

Sports experience changes the neural processing of action language

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Experience alters behavior by producing enduring changes in the neural processes that support performance. For example, performing a specific action improves the execution of that action via changes in associated sensory and motor neural circuitry, and experience using language improves language comprehension by altering the anatomy and physiology of perisylvian neocortical brain regions. Here we provide evidence that specialized (sports) motor experience enhances action-related language understanding by recruitment of left dorsal lateral premotor cortex, a region normally devoted to higher-level action selection and implementation—even when there is no intention to perform a real action. Experience playing and watching sports has enduring effects on language understanding by changing the neural networks that subserve comprehension to incorporate areas active in performing sports skills. Without such experience, sport novices recruit lower-level sensory-motor regions, thought to support the instantiation of movement, during language processing, and activity in primary motor areas does not help comprehension. Thus, the language system is sufficiently plastic and dynamic to encompass expertise-related neural recruitment outside core language networks.

expertise | premotor | action planning | motor stimulation | comprehension

The mechanisms for language processing in adulthood are thought to be extremely stable and impervious to change (1). Yet, it may be that plastic change within the language system can arise from non-language-related activities. Background knowledge of a topic certainly aids comprehension and retention of language related to that topic (2–4) and such knowledge can even eliminate comprehension differences between those lower and higher in verbal ability (5). But what if one is not just knowledgeable in a particular area, but also has direct experience with the behaviors described in the language one uses? Could such experience actually modulate the neural substrates called on to support effective comprehension? When individuals hear language about action, they activate neural networks involved in producing these actions (6). Given that athletes and novices rely on different cognitive and neural operations during overt action production (7–12), the resources called on to support language comprehension might differ as a function of an individual's experience with the communicated actions. In the current work we ask whether athletic experience carries implications beyond the playing field to a very different activity: language comprehension, and more specifically, to the understanding of language about actions one has experience with. Furthermore, if a relationship does exist, is it based on conceptual knowledge of the motor skill or a more direct ability to use the neural networks subserving action to enhance language comprehension?

To answer this question, ice-hockey players (professional and intercollegiate players; $n = 12$), fans (extensive hockey viewing, but no playing experience; $n = 8$), and novices (no ice-hockey playing/viewing experience; $n = 9$) passively listened to sentences describing ice-hockey actions or everyday actions during functional magnetic resonance imaging (fMRI) and then performed a language task outside the scanner that gauged their

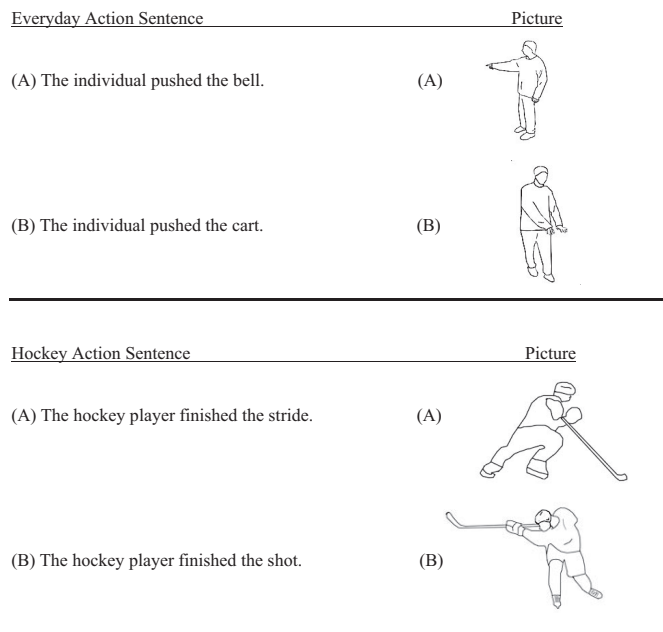


Fig. 1. Examples of the postscan comprehension task stimuli. Picture A serves as a match for Sentence A and a mismatch for Sentence B. Picture B serves as a match for Sentence B and a mismatch for Sentence A.

understanding of the heard scenarios (13–15). In this postscan comprehension task, subjects listened to sentences describing ice-hockey actions (e.g., “The hockey player finished the shot”) and everyday actions (e.g., “The individual pushed the door bell”). Following each sentence, participants were presented with a picture of an individual. Participants were told to judge as quickly as possible whether the pictured individual was mentioned in the sentence and to indicate this decision by pressing a “yes” or “no” button.

The pictures (Fig. 1) depicted either (i) an individual mentioned in the preceding sentence performing an action that matched the action implied in the sentence, (ii) an individual mentioned in the preceding sentence performing an action that was different from (or mismatched) the action implied in the sentence, or (iii) an individual not specifically mentioned in the sentence and performing an action that was not implied by

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Table 1. Significant regions for the whole-brain correlation maps for hockey action sentences at $P < 0.005$ (cluster-level corrected at $\alpha < 0.05$)

ROI (Brodmann's area)	Center of gravity			ROI size, mm ³	Correlation coefficient
	x	y	z		
<i>Left dorsal premotor cortex (BA6)</i>	-45	9	41	405	<i>0.608</i>
<i>Right dorsal primary sensory-motor cortex (BA4/1/2)</i>	22	-21	53	488	<i>-0.677</i>
Right post. cingulate gyrus (isthmus: BA30)	11	-35	2	1,178	-0.728
Left post. angular gyrus (BA39)	-39	-47	23	426	-0.644
Left post. cingulate gyrus (BA31)	-14	-54	27	639	-0.551
Left cuneus (BA18)	-14	-91	18	753	-0.597

Correlation coefficients are standardized r values ($N = 29$) and represent the average coefficient across all voxels in the cluster. Italicized regions are those that showed a significant relation between hockey experience and neural activity. Activity in these regions also significantly mediated the relation between hockey experience and hockey language comprehension (see Fig. 2). post., posterior.

the sentence. For this last category, pictures of hockey players were paired with everyday sentences (e.g., a picture of a hockey player skating followed the sentence, “The individual frosted the cake.”) and pictures of everyday individuals were paired with hockey sentences (e.g., a picture of a man buttoning his shirt followed the sentence, “The hockey player saved the shot.”). This last category required a “no” response (serving as fillers to equate “yes” and “no” responses across sentences) and was not analyzed. Trials of interest were those that required “yes” responses [i.e., (i) and (ii) above]. If individuals comprehend the actions described in the sentences (13–16), responses should be facilitated for pictured individuals whose actions match those implied in the sentence relative to those that do not. We termed this index of comprehension the action-match effect.

Results

Does hockey experience enhance language comprehension? All subjects, regardless of ice-hockey experience, showed a significant action-match effect for everyday sentences, responding faster to pictures of individuals performing actions that matched those implied in the sentences vs. pictures that did not [733 vs. 802 ms; $t(28) = 3.33$; $P < 0.01$; interaction with group, $F = 1$]. In contrast, only ice-hockey players [605 vs. 669 ms; $t(11) = 3.22$; $P < 0.01$] and fans [761 vs. 884 ms; $t(7) = 4.91$; $P < 0.01$] showed the action-match effect for hockey sentences. Novices' response times did not differ as a function of pictures that matched the sentence-implied hockey actions vs. those that did not [655 vs. 656 ms; $t(8) = 0.03$; $P = 0.98$]. Experience watching and playing ice-hockey facilitates comprehension specifically for hockey sentences.

To determine the neural mechanisms underlying this experience-dependent facilitation, we performed regression analyses relating neural activity during hockey-language listening to hockey experience and hockey-sentence comprehension. Importantly, we did not expect to find a significant relation among neural activity during everyday language listening, everyday sentence comprehension, and hockey experience because everyday language comprehension did not vary with hockey group. This expectation is not surprising given that everyone should have experience viewing and performing everyday actions.

We began by regressing, at the whole-brain level, our index of hockey sentence comprehension (i.e., the action-match effect) on neural activity measured while all subjects, irrespective of hockey experience, heard hockey sentences. Starting at the whole-brain level allows us to identify all possible neural areas that might mediate (or account for) the relation between hockey experience and hockey sentence comprehension. Neural activity in several brain regions was significantly related to hockey sentence comprehension (Table 1).

We next examined whether the regions that related to comprehension also significantly related to hockey experience. Here, and in the analyses below, we treated hockey experience as a categorical variable, rank ordering subjects based on hockey exposure from (1) no playing or viewing experience (novices), to (2) no playing, but viewing experience (fans), to (3) playing and viewing experience (players). This rank ordering, which captures relative degree of exposure to ice-hockey, allows us to test whether the facilitative effect of hockey experience on comprehension is mediated by neural activity during hockey language processing.⁸

Two brain regions related to hockey-sentence comprehension also related to the rank-ordering of hockey experience. Activation in left dorsal premotor cortex [Talairach center-of-gravity = ($\pm 45, 9, 41$)] positively correlated with hockey experience. Given that premotor regions are typically activated bilaterally within a larger network for action understanding in both action observation and language comprehension (17–19), we also looked at the analogous region in the opposite hemisphere. An interaction with hemisphere [$\beta = 0.29$, $t = 2.15$, $P < 0.05$] showed that this relation was limited to the left hemisphere ($r = 0.46$, $P < 0.02$). In addition, activation in right dorsal primary sensory-motor cortex [Tal ($\pm 22, -21, 53$)] negatively correlated with hockey experience ($r = -0.45$, $P < 0.02$). Not only did a lack of a hemisphere interaction ($P = 0.64$) support the bilaterality of the relation between primary sensory-motor activation and experience ($r = -0.43$, $P < 0.02$), but similarly high correlations were seen between average activation in right sensory-motor cortex and comprehension ($r = -0.68$, $P < 0.001$) and left primary sensory-motor cortex and comprehension ($r = -0.51$, $P < 0.005$). Thus, in the analyses from this point forward, we consider the relation among bilateral primary sensory-motor cortex activity, experience, and comprehension.

If the above relationships between neural activity and comprehension are because of experience-dependent changes in the neural networks subserving language understanding, then they should be limited to the domain in which subjects were selected to vary by experience (i.e., hockey). To test this hypothesis, we also correlated, at the whole-brain level, everyday action comprehension with neural activation while subjects listened to everyday sentences. Although one region, the left Lingual Gyrus [Tal ($\pm -10, -86, -9$)], related to comprehension ($r = 0.61$, $P < 0.001$), it was unrelated to hockey experience ($r = 0.15$, $P = 0.43$).

⁸Although our coding scheme was setup *a priori* to examine the relation between degree of exposure to ice-hockey and hockey-language comprehension, we also examined alternative ways of coding hockey experience. Considering fans and experts to be of equal experience, considering fans and novices to be of equal experience, or coding playing and watching experience separately did not produce the same significant correlations seen in the next paragraph and Fig. 2 between experience, comprehension, and neural activity, nor did it result in a better-fitting mediation model.

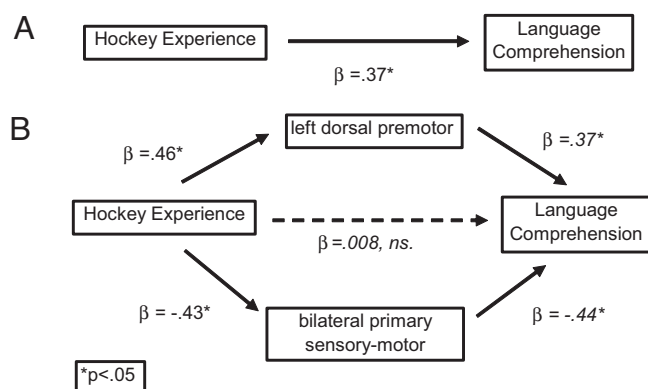


Fig. 2. Mediation analysis of the relation between hockey experience and hockey-action language comprehension. (A) A regression analysis established that hockey experience [as reflected in a rank ordering based on hockey exposure from (i) novices to (ii) fans to (iii) players] had a significant positive effect on hockey language comprehension ($\beta = 0.37$, $t = 2.06$, $P < 0.05$). (B) Hockey experience also had a significant positive effect on left dorsal premotor activity while passively listening to hockey-related sentences ($\beta = 0.46$, $t = 2.72$, $P < 0.02$) and bore a significant negative relation to bilateral primary sensory-motor activity while passively listening to hockey-related sentences ($\beta = -0.43$, $t = 2.49$, $P < 0.02$). Both left dorsal premotor activity ($\beta = 0.61$, $t = 3.98$, $P < 0.001$) and bilateral primary sensory-motor activity ($\beta = -0.64$, $t = 4.31$, $P < 0.001$) were significant predictors of language comprehension. When hockey experience, left dorsal premotor activity, and bilateral primary sensory-motor activity were simultaneously entered as predictors of hockey sentence comprehension, experience no longer significantly predicted comprehension ($\beta = 0.008$, ns), whereas both left dorsal premotor activity ($\beta = 0.37$, $t = 2.13$, $P < 0.05$) and bilateral primary sensory-motor activity ($\beta = -0.44$, $t = 2.55$, $P < 0.02$) remained significant in the equation. A Sobel test (45) of the reduction in the direct relation between experience and comprehension for both left premotor ($z = 2.24$, $P < 0.03$) and bilateral sensory-motor ($z = 2.80$, $P < 0.01$) was significant. This series of analyses provides support for our conclusion that hockey experience facilitates hockey language comprehension through relatively increased activity in left dorsal premotor cortex and relatively decreased activity in bilateral sensory-motor cortex.

Turning back to our original question regarding the impact of hockey experience on language comprehension, we asked whether the pattern of neural activity seen while listening to hockey language mediated (or accounted for) the impact of experience on hockey-language comprehension. As reflected above, the more hockey experience individuals had (ranging from novices to fans to players), the more effective they were in comprehending hockey sentences. This relation became nonsignificant when both left dorsal premotor and bilateral sensory-motor activity was also used to predict hockey comprehension. Only a strong positive relation between left dorsal premotor activity and hockey comprehension and a strong negative relation between bilateral primary sensory-motor activity and hockey comprehension remained (Fig. 2). The relation between experience and comprehension was fully mediated by left dorsal premotor and bilateral primary sensory-motor activity. Figs. 3 and 4 show the brain-behavior correlations, activation differences by hockey experience group, and the neural regions themselves.

Effective auditory comprehension of action-based language is accounted for by experience-dependent activation of the left dorsal premotor cortex, a region thought to support the selection of well learned action plans and procedures, often in response to learned symbolic associations (20–26). Evidence from lesion studies in monkeys (27) and humans (28, 29) shows that ablations of the dorsal premotor cortex prevents individuals from associating well learned motor response sequences (e.g., twisting a handle) with exogenous symbolic cues (e.g., a yellow placard or specific tone). Because subjects can independently perform the required movement se-

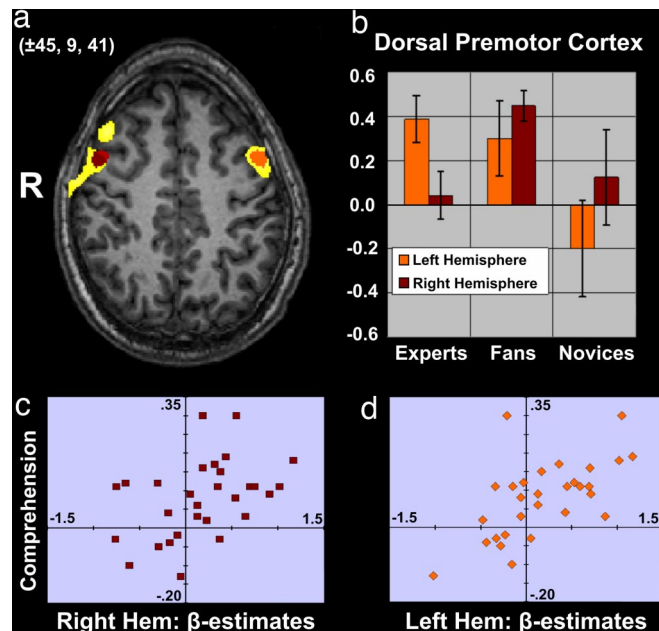


Fig. 3. Visualization of dorsal premotor cortex: positive correlation between neural activity when listening to hockey action sentences and postscan hockey language comprehension. (a) The left hemispheric region shown in orange was significant ($P < 0.005$, corrected) at the whole-brain level. To visualize this region as part of a larger premotor network, it is overlaid onto voxels showing a positive correlation at the reduced threshold of $P < 0.05$, uncorrected (depicted in yellow). To test for laterality, this region was reflected over the y axis (shown in the right hemisphere in dark red). (b) Activity in the left and right ROIs broken down by hockey experience group. The y axis represents neural activity in the form of β -estimates for the hockey action sentences. Error bars represent SEM. (c and d) Visualization of the linear relationship (across all 29 subjects) between neural activity (represented as β -estimates) (see Methods) and hockey language comprehension for each hemisphere (left hemisphere: $r = 0.61$, $P < 0.001$; right hemisphere: $r = 0.41$, $P = 0.05$).

quence and recognize the symbolic stimuli, the dorsal premotor cortex is believed to support the selection of well learned plans for action. The left hemisphere has been shown to drive this selection, regardless of the effector involved (24, 30–32). Substantial prior experience viewing and performing ice-hockey actions enhances hockey-language comprehension, likely by enabling individuals to associate linguistically described action scenarios with motor plans for execution. This ability, in turn, gives individuals the type of robust and multimodal representation that is the hallmark of optimal comprehension.

Interestingly, facilitated comprehension of hockey action sentences was not limited to those with significant hockey motor experience (i.e., hockey players); Ice-hockey fans also showed the action-match effect for hockey scenarios. This comprehension effect in fans was accompanied by activation in the left dorsal premotor cortex while listening to the hockey action scenarios that was significantly above baseline [$t(7) = 3.21$, $P < 0.02$]. Players showed a similar pattern [$t(11) = 3.50$, $P < 0.005$]. Novices did not ($P = 0.25$). Moreover, increased dorsal-premotor activation during hockey-sentence comprehension was seen bilaterally for the fans but not the players (Fig. 3). This observation is true whether one looks at the region of interest (ROI) directly contralateral to the left dorsal premotor region or the peak activation in the right prefrontal cortex more generally. The bilateral premotor activation may be indicative of more effortful action selection in fans vs. players, a view consistent with overall longer response times for the fans [mean (M) = 822 ms; SE = 73] vs. players (M = 637 ms; SE = 39) for hockey sentence comprehension [$t(18) = 2.44$, $P < 0.03$]. Nonetheless,

viduals. These filler trials were not analyzed. Across all trials, accuracy was high ($M = 89\%$, $SE = 3\%$) and did not differ as a function of hockey experience, sentence type, or picture.

The BOLD signal during fMRI was examined by using a multiple regression analysis of each functional time series (40) that took into account sentence type in a random-effects general linear model. The resulting hockey-action and everyday-action condition parameter estimates (β -estimates) in each voxel were submitted to separate whole-brain correlation analyses. In this analysis, the action-match effect [(Mismatch Reaction Time (RT) – Match RT)/Average RT] was regressed on the contrasts of hockey action sentences vs. baseline and everyday action sentences vs. baseline, respectively. This analysis was blind to hockey-experience-group membership. The statistical maps generated from these analyses were thresholded at $r(28) = 0.52$, $P < 0.005$, and subsequently corrected for multiple comparisons by using Monte Carlo simulation to yield a whole-brain significance of $\alpha = 0.05$ (41).

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