Social Interaction in Robotic Agents Emulating the Mirror Neuron Function

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Abstract. Emergent interactions that are expressed by the movements of two agents are discussed in this paper. The common coding principle is used to show how the mirror neuron system may facilitate interaction behaviour. Synchronization between neuron groups in different structures of the mirror neuron system are in the basis of the interaction behaviour. The robotics experimental setting is used to illustrate the method. The resulting synchronization and turn taking behaviours show the advantages of the mirror neuron paradigm for designing of socially meaningful behaviour.

1 Introduction

Recent neurophysiological, cognitive, and developmental research clearly shows that there are shared representations in the brain between perceived and generated actions, between actions produced by oneself and others (see for instance [20][27][15][16]). These shared representations, conveyed by the mirror neuron system, underlie the process of imitation, social learning, and prediction of the behaviour of conspecifics. Many attempts have been made to model the imitation process, for review see [26] and [17]. However, the imitation that has been modelled so far does not go further than one directional demonstrator-imitator interaction. In this paper we want to make an attempt to show the potential of the mirror neuron paradigm for social interaction, in particular for movement synchronisation, entrainment, and interchangeable turn-taking between two agents.

Entrainment of timing of social interaction has been investigated in multidisciplinary research on conversation. Conversation is an exchange of speech between two or more individuals. Although at first glance it looks like a chaotic process, conversation usually proceeds smoothly, by having the two parties take well timed turns. A number of authors have proposed that the listeners anticipate an upcoming end of a turn by perceiving eye gaze, body movement, or other semantic, syntactic, or prosodic queues from the speaker, for reviews see [8][9]. Conversely, listeners indicate their desire for turn ending. Speech is, in its essence, a motor act and it is likely that the mechanisms of speech and turn taking coevolved, perhaps building on the same preexisting structures and mechanisms for motor expression [29]. In their theoretical study Wilson and Wilson

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[29] argue that turn taking is likely to be successfully modelled by entrainment of endogenous oscillators.

Mutual entrainment of rhythmic activities has been theoretically studied as the basic mechanism of the organization of temporal order by Pavlidis [18]. Endogenous oscillators have been implicated in a range of cognitive processes, including perception, motor control, attention, memory, and consciousness [5].

In a robotic setup turn-taking behaviour is discussed in [6] and [12]. The turn taking behaviour in these studies takes place as a result of interaction of two dynamical recognizers - Elman type of recurrent neural networks that have widely been used to model dynamic systems. The training has been replaced by a genetic algorithm, which aims to produce "genetically different" agents. This will prevent from the low reliability of the interaction process based on neural learning [5].

We base our interaction behaviour on synchronization between neural latices that together simulate the mirror neuron-like functioning. The neuron firing in every lattice of neurons is modelled by an oscillatory model.

This paper is organized as following. In Section 2, we propose the biological background of the mirror neuron model and the common coding paradigm. Section 3 connects the biological modelling to concrete computational framework and shows how it is applied in a robot setting. The experimental setting and results of following and turn taking simulations are shown in Section 4. The discussion (Section 5) summarizes the results and puts this work in perspective.

2 Mirror Neuron System Model for Interagent Interaction

The Common coding paradigm postulates parity between perception and action, i.e. the action and perception arise simultaneously [11]. A core assumption of the Common coding paradigm is that actions are coded in terms of the perceivable effects (i.e. the distal perceptual events) that they should generate. It has the advantages to an information-processing paradigm which is unable to explain perception in many cases related to direct action [3][1]. Common coding paradigm has more solid foundations than Selection paradigm and Gibsons theory of direct perception [7] which fail to explain another group of phenomena like memory and imagination that can certainly originate an action by themselves [3][2].

A growing body of behavioural and neurophysiological studies support the grounding principles of the Common coding paradigm. As first evidence for direct matching between action perception and action execution came the discovery of 'mirror neurons' in the ventral premotor cortex of the macaque monkey [21][22][23]. Mirror neurons fire both when monkey carries out a goal-directed action and when it observes the same action performed by another individual [24], i.e. the perception and the action are likely coded in the same way, by the same structure. More recently, it was found that a subset of these mirror neurons also responds when the final part of a previously seen action is hidden and can only be inferred [28]. Therefore, the observation of an action activates action



Fig. 1. The common coding theory suggests that sensing and action is related to the activation in the same internal representations. Moreover they can be activated by endogenous factors. Common coding for two agents that share perceptual space as a basis for modelling interaction behaviour.

representations to the degree that the perceived action and the represented action are similar [14]. Specific neurons in this region respond to the representation of an action rather than to the action itself.

It can be inferred that the sensory and the motor activations that represent the same action or intention are related to the activation of the same area in the brain. One such an area is the ventral premotor cortex (PMv). Since the observed, executed, and imagined actions are related to an activation in a common representation, we schematically show this phenomena like the activation from the three events is projected to the common representation (Figure 1, the scheme of the individual agent). Actually, in case of an executed action the activation in the premotor and motor areas occurs in a very short for the behavioural time scale interval, i.e. practically co-occurs. The case of two agents that share perceptual space the common representation for perception and action for each agent will create a basis for an interaction behaviour, as shown in Figure 1.

Actually, there is more than one representational structure that gets active by the same event encountered by the sensory and the motor states. Most of the frontal motor areas receive robust sensory input (visual and somatosensory) from the parietal lobe. This pattern of connectivity supports relatively specialized fronto-parietal area for sensorimotor integration. A posterior area with mirror neuron properties is located in the rostral part of the inferior parietal lobule (IPL). Both areas form the mirror neuron system (MNS) The main visual input to the MNS originates from the posterior sector of the superior temporal sulcus (STS). Together, these three areas form a core circuit for imitation, one of the basic building components of social behaviour. The information flow from the parietal MNS, which is mostly concerned with the motoric description of an action, reaches back to the STS. By macaque, STS and the equivalent of IPL share patchy connections that overlap particularly well with the locations



Fig. 2. The information flow by the observation and the imitation of the same action in an individual agent. Double line arrows mark the path of the inverse model that signifies action observation. Single line arrows show the forward path for execution of imitated actions.

in which neurons respond specifically to complex body movements. The STS although considered to be a part of the 'mirror system' [19] do not show any motor activation itself. In spite of lacking mirror properties, STS neurons seem to 'understand' actions quite well, and it is plausible to assume that they send (via IPL) preprocessed signals about actions to the premotor areas that include information about the goal or the meaning of the observed action.

To construct a computational model that can facilitate the imitation and interaction functionality, we have modelled the three interconnected structures with lattices of neurons [4]. The direction of the connectivity between the structures differs while different actions take place. Figure 2 denotes the information flow by the observation and the imitation of the same action. The solid arrows show the part that has been considered in our model for achieving the imitation functionality.

3 Oscillatory Neural Dynamics of the Mirror Neuron System in the Robotics Setting

From the framework proposed in the previous Section becomes apparent that the mirror neuron model that materialises the Common coding paradigm is a useful tool for modelling interactive behaviour. Interagent interaction is initiated from the representation of the movement of each robot within the neural structures of the partner robot. To achieve the imitation functionality and create a model that is suitable for robotics, we have to make some simplification. We base our core scheme for imitation learning on conceptual model of Keisers and Perrett [13], whose experimental work has shown that there are anatomical connections between the macaque analogous of STS, IPL, and PF areas, and therefore a Hebbian learning rule can be applied. The particular network that has been used to simulate the imitation functionality is shown in Figure 3.

In this scheme the role of the STS neurons have the function to transfer the sensory (visual) stimuli and to account for the influence of the inhibitory neurons. For a robotic setting, modelling of the STS area can be reduced to the



Fig. 3. Core circuit for imitation

influence of the inhibitory neurons. Therefore, the sensory signals project directly to the IPL area which is associated with multisensory integration. The motor information, or the information from the movement of the wheels is co-activated in the simulated PMv area, which has sensorimotor integration functionality. The bidirectional projections between the two areas will insure that both areas represent the sensory and the motor signals.

The embodied implementation of this model is shown in Figure 4. The 8 range sensors of each robot project to the sensory integration area that resembles the functionality of the joined STS-IPL areas. The two wheels project to the sensorimotor integration area, which resembles the PMv, as shown in Figure 4.

Self-organization of rhythmic activity is a fundamental characteristic of biological systems. In addition, rhythmic activities are found in any level of the hierarchical structure, i.e. from the biochemical to the sociobiological level. At neuronal level, single neurons and networks respond with transient oscillations to strong input. The natural frequency, or eigenfrequency of the damped oscillation is a result of two opposing effects, often modelled by the combined effect of executory and inhibitory neurons.

We suggest to use entrainment of endogenous oscillators for modelling turn taking behaviour. The mirror neuron paradigm that allows the behaviour of each robot to be represented in the neuronal structures of its partner makes possible the oscillatory dynamics of the turn taking process to be modelled through the individual agents.

The mutual interaction between two robots has to emerge through selforganizing entrainment of oscillatory neurons. To check this hypothesis, the neurons of each robots mirror system are simulated as oscillators:

$$\theta(t) = \omega t \mod 2\pi \tag{1}$$

The above equation determines the change of rate of the phase with the time. is the the cycle of the limit cycle oscilation. The phase is periodic over the range.



Fig. 4. Robot architecture with mirror circuit

If a synaptic coupling H connects two neurons, their phase equations will be represented in the following way:

$$\frac{d\theta_1}{dt} = \omega_1 + H_1(\theta_2 - \theta_1) \tag{2}$$

$$\frac{d\theta_2}{dt} = \omega_2 + H_2(\theta_1 - \theta_2) \tag{3}$$

where indexes 1 abnd 2 refer to the first and the second neuron respectively. The learning rule for oscillatory networks was used, the original learning rule was shown in [4].

4 Experimental Setting and Results

The experimental setting consists of two parts. In the first part each robot has to build sensory-motor experiences by exploring an environment that consists of a circular arena inhabited by another robot, see Figure 5a. In the second part, interactive turn-taking behaviour emerges, based on the established oscillatory sensorimotor couplings.

Initially we constructed the experimental scenario that performs following behaviour for the training phase, see Figure 5a. Both robots consequently are taking the role of the follower, in order to establish adequate "mirroring" couplings between the lattices that resemble the IPL and PMv areas. Before training, IPL serves as multisensory integration area and PMv is primarily sensorimotor integration area. The hebbian connections between these lattices are modelled in the way that after training both areas will reflect the common sensorimotor representation that is the basis for interaction behaviour:

$$\Delta w_{lk}^{PM-IPL} = \alpha (PM_l - \overline{PM_l})(IPL_k - \overline{IPL_k})$$
(4)

where $\overline{PM_l}$ and $\overline{IPL_k}$ are the average activation values of units l and k over a certain time interval. IPL-PMv synaptic plasticity has the following dynamics:



Fig. 5. a) Scenario for following and turn taking behaviours based on the tag game. b) The shortest sensor measurement determines the relative position and direction of the partner robot for the following interaction.



Fig. 6. Movement imitation behaviours

the connection between them is strengthened if both of them are simultaneously active and weakened if the activation of one decreases.

In the initial experiment the robot-follower denotes its shortest distance reading, which signals the presence of the partner robot, as shown in Figure 5b. The placement of the distance sensors defines the relative heading of the partner robot. The robot-follower tends to synchronize its motion direction with the motion direction of the leading robot.

After neurons from the two lattices synchronize, the two simulated robots express a simple form of social behaviour. The leader robot performs movements with different complexity, and the follower (dashed lines) imitates it from its movement perspective, as shown in Figure 6.

At the second part of the experimental scenario, the emergent turn taking is to be shown. The role of the robot, being follower or leader at the present moment depends on which robot is 'within the visual field' of its partner. For the training phase, the tag game is simulated, by which the runner and the tagger functions change between the robots once the tagger reaches the runner.



Fig. 7. a) Neural activation during following behaviour. b) A typical desynchronization in the case of turn taking.

After training, the emergent turn taking has to take place which is expressed by symmetry breaking process after a period of synchronization. This way the leading robot can become a follower and later again the lead can be taken over by it. The turn taking, similar as by humans, takes place as a result of some subtle or explicit external stimulation. For the case of the tag game, the external stimulation is usually caused by losing the runner-robot from the perceptual field, caused by reaching the end of the arena or other reason for escape of the runner robot.

Figure 7 shows the neural activation during turn (the left plot) and in the period of turn taking (the plot on the right). The desynchronization of the neurons in central part of the right plot corresponds to the moment of losing the runner robot from the perceptual field. At that period previous follower changes its role to a runner, and vice versa.

5 Discussion

Social interaction has wide spectrum of expressions as synchronous movements, turn taking, gaze sharing, following, imitation and conversation. We have simulated simple interaction behaviours of following and turn taking. In the training phase, the simulated following and the simulated tag game helps to gather examples and establish the sensorimotor couplings between the two robots. In the test runs, there is not an external control that will cause the turn taking behaviour. The turn taking is caused by changing of synchronous firing of the oscillatory neurons. Although external events are initiating turn change, turn taking does not take place only by the same conditions as during the training - turn taking has emergent properties due to nonlinear oscillations and their interaction. This results resemble turn taking in speech: an upcoming end of a turn is anticipated by perceiving eye gaze, body movement, or other queues by the speaker, or indicated in a subtle manner by the listener. Important questions for designing a movement interactions lies in the respective computational role of each brain area that subserves the internal simulations and shared representations between self and others. We based our model on the simplified mirror neuron network, in which the mirroring functionality is obtained via the selforganization of synchronized neural firing in two robots that share perceptual space. The emergence is an important element, but better understanding of underlying processes and computations will increase the possibilities and reliability of the interaction behaviour. This work is in the process of extensive development.

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