation of the motor system. One avenue of investigation that aims to parse out unique effects of motor experience, exemplified by Casile and Giese (2006), is to directly examine the role of motor experience on action perception, independent of any visual experience. In this experiment, participants were trained to perform a novel movement while blindfolded. Their accuracy in identifying this movement via visually presented point light displays improved after motor training despite a lack of visual information during training. Interestingly, individual differences in visual recognition performance strongly correlated with participants' accurate performance of the newly learned action during blindfolded training.

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3 Similar effects of experience on brain activation over motor regions further
4 corroborate the behavioral effects of motor training. For example, Calvo-Merino and
5 colleagues (2006) measured brain activation during the observation of dance moves
6 professional dancers had previously performed relative to moves with which they had
7 visual familiarity but no motor experience (i.e., moves produced by a dance partner of
8 the opposite gender). They found increased premotor, parietal, and cerebellar activity
9 for those actions within the dancers' own motor repertoires relative to the visually, but
10 not motorically, familiar actions. Cross and colleagues (2006) further showed that
11 training dancers to perform novel dance moves increased premotor activity when
12 viewing the trained, but not untrained, actions. Similar effects of expertise on motor
13 activation and activation of larger mentalizing brain networks have been found for
14 chopstick use (Järveläinen, Schuermann, & Hari, 2004), handwriting (Quandt,
15 Marshall, Bouquet, Young, & Shipley, 2011), sports (Aglioti, Cesari, Romani, &
16 Urgesi, 2008; Kim et al., 2011), and music (Bangert et al., 2006).
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34 In contrast to the evidenced differences between motor experts and non-
35 experts in motor activation across domains, other research indicates comparable or
36 increased motor activity to motorically unfamiliar actions. For example, in a follow-up
37 to the Cross et al. study (2006), the researchers compared physical and
38 observational learning of dance moves and found increased activity in premotor and
39 parietal regions for both types of actions (relative to actions not performed/watched,
40 Cross et al., 2006; see also Cross et al., 2009, 2012). Thus, the current state of the
41 adult literature on experience is inconclusive. Development provides a unique
42 window in which to look at effects of experience on neural activation of the motor
43 system because infants have a more limited repertoire of experience on which to
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draw. Further, intervening in early development allows one to manipulate the timing of naturally emerging motor actions and study the direct effects of this manipulation.

A Developmental Perspective on Active Experience

Both behavioral and neuroimaging studies have been conducted with infants in the first two years of life to examine the role of active experience on action perception. In intervention studies, infants are trained to perform novel actions and their recognition of the goal of these actions is then assessed. For instance, three-month-old infants who were trained to perform reaching actions (using Velcro mittens), but not untrained infants, recognized the goal of a reaching action (Sommerville, Woodward, & Needham, 2005). Importantly, several studies have now contrasted active with observational experience. When three-month-old infants were given active versus observational training with reaching actions, only those who received active experience benefited from training (Gerson & Woodward, 2014a, b). Similarly, 10-month-old infants benefited from active, but not observational, training with cane-pulling actions (similar to cloth-pulling; Sommerville, Hildebrand, & Crane, 2008). Investigating the role of motor, relative to visual, experience is crucial for understanding the role that the motor system plays in action perception.

Other researchers have focused on neural measures to assess effects of active experience on the motor regions of the infant brain. The most common assessment of motor activity in the infant brain is the measure of the so-called “mu rhythm,” measured via electroencephalography (EEG). In both infants and adults, reduced power (relative to a baseline condition) in the alpha frequency bands (approximately 8-12 Hz in adults and 6-9 Hz in infants; Marshall, Bar-Haim, & Fox, 2002) has been found over motor regions of the brain (central and frontocentral electrodes) both when individuals perform and observe goal-directed actions

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3 (Cuevas, Cannon, Yoo, & Fox, 2014; Marshall & Meltzoff, 2012, 2014; Vanderwert,
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5 Fox, & Ferrari, 2013). Using this measure, some studies reveal striking parallels to
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7 the neuroimaging research on expertise in adults and the behavioral evidence in
8
9 infancy. For example, Paulus and colleagues (2012) found motor activity in 8-month-
10
11 olds during perception of sounds associated with previously learned actions but not
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13 sounds associated with objects on which infants did not act during a training phase
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15 (see also Lloyd-Fox, Wu, Richards, Elwell, & Johnson, 2013; van Elk, van Schie,
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17 Hunnius, Vesper, & Bekkering, 2008). This research, however, examined actions with
18
19 which infants already had previous experience and did not assess whether newly
20
21 acquired actions led to similar effects in the motor system.
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25 Further, as with adults, data concerning the role of experience in infancy are
26
27 not all consistent. In a recent study by Southgate and Begus (2013), there was no
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29 difference between motor activation during the observation of executable and non-
30
31 executable actions (e.g., movement of objects by a claw) in 9-month-olds. In a follow-
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33 up to the Paulus study (Paulus, Hunnius, & Bekkering, 2013), infants saw their
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35 parents shake a rattle that made a novel sound. These infants later showed motor
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37 activity to the sounds associated with the observed action. In this study, however, the
38
39 observed action was one that infants were capable of producing prior to the training
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41 study. Thus, it is still unknown whether such mapping via observation would occur
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43 with motorically unfamiliar actions.
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47 In sum, the question of whether active experience is crucial to processes
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49 underlying action perception is still open. In this experiment, we examine the
50
51 possibility that a close link between action production and perception early in
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53 development can be built upon with greater experience. According to this proposal,
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55 given that rattle shaking in the Paulus et al. (2013) study was within the motor
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3 repertoire of these infants, the subsequent motor activation to the sound associated
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5 with the observed shaking could be a function of the infant mirroring the shaking
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7 action during training. Rather than relying on assumptions of previous action
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9 experience, we created new experiences with a novel action and assessed the
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11 effects of this experience on neural responses to action perception. To do so, we
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13 manipulated young infants' experience with unfamiliar actions and measured the
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15 effects of motor versus observational learning of novel actions on the motor system.
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17 This training manipulation closely matches those used in behavioral research with
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19 infants that have found unique effects of active, above and beyond, observational
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21 experience (Gerson & Woodward, 2014a, b; Sommerville et al., 2008). Incorporating
22
23 an interventional training design with neural measures allows us, for the first time, to
24
25 examine causal claims about the effects of active versus observational experience on
26
27 action-perception links on a neural level and test the assumption that active
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29 experience is critical to motor activity during action perception early in development.
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31 We expected more motor activity to sounds associated with actively learned actions
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33 than observed actions that the infants had never produced (Elsner & Hommel, 2001;
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35 Hunnius & Bekkering, 2014; Woodward & Gerson, 2014). Further, we hypothesized
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37 that those infants who were better able to perform the learned action at the end of
38
39 training would have a more precise motor representation of the action and thus show
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41 greater motor activity to the sound associated with the learned action.
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47 **Methods**

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49 **Participants:** Twenty-six infants were scheduled based on previous research
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51 and seventeen infants were included in the final sample for this study. Nine infants
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53 began the study but were not included in final analyses due to not returning for the
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55 EEG session after training (n = 1) or not sitting through at least nine trials of each
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3 condition prior to movement and artifact rejection ($n = 8$). No infants were excluded
4
5 from final analyses due to outlying data points. As suggested in recent reviews
6
7 (Cuevas et al., 2014), we also examined whether data was similar when we removed
8
9 outliers from analyses. After calculating the Event Related Desynchronization (ERD:
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11 the relative decrease in power to the event versus baseline, as described in the
12
13 Results section below), any infant who had an ERD greater than three standard
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15 deviations above the mean at any site and in any condition. This excluded four
16
17 infants and we saw the same pattern of results (active ERD lower than observational
18
19 ERD). The final dataset of infants ranged in age from 9 months, 16 days to 10
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21 months, 16 days at the pre-training session (mean age = 10 months, 0 days).
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23 Research was approved by the local ethics board.
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27 **Materials and Procedure:** Infants' participation in this study consisted of three
28
29 phases: a pre-training behavioral session, behavioral training sessions at home, and
30
31 a post-training EEG session. In each of these sessions, the same two toys were used
32
33 during the behavioral portion (see Figure 1). Both toys afforded means-end actions
34
35 that resulted in a unique sound effect (a series of tones that lasted approximately two
36
37 seconds). One toy had a wooden puppet atop a rectangular box (10.5 X 10.5 X 17
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39 cm). When the box was hit with a staff-like tool (approximately 34 cm long), the
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41 puppet spun around and a sound was played. The second toy was a rectangular box
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43 (31 X 25 X 11.5 cm) with an indentation at the front. When a cane-like tool
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45 (approximately 40 cm) was used to pull a toy duck into the indentation, a sound was
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47 played. The particular sound associated with each toy was counterbalanced across
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49 infants. The toy on which the infant performed versus observed an action was also
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51 counterbalanced.
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3 **Pre-Training Session:** In the first session, infants observed the experimenter
4 perform the appropriate means-end action on one of the two above-described toys
5 (and heard the associated sound). After the experimenter performed the action, she
6 asked the parent to perform the action and made sure the parent was demonstrating
7 the action in a similar fashion (e.g., drawing the infant's attention to the toy during the
8 action and during the playing of the sound, if necessary). The infant was then taught
9 to perform the action on the other toy. The action was demonstrated for the infant
10 and the infant was encouraged to attempt the action his or herself. If the infant did
11 not act, the experimenter encouraged the parent to help move the child's hand
12 toward the tool and coach them through the action in scaffolded steps. After training
13 the child and the parent on each of the two actions, the experimenter gave the parent
14 a schedule for training throughout the following week. The experimenter also helped
15 the parent learn how to use the camera that they would take home to record home
16 training sessions.
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33 **Training Sessions:** Parents were asked to have their child perform the
34 trained action every alternate day for approximately five minutes each day between
35 the pre- and post-training sessions. On the alternate days, the parent demonstrated
36 the other action for the child. As during the pre-training session, parents were told to
37 draw their infant's attention to the action when observing and to help their child if
38 necessary when performing. Parents were asked to record all sessions so that
39 coding of activity could be assessed offline via digital video. Coding of these videos
40 indicated that parents typically practiced each action approximately three times
41 between the pre-training and post-training session (range: 3-5). The average length
42 of each training session was longer ($p = .02$) for the active training sessions (5:21
43 minutes) than for the observational training sessions (4:15 minutes). Despite the
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3 difference in length, infants viewed, on average, more successful actions per session
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5 ($p < .001$) during the observational training sessions ($M = 16.18$) than the active
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7 training sessions ($M = 9.64$).
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10 **Post-Training Session:** Post-training sessions occurred between six and
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12 eleven days (mean intersession length was 7.41 days) following the pre-training
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14 session. Infants were first familiarized with the room and the experimenters and then
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16 fitted with a 32 active electrode infant-sized EEG cap (Brain Products, Munich).
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18 Infants then sat on their parent's lap in a shielded room in front of a monitor that
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20 displayed abstract pictures that were randomly changed every 1600 to 5000ms
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22 (jittered timing between picture presentation). Pictures were an attempt to maintain
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24 infants' attention and were unrelated to the test stimuli. Audio stimuli were played
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26 from a central speaker every 2600 to 5000ms (jittered presentation time and
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28 unrelated to of the presentation of the pictures). Each audio stimulus lasted 2000ms
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30 and consisted of three different sounds: the sound associated with the performed
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32 action during training (which differed between infants due to counterbalancing), the
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34 sound associated with the observed action, and a novel sound. The order of sound
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36 presentation was pseudorandomized, with the constraint that each sound was always
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38 repeated two times (and never more than two times in a row). Each sound was
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40 presented twenty times throughout the session. Following the picture and sound
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42 presentation, infants had the opportunity to perform each of the two actions (while
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44 EEG was recorded). Infants first performed the action they had practiced at home
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46 and then the action they had only previously observed.
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51 52 **Coding and Analysis:**

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54 **Coding and Exclusion of Movement:** Each video was coded offline for infant
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56 movement. No sound was played during coding so that the coder was blind to the
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condition of trials that were identified as consisting of movement. All movements were coded and identified as gross movement (e.g., turning toward the parent), fine movement (e.g., moving one finger), or movement that resembled grasping. For final analyses, all trials with movement were removed. On average, 37 out of the 60 possible trials were removed per infant due to movement ($SD = 8.53$, range: 22-50).

Coding of Actions: At the end of the EEG session, all infants had the chance to perform each of the two actions. EEG collection was continuous throughout this portion. Five randomly selected segments of the EEG data collected during infants' action performance (for each infant) was used to identify the frequency bands used in our analyses (see below). We presented them with the object on which they had received active training first and then presented them with the object they had previously only observed. A trained coder then assessed, for each infant, his or her actions on each object. The videos were clipped and presented in random order so the coder was blind to the infant's experience with each object. The coder counted the number of times each action was successfully produced and whether the child produced the action successfully on his or her own or did so with the help of the parent or experimenter. Infants were assigned a score based on their actions on each object: Never Performed Alone (if they only ever performed the action with help or never performed the action at all) or Performed Alone (if they ever performed the action without help from the parent or experimenter). The scores were ordinal such that any infant who both performed the action with a parent and on their own was scored as Performed Alone. For actions on the actively learned object, nine infants were in the Never Performed Alone group and eight infants were in the Performed Alone group. For actions on the observed object, eleven infants were in the Never Performed Alone group and six were in the Performed Alone group.

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3 **EEG Analysis:** Thirty-two active electrodes were arranged in the 10-20
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5 system and referenced online to FCz. The signal was amplified using a 32-channel
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7 BrainAmp DC EEG amplifier, band-pass filtered (.1-125 Hz), and digitized at 500 Hz.
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9 We kept impedances below 60 k Ω . We analyzed the data using FieldTrip, an open
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11 source Matlab (version 7.0, TheMathWorks, Inc.) toolbox developed at the Donders
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13 Institute for Brain, Cognition and Behaviour (<http://www.ru.nl/neuroimaging/fieldtrip>).
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15 Trials were identified by the onset of the each sound and lasted for the length of the
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17 sound (i.e. about two seconds). Trials during which infant movement was coded were
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19 excluded from further analyses. We then visually inspected the remaining trials to
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21 exclude EEG artifacts such as noisy channels. On average, approximately nine trials
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23 (range 3-20) remained for each condition for each subject after movement and
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25 artifact rejection. The number of trials analyzed did not differ between conditions.
26
27 When including only those infants who had 6 or more trials per condition
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29 (approximately 100/170 data points), results remained consistent (i.e., a significant
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31 effect of condition remained, $p = .05$). A bandpass filter was used with a frequency
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33 range of 1 to 30 Hz. A fast Fourier transform was then conducted using a multitaper
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35 method with a Hanning taper and a 2 Hz smoothing box to determine spectral power
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37 estimates for each condition from 1 to 30 Hz.
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43 **Results**

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45 In order to verify that the mu rhythm was in the 6-9 Hz range for our sample
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47 (as suggested by previous research; Cuevas et al., 2014; Marshall & Meltzoff, 2011),
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49 we compared power values across the 3-30 Hz range during periods of infant
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51 movement (i.e., when infants performed the actions at the end of the EEG session)
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53 relative to periods that were free of movement (i.e., collapsing across conditions to
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55 include all trials of sound presentation that were not removed due to movement). Mu
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3 rhythm has been defined as the frequency bands that are suppressed during
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5 movement relative to non-movement over motor regions. In order to account for
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7 variability in power ranges across infants, a proportional score was used to calculate
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9 relative power per band from 2-30 Hz: (power in each 1 Hz band)/(average power
10
11 across 2-30 Hz bands). As expected, a decrease in power for the movement, relative
12
13 to the still, phases of the experiment in the 6-9 Hz range confirmed our choice of
14
15 frequency bands for analyses (see Figure 2).
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19 Event-related desynchronization (ERD) was defined as (event-related power –
20
21 baseline power)/(baseline power) with power during presentation of the novel sound
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23 serving as baseline (Pfurtscheller, 2003; Vanderwert et al., 2013). Based on previous
24
25 research (Marshall & Meltzoff, 2012), mu power was calculated as power in the 6-9
26
27 Hz frequency range over frontocentral sites (C3, C4, Cz, FC1, and FC2). ERD was
28
29 calculated separately for time-locked to sounds associated with performed actions
30
31 and sounds associated with observed actions. A generalized linear model (GLM) with
32
33 ERD as the dependent variable and Condition (performed or observed), Region (C or
34
35 FC), and Site (1/3 [i.e., FC1 or C3], 2/4 [i.e., FC2 or C4], z [i.e., Cz]) as within
36
37 subjects factors revealed a main effect of condition, $F(1,152) = 6.20$, $p = .014$, $\eta_p^2 =$
38
39 $.039$, and no other main effects or interactions. The difference between conditions
40
41 was a function of a lower ERD in the performed than the observed condition.
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45
46 Identical GLM analyses that replaced frontal, parietal, and occipital regions
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48 with the central and frontocentral regions in the above-described GLM (e.g., ERD as
49
50 the dependent variable, Condition [performed or observed] and Site [3, 4, z] as within
51
52 subjects factors for Frontal sites) revealed no significant differences between
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54 conditions, $ps > .30$; $\eta_p^2 \leq .01$, indicating that the effect was localized to frontocentral
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3 sites (see Figure 3; *note: when outliers were excluded for these other regions, the*
4
5 *results remained the same*).
6

7
8 In an additional GLM (with Site and Region as within subjects factors), raw
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10 power data (i.e., not ERD corrected for baseline, so that activation to the novel
11
12 sound, previously baseline, could be compared directly with activation to the other
13
14 sounds) was compared between the sound associated with the performed and
15
16 observed action versus the novel sound. A significant decrease in power was found
17
18 in response to the sound associated with the actively trained action, $F(1,152) = 4.29$,
19
20 $p = .04$, $\eta_p^2 = .027$ (see Figure 4). There was no difference in power between the
21
22 observed versus novel sound, $p = .84$, $\eta_p^2 < .001$.
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26 In order to examine individual differences in training effectiveness, a final set
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28 of GLMs were conducted with condition as a fixed factor and region and site as
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30 within-subjects factors, ERD to the active sound as dependent variable, and ability to
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32 perform the learned action in the last session (Performed Alone or Never Performed
33
34 Alone) as a between-subjects factor. This revealed a significant interaction between
35
36 condition and performance ability, $F(1,142) = 3.24$, $p = .034$, $\eta_p^2 = .031$ (see Figure 5),
37
38 such that infants who had performed the action alone showed a significant difference
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40 in ERD to the sounds associated with the actively learned versus observed actions
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42 (mean difference = .64 [SEM = .19], $p = .001$), whereas infants who never performed
43
44 the action alone did not (mean difference = .08 [SEM = .18], $p = .68$). Because, at
45
46 the end of the final session, infants also had the chance to perform the action they
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48 had previously only observed, we also measured differences in infants' ability to
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50 perform this action at the end of training (though they only received observational
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52 training with this action). When the ability to perform the observed action was added
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54 as a covariate to the above GLM, no such interaction emerged, $p = .06$, $\eta_p^2 = .02$.
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3 This suggests that variability in learning the actively experienced action, but not
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5 variability in learning a similarly difficult (but only previously observed) action, drove
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7 these effects.
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9 10 **Discussion**

11 The results of the current study can be summarized in two points: First, motor
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13 experience with novel actions uniquely activates the neural motor system in response
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15 to perceiving the effect of this action, whereas observational experience does not.
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17 Second, this effect is directly related to individual differences in the degree of motor
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19 learning during active training. These findings suggest that active experience is
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21 critical to modulating motor activity during action perception early in development.
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23 These findings provide crucial information for the theoretical debates concerning the
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25 role of experience in action perception and motor activation. The effects of short-term
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27 training of novel motor actions on neural responses to the sensory consequences of
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29 those actions have never before been studied in infants. The training paradigm
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31 demonstrates a direct, causal effect of active experience on the neural correlates of
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33 action perception. Further, the fact that neural responses were related to infants'
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35 ability to produce the actively learned, but not observationally learned, action
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37 demonstrates specific effects on neural functioning of learning particular actions. This
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39 is the first study to teach infants novel motor actions and find a relation between each
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41 infant's ability to produce that action at the end of training and activation of their
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43 neural motor system when presented with the sensory consequence of that particular
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45 action. Finally, the measure of motor activation to audio presentation of action effects
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47 ruled out any possible effects of visual processing of the action, thus ensuring that
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49 the activation was not a function of visual processing.
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3 The discrepancy between our findings and those that suggest benefits from
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5 observational experience raise an important question: How can we move beyond
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7 initial motor experience to learn more broadly about actions and their effects in the
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9 environment? Infancy provides a unique window through which to look at the
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11 differential roles of motor and observational experience because of infants' limited
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13 repertoire of motor actions. Behaviorally, we saw no difference in the number of
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15 infants who could perform the actively versus observationally learned actions at the
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17 end of training. Importantly, there were no demonstrable benefits of learning from
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19 observation either on a group level or when infants were split based on whether they
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21 could perform the observationally learned action. Because action performance was
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23 measured at the end of the EEG session, it is an open question whether infants who
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25 could perform the observationally learned action would have shown a similar
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27 response to the sounds associated with that action (as those associated with the
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29 actively learned action) if they had had the chance to perform the action prior to the
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31 post-training EEG session. The precise nature of the motor representation created for
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33 each of these actions through training and how this relates to motor activity in the
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35 brain should be examined further in future research.
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40 In this study, infants had a week's worth of active and observational training
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42 with new, two-step actions. In natural contexts, longer-term learning of motor actions
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44 likely serves as a base for and contributes to infants' subsequent observational
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46 learning (Gerson, 2014; Woodward & Gerson, 2014). In this way, later in
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48 development, similar motor activity is likely to be seen for observed actions within
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50 and outside one's motor repertoire, as long as the person can achieve the goal of the
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52 action (in some way) using movements already within his or her motor system (see
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54 Aziz-Zadeh, Sheng, Liew, & Damsio, 2012; Woodward & Gerson, 2014).
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3 Southgate has presented a similar hypothesis to reconcile controversial
4 evidence of motor activation for actions within and outside infants' motor repertoire
5 (Southgate, 2013). In the Begus and Southgate (2013) experiment, the authors found
6 no indication of a difference in infants' motor activation in response to observing claw
7 actions and hand actions on an object. Similar activation for claw and hand actions
8 would be expected if infants were able to recognize the path of the claw and relate
9 this to an action they could use to achieve this goal (reaching with a hand). Reaching
10 with the hand is an action that is repeatedly practiced by 9-month-olds in their
11 everyday life and would thus easily be brought forward as an alternative action. It is
12 possible that increased experience with a variety of simple actions leads to motor
13 activation to the observation of a broader range of actions that can achieve similar
14 goals through the combination or substitution of the simple actions (Buccino & Riggio,
15 2006). After having gained motor experience with a particular action (e.g., shaking),
16 an infant can potentially map this motor representation, as in Paulus et al., to novel
17 effects and goals (e.g., a new sound) that become associated with the motor action
18 through observational, rather than active, experience (Paulus et al., 2013; cf. de
19 Klerk, Johnson, Heyes, & Southgate, 2014).

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41 Whether the observed effect was a function of a *causal* link between the motor
42 action and the sound is an open question. That is, it is unknown whether the motor
43 action necessarily needed to result in the sound in order for the association between
44 the sound and the motor system to be formed. For example, if infants were to
45 activate their own motor systems via play with another object while they viewed their
46 parents perform the (motorically unfamiliar) action that resulted in the sound, their
47 motor system might then later respond to the sound via association with the action
48 the infant had produced (that was unrelated to the sound except in "coincidental"
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3 timing). This possibility would be consistent with the theoretical proposal put forth by
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5 Paulus and colleagues (2008) and is a question for future research. The current
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7 findings are consistent with the unique behavioral effects of active relative to
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9 observational experience early in development (Gerson & Woodward, 2014a;
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11 Hunnius & Bekkering, 2014; Sommerville et al., 2008). Whether and how this change
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13 in motor activity is related to the change in perception of the goal of the observed
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15 action is an open question. To date, as far as we are aware, no studies have
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17 simultaneously measured individual differences in motor activation and perception of
18
19 the effect of an action. Instead, similar relations between expertise and behavioral
20
21 measures and expertise and motor activation have been hypothesized to support
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23 links between the behavioral and neural measures (e.g., Marshall & Meltzoff, 2014;
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25 Woodward & Gerson, 2014). Examining this relation more directly is important for
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27 uncovering the true nature of the link between motor activity and action
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29 understanding.
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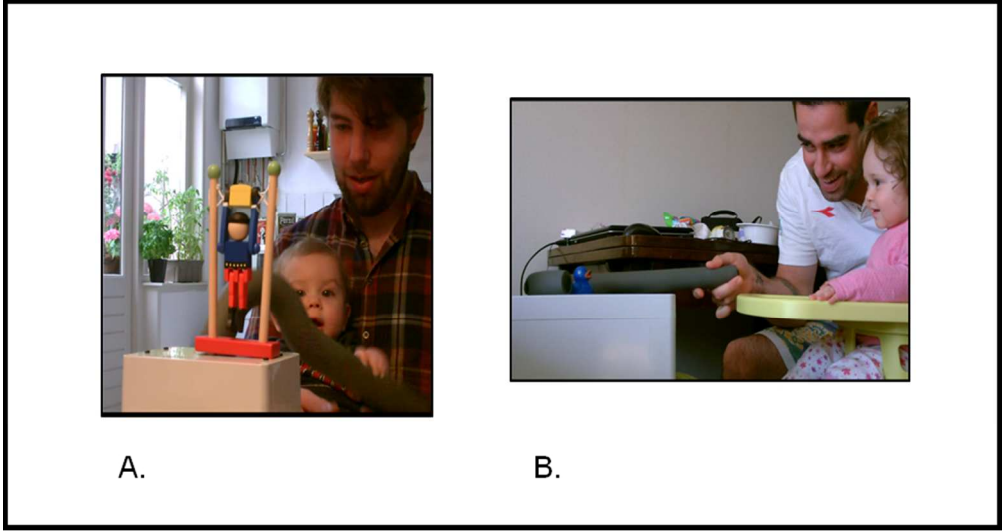


Fig. 1.
(A) An infant learning to perform a new action at home: hitting a toy with a stick and (B) an infant observing her dad perform a new action at home: pulling a duck with a cane
163x87mm (150 x 150 DPI)

view Only

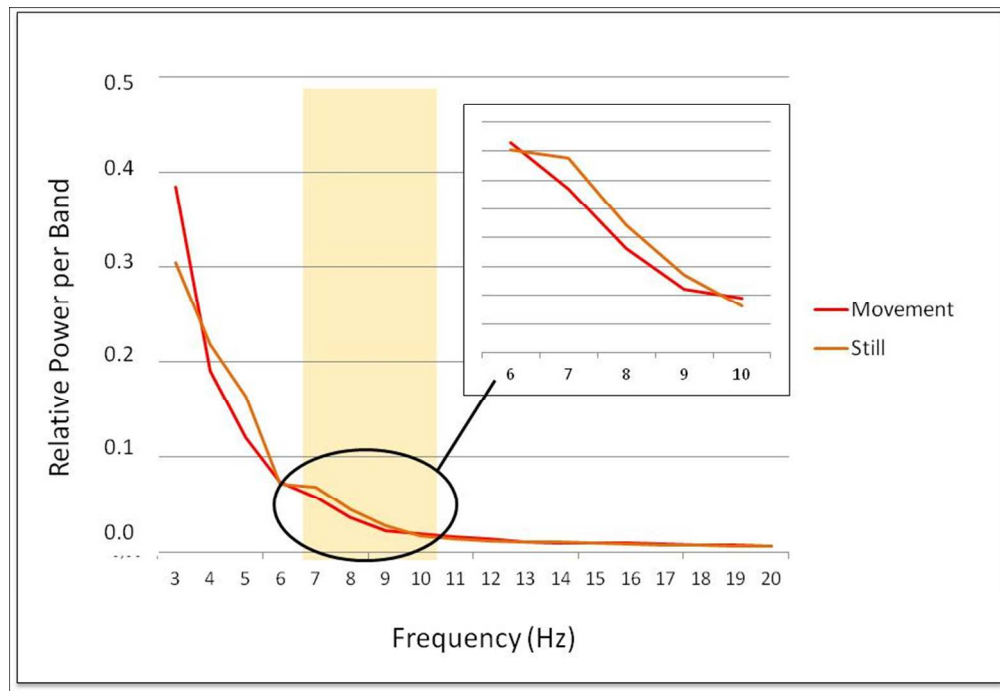


Fig. 2.
Relative power values as a function of frequency (Hz) depicted for still periods (collapsed across conditions) and execution of actions. The yellow shaded area illustrates the Alpha-frequency range.
208x143mm (120 x 120 DPI)

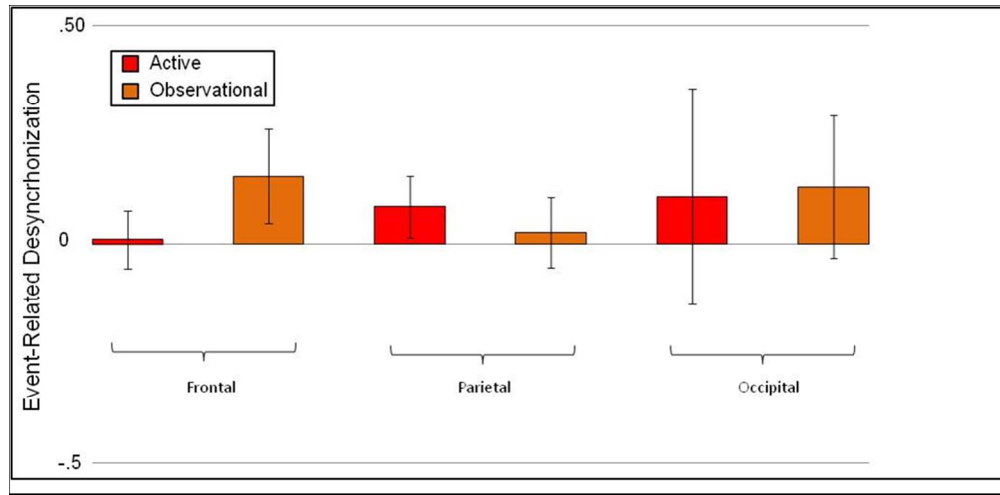


Fig. 3.

ERD (in frontocentral electrodes) to active and observational sounds was not significantly different in frontal, parietal, or occipital regions. ERD is defined as $(\text{event-related power} - \text{baseline power}) / (\text{baseline power})$ with power (6-9 Hz) during presentation of the novel sound serving as baseline. (outliers excluded; error bars represent standard errors)

220x107mm (120 x 120 DPI)

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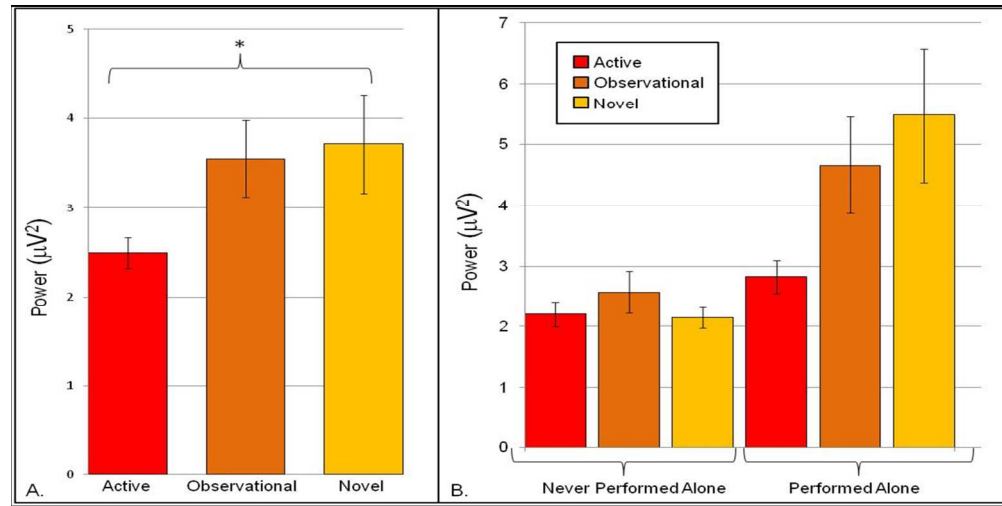


Fig. 4.
Raw power to active and observational sounds collapsed across all subjects (A) and split by infants' ability to perform the actively learned actions (B). (error bars represent standard errors, * $p < .05$)
236x148mm (150 x 120 DPI)

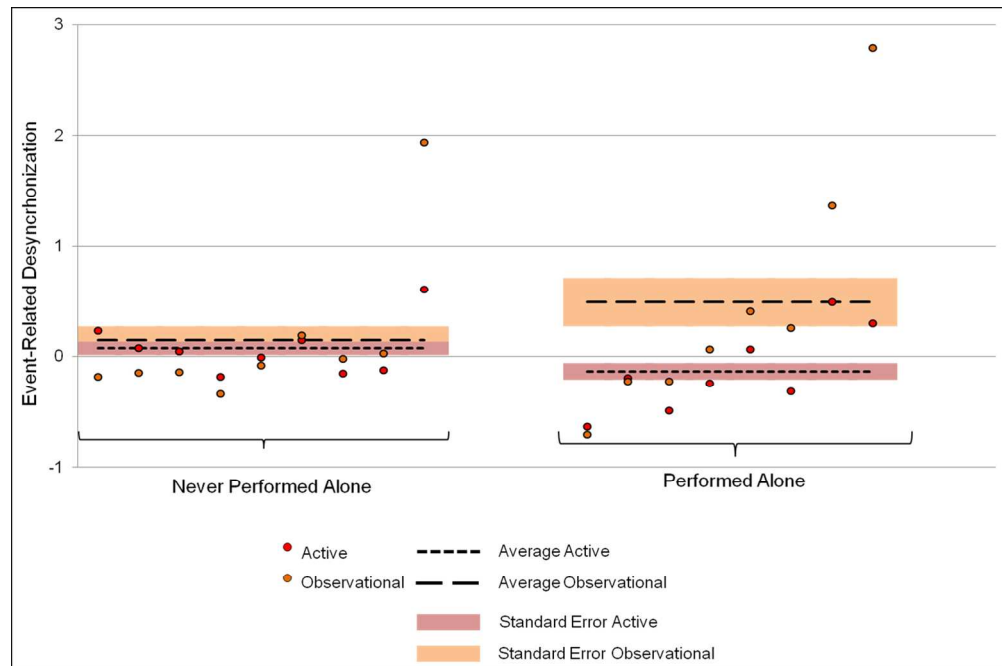


Fig. 5.

ERD (in frontocentral electrodes) plotted per subject, split as a function of infants' ability to produce the actively learned action at the end of training. Means across subjects within each group are plotted as lines and the shaded area represents the standard error. ERD is defined as (event-related power - baseline power)/(baseline power) with power (6-9 Hz) during presentation of the novel sound serving as baseline.
 254x168mm (150 x 150 DPI)