



# Towards a neurocomputational model of speech production and perception

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## Abstract

The limitation in performance of current speech synthesis and speech recognition systems may result from the fact that these systems are not designed with respect to the human neural processes of speech production and perception. A neurocomputational model of speech production and perception is introduced which is organized with respect to human neural processes of speech production and perception. The production–perception model comprises an artificial computer-implemented vocal tract as a front-end module, which is capable of generating articulatory speech movements and acoustic speech signals. The structure of the production–perception model comprises motor and sensory processing pathways. Speech knowledge is collected during training stages which imitate early stages of speech acquisition. This knowledge is stored in artificial self-organizing maps. The current neurocomputational model is capable of producing and perceiving vowels, VC-, and CV-syllables (V = vowels and C = voiced plosives). Basic features of natural speech production and perception are predicted from this model in a straight forward way: Production of speech items is feedforward and feedback controlled and phoneme realizations vary within perceptually defined regions. Perception is less categorical in the case of vowels in comparison to consonants. Due to its human-like production–perception processing the model should be discussed as a basic module for more technical relevant approaches for high-quality speech synthesis and for high performance speech recognition.

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## 1. Introduction

Current speech recognition systems are easily outperformed in the case of (i) non-restricted vocabulary, (ii) if the speaker is not well-known by the system and (iii) if noise reduces the speech signal quality (e.g. Benzeghiba et al., 2007; Scharenborg, 2007). Current corpus-based speech synthesis systems are limited as well, especially concerning (i) flexibility in modeling different speaker and voice characteristics and concerning (ii) segmental as well as prosodic naturalness (e.g. Clark et al., 2007; Latorre et al., 2006). These limitations may be attributed to the fact that speech recogni-

tion as well as speech synthesis systems currently are not modeled with respect to the basic human neural processes of speech production and speech perception.

A variety of brain imaging studies clarify the role of different subcortical and cortical brain *regions* for speech production (e.g. Murphy et al., 1997; Kuriki et al., 1999; Wise et al., 1999; Bookheimer et al., 2000; Rosen et al., 2000; Scott et al., 2000; Benson et al., 2001; Huang et al., 2001; Blank et al., 2002; Vanlancker-Sidtis et al., 2003; Ackermann and Riecker, 2003; Hillis et al., 2004; Shuster and Lemieux, 2005; Kemeny et al., 2005; Riecker et al., 2006; Sörös et al., 2006) as well as for speech perception (e.g. Binder et al., 2000; Hickok and Poeppel, 2000; Fadiga et al., 2002; Wilson et al., 2004; Boatman, 2004; Poeppel et al., 2004; Rimol et al., 2005; Liebenthal et al., 2005; Uppenkamp et al., 2006; Zekveld et al., 2006; Obleser et al., 2006, 2007). Other studies focus on the interplay of speech

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production and perception (Heim et al., 2003; Okada and Hickok, 2006; Callan et al., 2006; Jardri et al., 2007) but only few among them introduce *functional* neural models which explain and emulate (i) the complex neural sensorimotor processes of speech production (Bailly, 1997; Guenther, 1994, 1995, 2006; Guenther et al., 2006) and (ii) the complex neural processes of speech perception including comprehension (McClelland and Elman, 1986; Gaskell and Marslen-Wilson, 1997; Luce et al., 2000; Grossberg, 2003; Norris et al., 2006; Hickok and Poeppel, 2004, 2007).

It is the aim of this paper to introduce a biologically motivated approach for speech recognition and synthesis, i.e. a computer-implemented neural model using artificial neural networks, capable of imitating human processes of speech production and speech perception. This production–perception model is based on neurophysiological and neuropsychological knowledge of speech processing (Kröger et al., 2008). The structure of the model and the process of collecting speech knowledge during speech acquisition training stages are described in detail in this paper. Furthermore it is described how the model is capable of producing vowels and CV-syllables and why the model is capable of perceiving vowels and consonants categorically.

## 2. The structure of the neurocomputational model

While the *structure* of this neurocomputational model is based on neurophysiological and neuropsychological facts (Kröger et al., 2008), the speech *knowledge* itself is gathered by training artificial neural networks which are part of this model (Kröger et al., 2006a,b). The organization of the model is given in Fig. 1. It comprises a cortical and a subcortical–peripheral part. The cortical part is subdivided with respect to neural processing within the frontal, the temporal, and the parietal cortical lobe. Functionally the model comprises a production and a perception part. In its current state the model excludes linguistic processing (mental grammar, mental lexicon, comprehension, conceptualization) but focuses on sensorimotor processes of speech production and on sublexical speech perception, i.e. sound and syllable identification and discrimination.

The *production part* is divided into feedforward and feedback control (see also Guenther, 2006). It starts with the phonemic representation of a speech item (speech sound, syllable, word, or utterance) and generates the appropriate time course of articulatory movements and the appropriate acoustic speech signal. The phonemic representation of a speech item is generated by higher level lin-

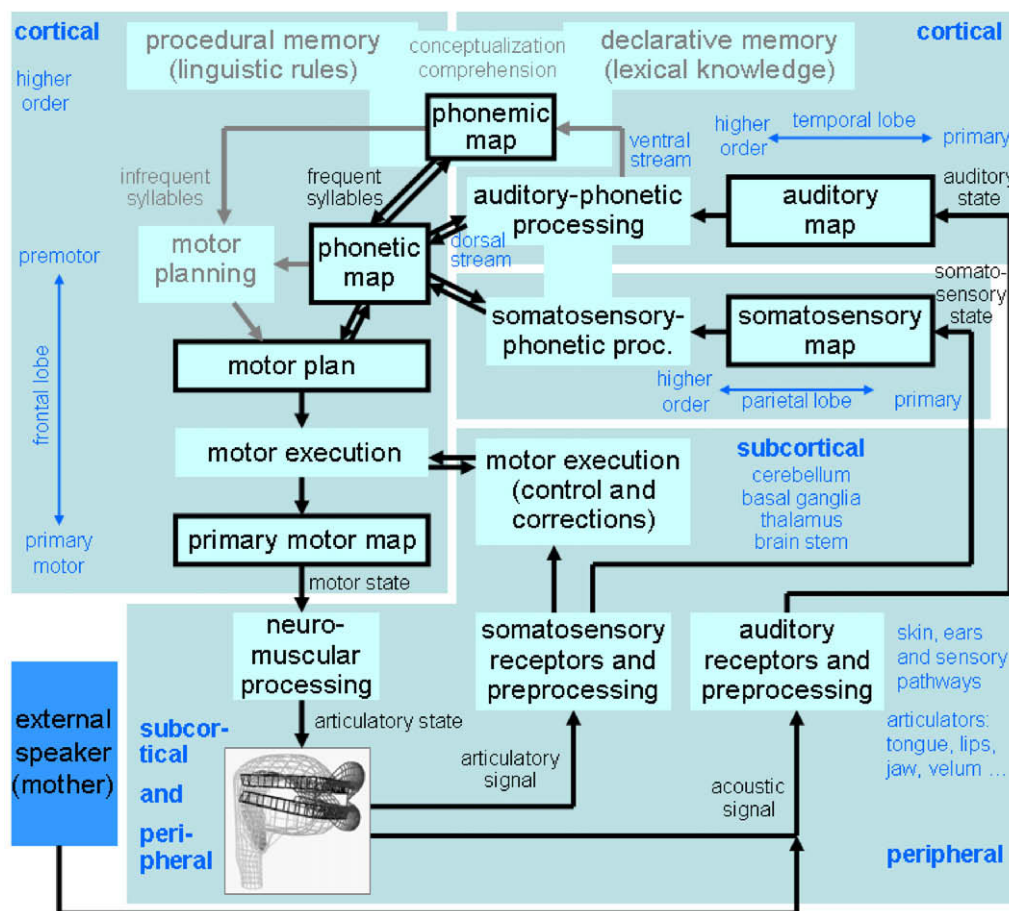


Fig. 1. Organization of the neurocomputational model. Boxes with black outline represent neural maps. Arrows indicate processing paths or neural mappings. Boxes without outline indicate processing modules. Grey letters and grey arrows indicate processing modules and neural mappings which are not computer-implemented in the current version of the model.

guistic modules (Levelt et al., 1999; Dell et al., 1999; Indefrey and Levelt, 2004) subsumed as widely distributed frontal–temporal procedural and declarative neural processing modules (Ullman, 2001; Indefrey and Levelt, 2004) which are not specified in detail in this model. Subsequently each phonologically specified syllable (i.e. a phonemic state; a neural activation pattern on the level of the phonemic map) is processed by the *feedforward control* module. In the case of a *frequent syllable*, the sensory states (auditory and somatosensory state) and the motor plan state of the syllable (which are already learned or trained during speech acquisition; see below) are activated via the phonetic map. The *phonetic map* (Fig. 1) can be interpreted as the central neural map constituting the mental syllabary (for the concept of mental syllabary, see Levelt and Wheeldon, 1994; Levelt et al., 1999). For each frequent syllable a phonemic state initiates the neural activation of a specific neuron within the phonetic map, which subsequently leads to activation patterns of the appropriate sensory states and the appropriate motor plan state. In the case of *infrequent syllables* the motor plan state is assembled within the motor planning module on the level of sub-syllabic units, e.g. syllable constituents like syllable onset and syllable rhyme or single speech sounds (Varley and Whiteside, 2001). This path is not implemented in our model at present. On the level of the *motor plan map* a high level motor state (motor plan) is activated for each speech item under production (current speech item). This high level motor state defines the temporal coordination of *speech gestures* or *vocal tract action units* (Goldstein et al., 2006; Saltzman and Munhall, 1989; for a general description of goal-directed action units, see Sober and Sabes, 2003; Todorov, 2004; Fadiga and Craighero, 2004). The motor plan of a speech item is processed by the motor execution module in order to define the spatio-temporal trajectories of articulator movements. Thus the motor execution module calculates the concrete specification of each speech gesture on the level of the *primary motor map* (cf. Ito et al., 2004; Sanguinetti et al., 1997; Saltzman, 1979; Saltzman and Munhall, 1989; Saltzman and Byrd, 2000). For example, a labial closing gesture involves coordinated movement of at least the lower jaw, the lower and upper lips. Thus each of these articulators must be controlled synergetically for the realization of a speech gesture. Subsequently the movement of an articulator is executed by activating the motor units controlling this articulator via the neuromuscular processing module.

The (lower level) primary motor map comprises 10 articulatory parameters (Kröger et al., 2006b). Each articulatory parameter value is coded by two neurons with complementary activation (see below) leading to 20 neurons to encoding the primary motor commands for each point in time. The conversion of physical parameter values (e.g. displacement of an articulator) into neuromotor activation patterns is done (i) by mapping the physical displacement range for each parameter onto a neural activation range  $[0, 1]$  (i.e. no activation to full activation of a neuron) and (ii) by defining two neurons for each

parameter with complementary activation ( $a_2 = 1 - a_1$ ) in order to hold the overall activation  $a$  ( $a = a_1 + a_2$ ) constant ( $=1$ ) for each parameter value. The size of the (higher level) motor plan map depends on the length of the utterance under production. In the case of V-, CV-, and VC-items three vocalic higher level parameters (high–low, front–back, rounded–unrounded) and four higher level consonantal parameters (labial, apical, dorsal, exact closing position) are controlled. These vocalic parameters and the consonantal parameter closing position are encoded using two neurons with complementary activation each, while the three remaining consonantal parameters are encoded by one neuron each in order to reflect the activation of a specific vocal tract organ. Thus the motor plan map for V-, CV-, and VC-items consists of 11 neurons. Since a motor plan encodes a motor or sensory V-, CV-, or VC-item of a transition for C (encoded by four time labels) and a steady state portion for V (encoded by one time label) the (lower level) primary motor state of these items is encoded by five consecutive time labels. Thus the appropriate number of primary motor map neurons for a whole speech item is  $5 \times 20 = 100$  neurons plus 10 neurons for coding five time intervals describing the temporal distance from label to label.

A computer-implemented numerical *articulatory vocal tract model* generates the time course of vocal tract geometries and subsequently the *acoustic vocal tract model* generates the acoustic speech signal. A three-dimensional articulatory–acoustic model is used here which is capable of generating high-quality articulatory and acoustic speech signals (Birkholz and Jackel, 2004; Birkholz and Kröger, 2006, 2007; Birkholz et al., 2006, 2007 and Kröger and Birkholz, 2007). These articulatory and acoustic signals are used for feedback control.

The articulatory and acoustic signals generated by feedforward control are continuously monitored or controlled. For this *feedback control* the articulatory and acoustic signals are converted into neural signals by auditory and somatosensory (i.e. tactile and proprioceptive) receptors. Somatosensory feedback signals (relative positions of articulators to each other and position and degree of vocal tract constrictions, see Saltzman and Munhall, 1989; Shadmehr and Mussa-Ivaldi, 1994; Tremblay et al., 2003; Nasir and Ostry, 2006) are used for controlling motor execution. In addition sensory (i.e. somatosensory and auditory) signals are converted into *higher level cortical sensory states*, which represent the current speech item. These auditory and somatosensory (feedback) states of a currently produced speech item are processed by comparing them with the appropriate prelearned auditory and somatosensory state, activated by feedforward control before the current speech item is produced. This comparison is done on the level of the somatosensory and auditory processing modules. If the prestored (or feedforward) sensory state and the feedback sensory states indicate a reasonable difference an error signal is activated for correcting the motor plan during the ongoing feedforward control.

The conversion of physical or psychophysical sensory parameter values (e.g. bark scaled formant values) into neural activation patterns is done (i) by mapping the whole physical parameter range onto the “neural” range  $[0, 1]$  (i.e. no activation to full activation of a neuron) and (ii) by defining two neurons per parameter with complementary activation (see above for the primary motor map). Since auditory states are processed as whole patterns, parameter values for our V-, CV-, and VC-items (see above) are obtained at five positions (labels) in the acoustic signal. Three formants were processed leading to  $3 \times 5 = 15$  parameter values and thus to 30 neurons per item for the auditory state map. 10 proprioceptive and nine tactile parameters were processed (Kröger et al., 2006b) leading to 19 parameter values and thus 28 neurons for each item. Only one tactile and proprioceptive state is coded for the whole speech item representing the gestural target region of the vocalic part in the case of a V-item and representing the gestural target regions of the vocalic and the consonantal part in the case of VC- or VC-items (overlay of tactile and proprioceptive patterns).

The *perception part* of the neurocomputational model starts from an acoustic speech signal, generated by an external speaker (Fig. 1). This signal is converted into neural signals by auditory receptors and is further processed into a cortical higher level auditory signal via the same auditory pathway that is used for the feedback control of speech production (self-productions). Speech perception comprises two pathways (cf. Hickok and Poeppel, 2004, 2007). The *auditory-to-meaning pathway* (*ventral stream*) directly activates neural states within the mental lexicon by the high level cortical auditory state for a speech item (e.g. a word). This pathway is not included in our model, since high level mental lexical representations are out of the scope of this study. The *auditory-to-motor pathway* (*dorsal stream*) activates the phonetic state of the current speech item (e.g. sound or syllable) within the cortical frontal motor regions. This pathway is included in our model and it will be shown below that this pathway is capable of modeling categorical perception of speech sounds and is capable of modeling differences in categorical perception of vowels and consonants.

The structure of the neurocomputational model differentiates neural maps and neural mappings. *Neural maps* are ensembles of neurons which represent the phonemic, phonetic, sensory or motor speech states. These maps are capable of carrying states of different speech items by different neural activation patterns. These activations change from speech item to speech item under production or perception. *Neural mappings* represent the neural connections between the neurons of neural maps (Fig. 2). These connections can be excitatory or inhibitory. The degree of excitatory or inhibitory connection is described by link weight values. These values  $w_{ij}$  characterize the neural connection between each pair of neurons. They define the degree of activation of a connected neuron  $b_j$  within a neural map 1 (comprising  $M$  neurons  $j = 1, \dots, M$ ) resulting from the degree of acti-

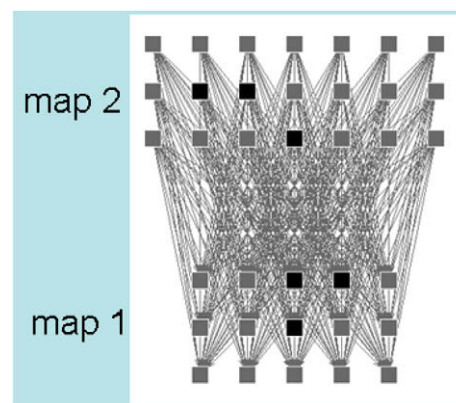


Fig. 2. One-layer feedforward network connecting two neural maps 1 and 2. Grey lines indicate the neural connections connecting each neuron of map 1 with each neuron of map 2.

vation of all connecting neurons  $a_i$  within a neural map 2 (comprising  $N$  neurons  $i = 1, \dots, N$ ) (see Eq. (1) and Fig. 2).

$$b_j = \text{actfunc} \left( \sum_{i=1}^N a_i w_{ij} \right) \quad \text{for } j = 1, \dots, M \quad (1)$$

Here *actfunc* is the activation function (a sigmoid function in the case of our modeling; see Zell, 2003) which represents the total activation of neuron  $b_j$  in map 1 as function of the sum of activations from all neurons  $i$  within map 2. The link weight values  $w_{ij}$  are limited to the interval  $[-1, +1]$  (i.e. maximal inhibitory to maximal excitatory link weight value).

The link weight values reflect the whole knowledge inherent in the training data and thus the knowledge gathered during the training procedures. Link weight values are adjusted during training stages, i.e. during speech acquisition stages (see below). They are allowed to be modified continuously in order to reflect new knowledge gained over life time.

One-layer feedforward networks (Fig. 2) are of limited power and are used in our model exclusively for calculating articulatory joint-coordinate parameters from articulatory tract-variable parameters (cf. Kröger et al., 2006c). In this paper we will focus on the central phonetic map and the multilateral co-activation of phonemic states, sensory states, and motor plan states via the phonetic map. This multilateral co-activation is achieved by using self-organizing maps or networks (Kohonen, 2001 and Fig. 3). Each neuron of the central self-organizing map (i.e. the phonetic map) represents a speech item. Different phonetic submaps (i.e. different parts within the phonetic map) are defined for each class of speech items, i.e. for vowels, for CV-, and for VC-syllables. Multilateral co-activation of phonemic, sensory, and motor plan states for a speech item via the phonetic map means that an activated neuron of the phonetic map (representing a currently perceived or produced speech item) leads to a co-activation of neural activation patterns within the phonemic, motor plan, or sensory side

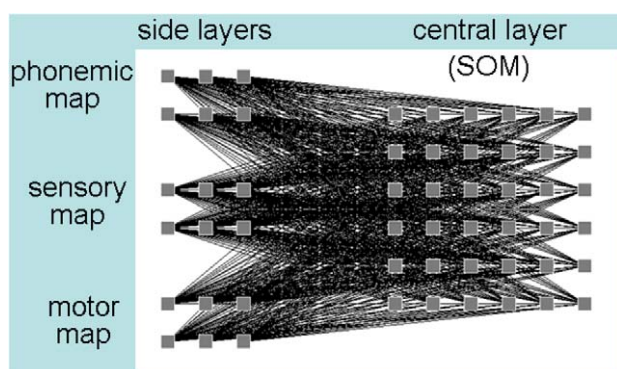


Fig. 3. Self-organizing network connecting three neural maps (side layer maps) by a central self-organizing map (SOM or central layer map). Black lines indicate the neural connections, connecting each neuron of each side layer map with each neuron of the central map.

layer maps representing this current speech item. The set of link weight values of the connections between all neurons of the phonemic, motor plan, or sensory side layer map and a neuron within the central phonetic map characterize the phonemic, motor plan, or sensory state of the speech item represented by this neuron within the phonetic map. Activation patterns of neurons within the side layer maps induced by an activation pattern of the phonetic map as well as activation patterns of the phonetic map induced by an activation pattern of one of the side layer maps are calculated in the same way as it is described above for simple one-layer feedforward networks (Eq. (1)).

The structure of the neurocomputational production–perception model introduced here is based on the structure of the DIVA model introduced by Guenther (2006) and by Guenther et al. (2006). The approach described in this paper as well as the Guenther approach comprise a feedforward and a feedback control path. Both approaches comprise self-organizing networks for processing neural states and comprise neural maps for storing phonemic, motor, and sensory states representing speech items. Both approaches introduce pre-linguistic and early linguistic language-specific training (i.e. babbling and imitation training, see below) in order to shape the neural mappings within the computational models and both approaches include the concept of a mental syllabary (Levelt and Wheeldon, 1994; Levelt et al., 1999) and basic ideas of the mirror neuron concept (Fadiga and Craighero, 2004; Rizzolatti and Craighero, 2004) since both approaches claim a simultaneously occurring activation of sensory and motor states for speech items.

But there are three major differences between both approaches. Firstly, the DIVA approach does not *separate motor planning and motor execution* as is introduced here. This separation results from the fact that for all types of voluntary movements (actions) just the goal of an action (e.g. grasping a definite object or pressing a sequence of buttons) and the temporal overlap or temporal sequencing of actions are determined on the planning level while the details of movement execution are determined on lower

neural levels (Kandel et al., 2000; Kröger et al., 2008). In the case of speech production *vocal tract action units* or *speech gestures* are well established as basic units of speech production (Browman and Goldstein, 1989, 1992; Goldstein et al., 2006, 2007) separating motor speech planning – i.e. the level of action scores (Goldstein et al., 2006) – and motor speech execution (Saltzman and Munhall, 1989; Goldstein et al., 2006) – i.e. the detailed determination of all articulator movements. The practical importance of dynamically defined speech action units becomes apparent if modeling of segmental reduction effects resulting from high speech rate (Kröger, 1993) or if modeling of speech errors (Goldstein et al., 2007) is attempted. Secondly, the DIVA model does not explicitly introduce a *phonetic map* or at least a map, reflecting the self-organization of speech items between sensory, motor, and phonemic representation; and the DIVA model does not explicitly claim *bidirectional mappings* between phonemic, sensory, and motor representations. But the assumption of bidirectional associations is essential in our production–perception model. Production is modeled in our approach using neural connections from the phonemic map directed towards the motor and sensory maps via the phonetic map and perception is modeled in our approach using the neural connections from sensory maps directed toward phonemic map via the phonetic map. Furthermore, the phonetic map itself is a central concept in our approach. On the one hand, the phonetic map introduces a *hypermodal description* of speech items which connects the sensory and motor representations of a speech item as is claimed in the mirror neuron theory. Our simulation results indicate that it is very feasible to introduce this level of neural self-organization (phonetic map) since it elucidates the ordering of speech items with respect to phonetic features (*phonotopy*, see below). Furthermore the notion of the phonetic map is important for modeling speech perception since perceptual discrimination is defined in our approach as a distance between activated states on this neural level (see below). Thirdly, the DIVA model is a production model not aiming for modeling *speech perception*. But according to the arguments given above the modeling of speech production and speech perception as two closely related processes is of great importance. This is achieved in our approach.

### 3. Gaining speech knowledge: training stages for speech acquisition

Speech knowledge is gained during training stages which model *basic stages of human speech acquisition*. This knowledge is stored within the mappings of the model, i.e. by the link weight values connecting the neurons within the production–perception model. Link weight values are adjusted during training stages. Two basic training stages can be differentiated, i.e. the babbling and the imitation stage (Oller et al., 1999).

For *babbling training* the training sets comprise pre-linguistic speech items, i.e. proto-vocalic and proto-syllabic

speech items. The model randomly produces proto-vocalic and proto-syllabic speech items and listens to its own productions using the auditory feedback path (Fig. 1). The link weights between the sensory maps and the motor plan map are adjusted via the phonetic map during this training stage. No phonemic activation occurs since these training items are pre-linguistic. The knowledge which is gained during babbling training is language independent *general phonetic knowledge*. During this training stage the neuro-computational model learns the *sensorimotor interrelationship* of the vocal tract apparatus and its neural control, i.e. the interrelationship between various motor plan states and their resulting somatosensory and auditory states. The babbling training can be subdivided into training stages for training of proto-vocalic states and for proto-syllabic CV- and VC-states.

The *proto-vocalic babbling training set* comprises a set of proto-vocalic states which exhibit a quasi-continuous variation of the vocalic motor plan parameters low–high and front–back. The training set used here comprises 1076 proto-vocalic states which cover the language independent articulatory vowel space between the cardinal vowel qualities [i], [a], and [u] (Fig. 4). Each proto-vocalic motor plan state is defined by the vocalic parameters back–front and low–high. Thus the proto-vocalic training stimuli form a two-dimensional plane within the F1–F2–F3 acoustic vowel space (Fig. 4). Other motor parameters like tongue body height, tongue body horizontal and vertical position, and like lip opening are functions of these two motor plan parameters (Kröger et al., 2006c). Even lip protrusion is a function of the parameter front–back in the case of this preliminary training set since the training set does not include rounded front proto-vowels (e.g. [y]-like vowel qualities).

The *proto-syllabic CV and VC babbling training sets* are based on a set of 31 proto-vocalic motor plan states which are characterized by the motor plan parameters back–front and low–high and which are covering the whole articulatory vowel space. Labial, apical, and dorsal opening and closing gestures (proto-CV- or proto-VC-gestures) starting or ending with a full closure are superimposed on these proto-vocalic states. Three proto-consonantal places of articulation (front–mid–back) are defined per gesture. This leads to a total amount of 279 training items for each of the two proto-syllabic training sets (proto-CV- and proto-VC-training set). The articulatory velocity of the gesture-executing articulator for all closing and opening gestures is proportional to the distance between actual articulator position and gestural target position. This leads to an exponential time function for the displacement of this articulator. A gesture is ending if the articulator-target distance is below 10% in the case of opening or proto-CV-gestures or if a full closure is reached in the case of closing or proto-VC-gestures (target of closing gestures is beyond the full closure position). In summary the motor plan state for these proto-CV- and proto-VC-gestures is defined by (i) two vocalic parameters (back–front and low–high), by (ii) the gesture-performing articulator (labial, apical, or dorsal)

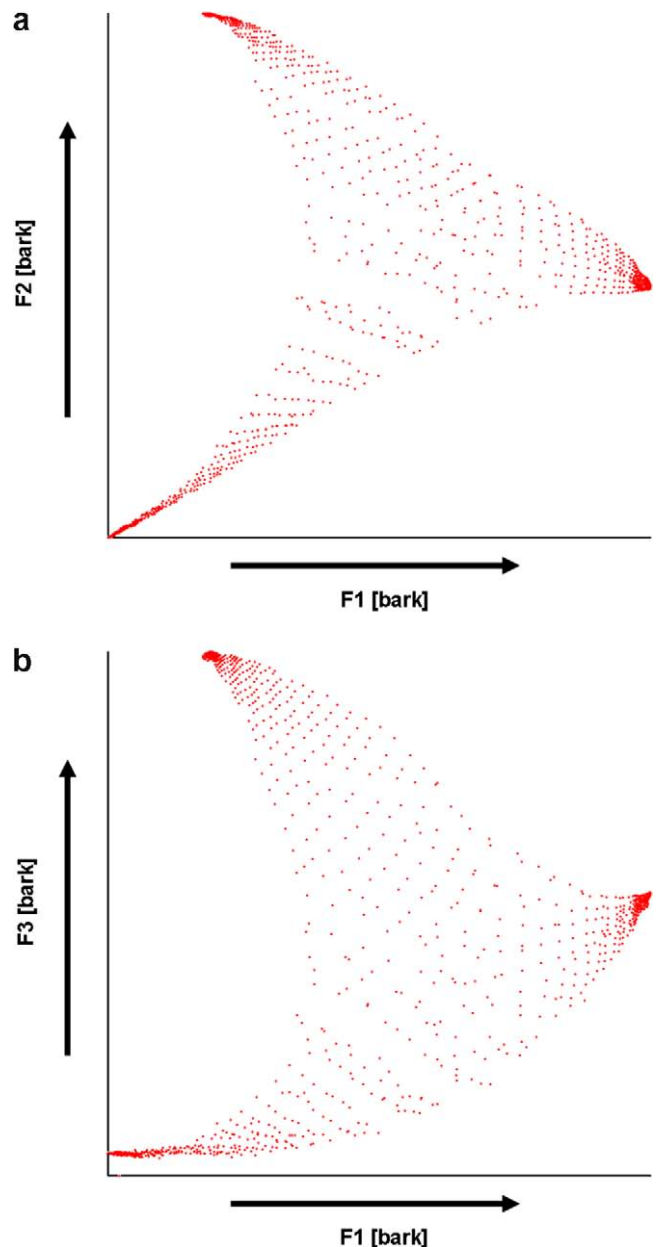


Fig. 4. Position of all auditory patterns of the proto-vocalic training stimuli (grey points) in the normalized and bark-scaled (a) F1–F2 and (b) F1–F3 vowel space.

and by (iii) the exact proto-consonantal closing position (front–mid–back). The lower level (or primary) motor parameters and their time courses are calculated from these motor plan parameters by the motor execution module. The appropriate auditory state of these opening and closing gestures (proto-CV- and proto-VC-syllables) is the time courses of the first three formants F1, F2, and F3 (Fig. 5 and see Section 2) and the appropriate somatosensory state for each proto-syllabic motor plan state comprises tactile information of the proto-consonantal closure and proprioceptive information of the proto-vocalic opening.

Proto-vocalic, proto-CV-syllabic, and proto-VC-syllabic babbling training is performed independently from each

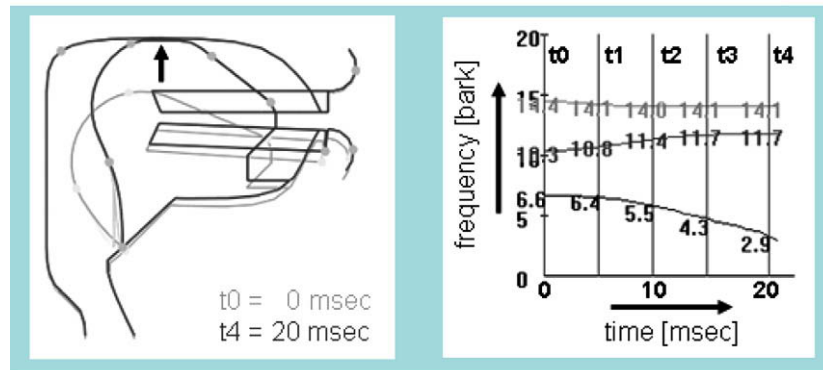


Fig. 5. Auditory state (right side) for a dorsal closing gesture (left side).

other. Three self-organizing maps (size:  $M = 15 \times 15 = 225$  neurons) form three phonetic submaps and are trained by using the three training sets described above. Training leads to an adjustment of link weight values  $w_{ij}$  between the  $N$  side layer neurons  $a_i$  and the  $M$  central layer neurons  $b_j$ . The side layers consist of the motor plan map ( $i = 1, \dots, K$ ) and the sensory (auditory and somatosensory) maps ( $i = K + 1, \dots, N$ ) while the central layer represents the phonetic map ( $j = 1, \dots, M$ ). Link weight values are initialized by random values within the interval  $[0, 1]$  (i.e. no activation to full activation). The link weights  $w_{ij}(t_{\text{init}})$  are initialized using random values between 0 and 1 (Eq. (2)). This adjustment of the link weights is done incrementally, i.e. step by step, using Hebbian learning (Eq. (3)). When a new stimulus  $I$  with  $I = (x_0, \dots, x_N)$  is presented, the winner neuron  $b_{\text{winner}}$  is identified in the central layer by calculating the minimum of Euclidian norm between  $I$  and  $W_j$ ,  $j = 1, \dots, M$ ; i.e.  $\text{winner} = \arg \min_j (\|I - W_j\|)$ , where  $W_j$  is a vector containing the link weights of all links from the central layer neuron  $b_j$  to the side layer neurons  $a_i$ , i.e.  $W_j = (w_{1j}, \dots, w_{Nj})$ . Once the winner neuron  $b_{\text{winner}}$  is identified the link weights for a step  $t$  with  $t_{\text{init}} < t < t_{\text{max}}$  are updated as

$$w_{ij}(t_{\text{init}}) = \text{rand}(0, 1) \quad (2)$$

$$w_{ij}(t + 1) = w_{ij}(t) + N_{\text{winner},j}(t) \cdot L(t) \cdot (I_i - w_{ij}(t)) \quad (3)$$

where  $0 < L(t) < 1$  is a constantly decreasing learning factor defined as

$$L(t) = 0.00001 + (L_{\text{init}} - 0.00001) \left(1 - \frac{t}{t_{\text{max}}}\right) \quad (4)$$

and  $N_{\text{winner},j}(t)$  is a neighborhood kernel (see Eq. (5)). Only the link weights of the neurons in the neighborhood around the winner neuron are updated. A 1-neighborhood is defined as all 8 neurons around the winner neuron, if they exist. A  $(n + 1)$ -neighborhood contains all neurons of a  $n$ -neighborhood and their 1-neighbors, if they exist. Thus a neighborhood kernel  $N_{\text{winner},j}(t)$  is defined as

$$N_{\text{winner},j}(t) = \begin{cases} 1 & \text{if } b_j \in r(t)\text{-neighborhood} \\ 0 & \text{if } b_j \notin r(t)\text{-neighborhood} \end{cases} \quad (5)$$

with neighborhood radius of  $b_{\text{winner}}$ . The additional step dependent function  $r(t)$  is introduced to get a constantly decreasing neighborhood radius (see Eq. (6)).

$$r(t) = 1.0 + (r_{\text{init}} - 1.0) \left(1 - \frac{t}{t_{\text{max}}}\right) \quad (6)$$

For the babbling training an initial neighborhood radius  $r_{\text{init}} = 12$  and an initial learning rate  $L_{\text{init}} = 0.8$  are chosen.

Proto-vocalic and proto-syllabic test sets were defined for testing the proto-vocalic and proto-syllabic training results. The proto-vocalic test set comprises 270 proto-vocalic states which cover the language independent articulatory vowel space between the cardinal vowel qualities [i], [a], and [u]. This proto-vocalic test set is organized in the same way as the proto-vocalic training set but the test set exhibits a much lower density within the articulatory or auditory vowel space. This also results in different training and test items. Both proto-syllabic test sets are based on a set of 22 quasi-vocalic motor plan states covering the whole language independent articulatory vowel space. Both proto-syllabic test sets are organized in the same way as the proto-syllabic training sets but the test sets exhibit a lower density within the articulatory or auditory vowel space for the proto-vocalic starting or ending positions of the VC- or CV- proto-syllables. Both proto-syllabic test sets comprise 198 items. The test items were different from the training items defined above.

An estimation of the quality of the proto-vocalic and the proto-syllabic training results is done by calculating a mean error over all test set items for estimating an articulatory state of a test set item from its auditory state. The calculation of the error value for each test item comprises six steps: In a first step the motor plan state of a test item is applied to the motor execution module for calculating the appropriate articulatory patterns (i.e. the time course of articulatory parameters for a speech item) by using the feedforward part of the model. This calculated articulatory pattern is called *initial articulatory pattern*. In a second step the appropriate auditory state pattern is calculated by using the output of the three-dimensional articulatory-acoustic model for the initial articulatory pattern and by

applying this output to the auditory feedback pathway of the model. In a third step the motor plan state is recalculated from the auditory state pattern calculated in the second step. Note that the trained self-organizing network is used for this step. This step leads to an *estimated motor plan state* which results from the sensorimotor knowledge stored within the self-organizing network, i.e. which results from the learning or training procedure. In a fourth step the *estimated articulatory pattern* is calculated for the estimated motor plan states by reusing the feedforward part of the model. In a fifth step the estimated and initial articulatory patterns are compared. An error value is calculated for each test item which is the difference between estimated and initial articulatory pattern. This difference is normalized with respect to the initial articulatory pattern. In a sixth step the mean error over all test set items is calculated for the trained network.

500,000 training steps are sufficient for predicting associated articulatory states from the auditory states of the test items with a precision below 2% error rate on the primary motor level in the case of the proto-vocalic training (using the proto-vocalic training set) and 280,000 training steps are sufficient for predicting the articulatory states from the auditory states with a precision below 5% error rate in the case of both proto-syllabic trainings (using both proto-syllabic training sets). Thus the complete babbling training requires less than five minutes on standard PC's.

The resulting link weight values for the neurons connecting the self-organizing phonetic maps with the motor plan and auditory map are graphically displayed for the

proto-vocalic training in Fig. 6 and for the proto-CV-syllabic training in Fig. 7. It appears that motor plan states are organized with respect to phonetic categories. In the case of the vocalic phonetic submap vocalic states are ordered continuously with respect to the motor plan parameters high–low and front–back. Experimental evidence for this kind of ordering is given by Obleser et al. (2006). In the case of the syllabic submap three regions occur which represent the gesture-performing articulator (labial, apical, and dorsal), i.e. an ordering occurs with respect to the motor-plan parameter gesture-performing articulator. This neural behavior resulting from self-organization of vocalic and consonantal or syllabic states with respect to phonetic categories (high–low, front–back, gesture-performing articulator) can be labeled as *phonetotopy* in parallel to tonotopy for the cortical ordering of auditory states with respect to their fundamental frequency (Kandel et al., 2000, p. 609) or in parallel to somatotopy for the ordering of somatosensory states with respect to their location on the body surface (Kandel et al., 2000, p. 460f).

It should be kept in mind at this point that the general phonetic sensorimotor knowledge stored in these phonetic maps is knowledge of sensorimotor relations exclusively generated by the three-dimensional articulatory and acoustic vocal tract model. Thus it is important for the performance or quality of neurocomputational models of speech production and perception that these models comprise realistic articulatory and acoustic vocal tract models as front–end modules which are capable of generating high-quality articulatory and acoustic signals, since the sig-

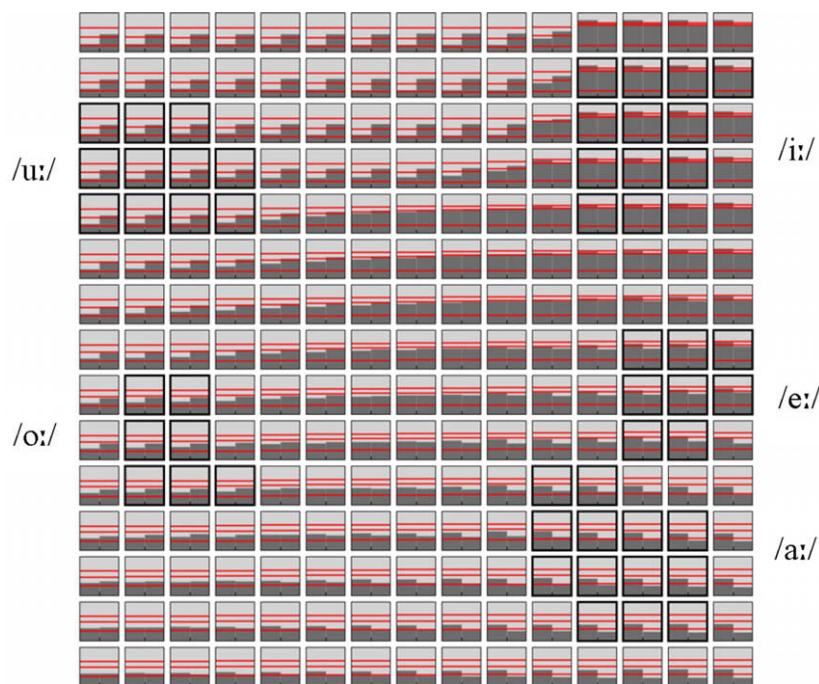


Fig. 6. Motor plan and auditory link weight values after vocalic babbling and imitation training for each neuron within the vocalic phonetic map ( $15 \times 15$  neurons). Link weight values are given for two motor plan parameters within each neuron box: back–front (left bar) and low–high (right bar). Link weight values are given for three auditory parameters: bark scaled F1, F2, and F3 (horizontal lines within each neuron box). The outlined boxes indicate the association of neurons with vowel phoneme categories. These associations are established during imitation training (see text).



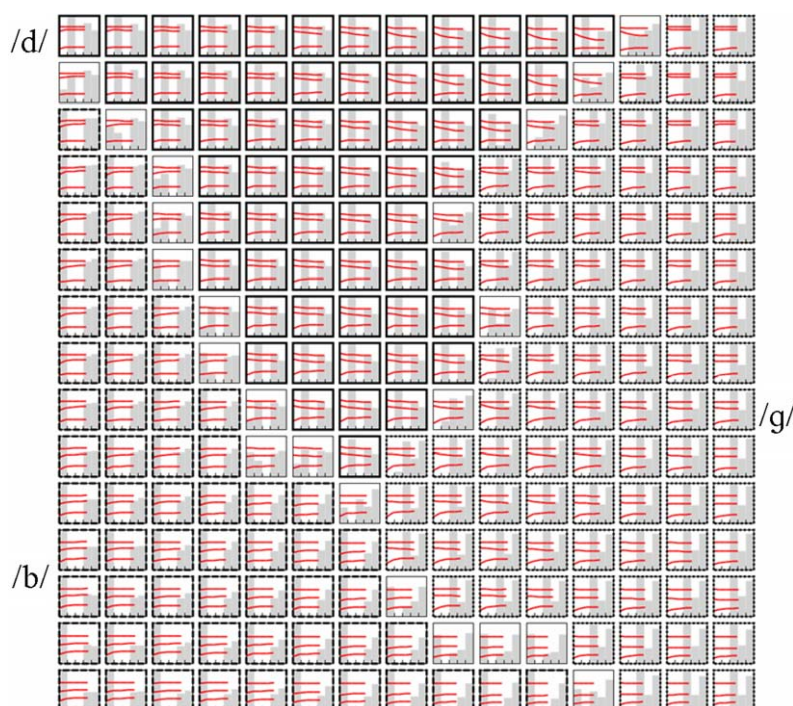


Fig. 7. Motor plan and auditory link weight values after CV-syllabic babbling and imitation training for each neuron within the CV-phonetic map ( $15 \times 15$  neurons). Link weight values are given for five motor plan parameters within each neuron box. First three columns: vocal tract organ which performs the closing gesture (labial, apical, dorsal); two last columns: back–front value (fourth column) and low–high value (fifth column) of the vowel within the CV-sequence. Link weight values are given for three auditory parameters: bark scaled F1, F2, and F3 (formant transitions within each neuron box). The outlined boxes indicate the association of neurons with consonant phoneme categories /b/, /d/, and /g/; each of these three regions comprises the appropriate consonant in all vocalic contexts. These associations are established during imitation training (see text).

nals generated by the articulatory–acoustic model are the basis for the calculation of all sensory signals.

After babbling training the neurocomputational model is capable of reproducing (or imitating) the motor plan state (i.e. the articulation) of any pre-linguistic speech item – in our case of any proto-vowel, proto-CV-syllable and proto-VC-syllable (with C = proto-consonantal closing gestures) – from their acoustic (or auditory) state patterns. Thus the neurocomputational model is now ready for language-specific *imitation training*. For imitation training the training sets comprise *language-specific* speech items; in our case vocalic and syllabic speech items. Beside the adjustment of link weights of the mapping between the phonetic map and the sensory maps and of the mapping between the phonetic map and the motor plan map, which is mainly done during babbling training, now in addition the link weights of the mapping between the phonetic map and the phonemic map are adjusted. Language-specific imitation training results in (i) specifying *regions of typical phoneme realizations (phone regions)* within the phonetic map, i.e. in specifying regions of neurons within the phonetic map, which represent typical realizations of a phoneme or of a syllable phoneme chain (see Figs. 6 and 7) and in (ii) fine-tuning of the sensorimotor link weights already trained during babbling. This fine-tuning mainly occurs at the phone regions. Thus the knowledge which is gained during imitation is language dependent. In other words

during this training stage the neurocomputational model mainly learns to link neurons which represent different phonemes or phonemic descriptions of syllables with the motor plan states and with the sensory states of their appropriate typical realizations. In parallel to babbling training also imitation training can be subdivided into training procedures for vowels, CV- and for VC-syllables.

The *vowel imitation training set* comprises a set of 100 acoustic vowel realizations per phoneme for a typical five vowel phoneme system /i/, /e/, /a/, /o/, and /u/ (e.g. Bradlow, 1995 and Cervera et al., 2001). A three-dimensional Gaussian distribution was chosen for each phoneme for distributing the 100 realizations per phoneme over the F1–F2–F3-space (Fig. 8 for the F1–F2-space). The distribution of the phoneme realizations in the acoustic vowel space (F1–F2–F3-space) is chosen as realistically as possible. The acoustic vowel realizations within the acoustic vowel space slightly overlap. These 500 vowel realizations are supposed to be realizations given by different external speakers, but matched with respect to the models babbling vowel space. It should be noted that vowel phonemes normally are learned in the context of words during speech acquisition. This is replaced in this model by training of isolated vowels by reason of simplicity. More complex training scenarios are beyond the scope of this paper.

During vowel imitation training each external acoustic (or auditory) vowel item is processed by the proto-vocalic

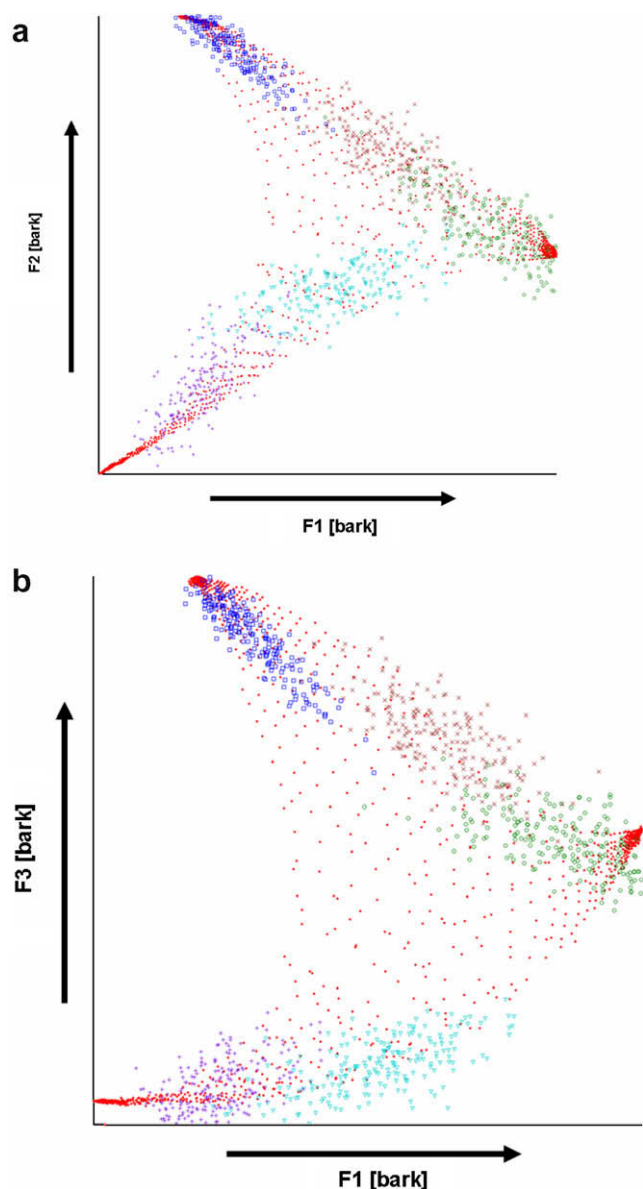


Fig. 8. Positions of all auditory patterns of the language-specific vocalic training stimuli (phone clouds: 100 realizations per phoneme /i/ (square), /e/ (cross), /a/ (circle), /o/ (triangle), and /u/ (plus)) in the normalized and bark-scaled (a) F1–F2 and (b) F1–F3 vowel space. The patterns (or phone clouds) are added to the proto-vocalic training stimuli (points).

babbling network in order to estimate the appropriate motor plan parameters. Thus the model is capable of re-articulating (imitating) these externally produced vowels and the model is capable of generating the appropriate internal auditory feedback states. In natural human speech acquisition scenarios the imitated vowel item is then judged as right or wrong (i.e. is accepted or not accepted) by an external listener; i.e. the produced item is awarded or not by the external listener, e.g. by communication between carer and toddler. If the item is accepted as a proper realization of the intended phoneme, its motor and sensory states can be linked to the neuron representing this phoneme in the phonemic map. In the case of our model all

realizations (re-articulations or imitations) can be accepted and thus can be added to the imitation training data set since the acoustic realizations of all re-articulations (or imitations) occur within the phoneme realization clouds (Fig. 8). Thus the vocalic imitation training set comprises 500 items of appropriate phonemic, motor plan, and sensory states. These data are the basis for imitation training.

The *syllable CV and VC imitation training sets* are based on a set of a labial, apical, and dorsal closing and opening gesture ending or starting at 31 different vowel realizations per vowel phoneme. That leads to 31 acoustic realizations for each of the phonemic CV- or VC-syllables (i.e. /bi/, /di/, /gi/, /be/, /de/, /ge/, /ba/, /da/, /ga/, /bo/, /do/, /go/, /bu/, /du/, and /gu/) and results in 465 training items. Each of these externally produced acoustic items are imitated in the same way as described above for the vowel items. Thus 465 training items of appropriate phonemic, motor plan, and sensory states for CV- or VC-stimuli are generated.

Only 5,000 training steps for vowels and only 5,000 training steps for CV- and VC-syllables had to be added to the proto-vocalic and proto-syllabic babbling training for obtaining clear *phoneme realization regions (phone regions)* within the phonetic maps (see the outlined neuron boxes in Figs. 6 and 7). A neuron of the phonetic map is defined to be a part of a phone region if the phonemic link weight value for this neuron of the phonetic map and the appropriate neuron of the phonemic map is above the level of 0.95. Thus for the neurons which form a phone region within the phonetic map, strong excitatory connections exist towards the neuron representing the appropriate phoneme within the phonemic map.

For imitation training the imitation training sets are used in addition to the ongoing applied babbling training set. The network is not reset; the already trained babbling network is used as a basis for further training. The algorithms for adjusting the network link weight are identical for babbling and for imitation training. Thus the succession from babbling to imitation training needs not to be abrupt. Imitation training can start in parallel to babbling training if some sensorimotor knowledge, i.e. if some knowledge how to estimate motor plan states from auditory states, is already available from early babbling training. The complete imitation training requires less than one minute on a standard PC for the training sets used here.

#### 4. Producing and perceiving vowels and CV-syllables

It should be emphasized that babbling and imitation training is not only the basis for learning to *produce* speech items of a target language. Since the sensory states of all self-productions are perceived by the feedback loop during babbling training and since external acoustic speech items as well as self-productions are perceived during imitation training it can be hypothesized that babbling and imitation training are also important for learning to *perceive* speech items of a target language.

The *production pathway* (phonemic map → phonetic map → motor plan map → primary motor map → articulation) has been introduced in Section 2. The speech items which were trained in this study can be labeled as frequent syllables. The description of the processing of infrequent syllables is beyond the scope of this paper. Our training results given above indicate strong neural connections from a phonemic state within the phonemic map to a set of neurons within the phonetic map. Each of these sets of neurons within the phonetic map represent a region of phoneme realizations (phone regions) and thus represent *production variability* since neighboring neurons within the phonetic map represent slightly different motor and sensory states (for natural variation in vowel realizations, see Perkell et al., 1993). If a phonemic speech item is activated (phonemic map) this leads to an activation of *several* neurons within the phonetic map (see the outlined boxes or phone regions for example for the vocalic phonetic map; Fig. 6). Thus in our model the maximal activated neuron within the phonetic map can differ from realization to realization. Therefore the motor plan and the subsequent articulatory realization of a phonemic item are allowed to vary within a perceptually acceptable region. These regions for phonemic items are the phoneme realization regions or phone regions and they are language-specific and are defined during imitation training (see Figs. 6 and 7).

Furthermore *coarticulation* is introduced in our neurocomputational model. Two sources of coarticulation are implemented in our model. Firstly, coarticulation results from the fact that the exact coordination of articulators for executing a speech gesture is controlled by the motor execution module and that a speech gesture is not encoded in all details on the motor plan level. That leads to variability in gesture execution with respect to context. For example the realization of /b/ in /ibi/ or /aba/ is different in our model. In /aba/ the lower jaw is more involved in the execution of the labial closing gesture than in /ibi/ because of the wide mouth opening occurring in /a/ in comparison to /i/. Because of this wide mouth opening in /a/ it would be ineffective to execute the closing gesture in /aba/ just by using the lips. It is more effective to add a synergetic elevation of the lower jaw. Thus, the lower jaw elevation and the lower lip elevation form a labial closing gesture in a *synergetic* way. Secondly, coarticulation results from the fact that gesture specifications can vary even on the level of the motor plan. For example lip protrusion is allowed to vary for a consonantal labial closing gesture since lip protrusion is a *non-relevant phonemic feature* in the case of a labial closing gesture in our target language. Since the labial closing gesture within a CV-syllable temporarily overlaps with the following vocalic gesture (e.g. for a gesture for realizing an /i/ or /u/) our simulations show anticipatory lip protrusion on the motor execution level in /pu/ while lips are not protruded during the labial closure in /pi/.

In the case of language-specific perception of speech items it can easily be shown that the neurocomputational

model trained thus far for vowels and simple CV- and VC-syllables is capable of producing *categorical perception* for vowels and in an even stronger way for consonants (i.e. voiced plosives in the case of our model). The *auditory pathway* for perception of external speech items (auditory receptors → auditory map → phonetic map → phonemic map) has already been introduced in Section 2 (auditory-to-motor pathway, see Hickok and Poeppel, 2000, 2004). Thus the phonetic map is not only a central neural representation in speech production but also in speech perception at least for sublexical speech units like speech sounds and syllables. In order to show that the current neuron-computational production–perception model perceives vowels (for the five vowel system /i/, /e/, /a/, /o/, and /u/) and consonants (for the voiced plosives /b/, /d/, and /g/) in a speech-like categorical way, speech identification and discrimination experiments were carried out using the model. In order to be able to perform these experiments using the model, 20 different instances of the model were trained using (i) different sets of training data due to different randomization procedures for determining the vocalic items within all training sets, using (ii) a different ordering of training stimuli during each training stage, and using (iii) different sets of randomly generated initial link weight values for each of the 20 instances. The resulting 20 instances of the model are called *virtual listeners*.

Identification of an external acoustic stimulus is performed in our model by a virtual listener by identifying the most excited neuron within the phonemic map. Discrimination of two external acoustic stimuli is performed in our model by calculating the most activated neuron on the level of the phonetic map for each acoustic stimulus and subsequently by calculating the city block distance between these both neurons for each virtual listener. The phonotopic ordering of speech items on the level of the phonetic map (see above) is a first hint that distance between speech items (states) on the level of this map indicates phonetic similarity or dissimilarity. Moreover we assume that the sensory resolution of two states (i.e. the capability for discrimination between these states) is governed by the spatial distance of these two states on the level of the phonetic map. This assumption holds for tonotopic ordering and thus for F0-discrimination of auditory stimuli (see the discussion of tonotopic cortical maps, Kandel et al., 2000, p. 609) and this assumption also holds for somatotopic ordering and thus for the spatial discrimination of tactile stimuli (see the discussion of somatotopic maps, Kandel et al., 2000, p. 460ff). Consequently it can be hypothesized that two stimuli can be discriminated if the distance of the activated neurons representing the stimuli on the level of the phonetic map exceeds a certain neuron distance within this map and it can be hypothesized that discrimination becomes stronger with increasing neuron distance.

Vocalic and consonantal identification and discrimination tests were performed on the basis of *quasi-continuous acoustic stimulus continua* (for an introduction to speech

perception experiments, see e.g. Raphael et al. (2007)). The stimulus continua generated for these tests model an /i/-/e/-/a/-continuum for vowels and a /ba/-/da/-/ga/-continuum for CV-syllables (Figs. 9 and 10). The resulting identification and discrimination scores are given in Figs. 11 and 12. It can be seen that the measured identification scores (measured for the 20 virtual listeners by identifying the most excited neuron within the phonemic map via the phonetic map for each stimulus) indicate more abrupt phoneme boundaries in the case of consonants than in the case of the vowels. Additionally it can be seen that the measured discrimination scores (measured for the same 20 virtual listeners by estimating the distance for both stimuli on the level of the phonetic map; see above) indicate higher discrimination scores at least for consonant perception. Beside *measured discrimination* (naturally perceived discrimination) also *calculated discrimination* scores are shown in Figs. 11 and 12. Calculated discrimination scores are theoretical constructs (see Liberman et al., 1957). They are calculated from (measured) identification scores for each single (virtual) listener. Thus calculated discrimination is a discrimination of stimuli which merely results from differences in identification of these stimuli. The probability  $p_{discr}$  for a certain percentage of calculated discrimination of two stimuli  $a$  and  $b$  is based just on the identification probabilities  $p_{id}$  of these two stimuli for each phonemic category  $i = 1, 2,$  or  $3$  (with  $1 = /b/$ ,  $2 = /d/$ , and  $3 = /g/$  in case of consonants and with  $1 = /i/$ ,  $2 = /e/$ , and  $3 = /a/$  in the case of vowels, see Eq. (7) and Liberman et al., 1957, p. 363).

$$p_{discr} = 0.5 + 0.5 \cdot \sum_{i=1}^3 (p_{id}(a, i) - p_{id}(b, i))^2 \quad (7)$$

Consequently calculated discrimination just indicates that part of discrimination of stimuli which results from the ability of subjects to classify stimuli to different categories. Calculated discrimination or *discrimination based on identification* (Liberman et al., 1957; Eimas, 1963) and its difference to (naturally) measured discrimination is discussed as an important feature of categorical perception (Damer

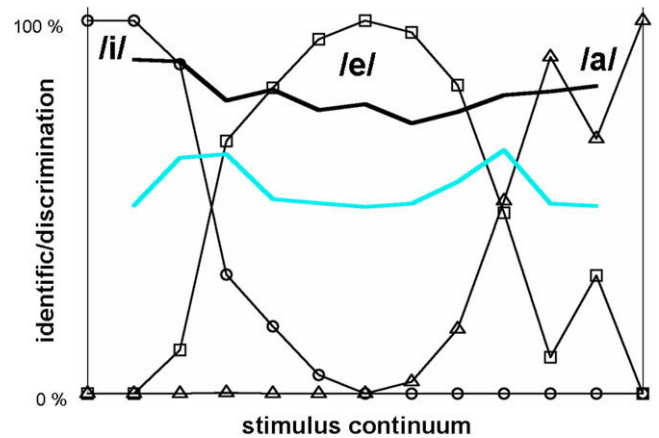


Fig. 11. Measured identification scores (non-bold black lines) and measured (bold black line) and calculated (bold grey line) discrimination score for the vocalic /i/-/e/-/a/ stimulus continuum for 20 virtual instances of the model.

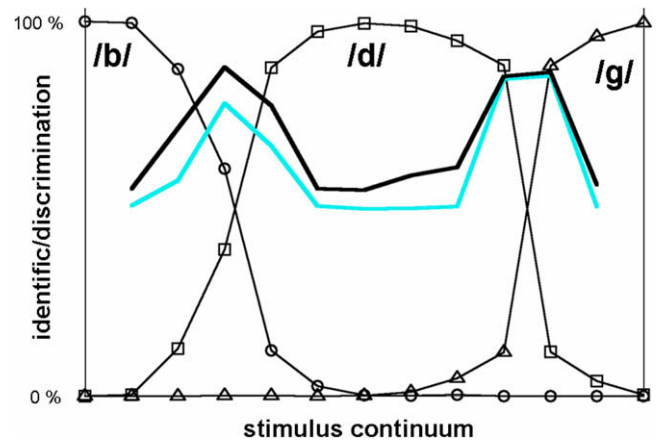


Fig. 12. Measured identification scores (non-bold black lines) and measured (bold black line) and calculated (bold grey line) discrimination scores for the consonantal /ba/-/da/-/ga/ stimulus continuum for 20 virtual instances of the model.

and Harnad, 2000). Calculated discrimination indicates discrimination which is just based on discrete linguistic or phonemic categorical knowledge, while measured discrimi-

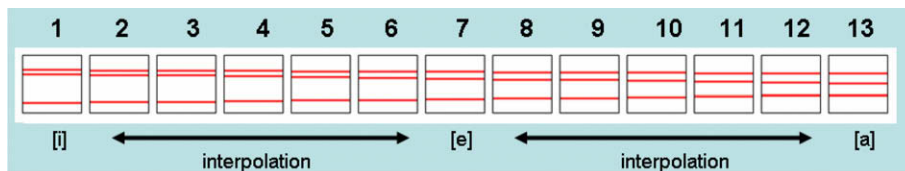


Fig. 9. Bark-scaled formant pattern for 13 vocalic stimuli (/i/-/e/-/a/-continuum) for the vocalic perceptual identification and discrimination tests.

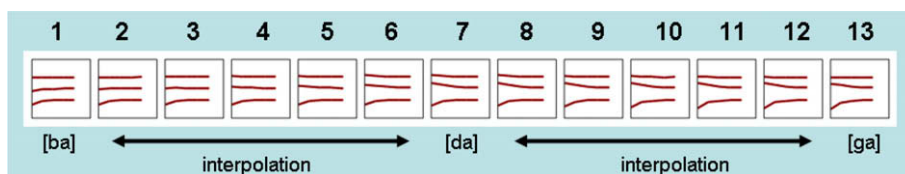


Fig. 10. Bark-scaled formant pattern for 13 CV-stimuli (/ba/-/da/-/ga/-continuum) for the consonantal perceptual identification and discrimination tests.

nation scores indicate the *complete* discrimination of two stimuli based on all available auditory information given by these stimuli; not just the linguistic, phonemic, or categorical information, needed for (categorical) identification. It can be seen from Figs. 11 and 12 that measured discrimination rates are always higher than calculated discrimination rates. That is in agreement with identification and discrimination scores extracted from identification and discrimination experiments carried out with humans and can be interpreted in the way that acoustic speech stimuli always convey categorical (linguistic) and non-categorical (para-linguistic or non-linguistic extra) information. While measured and calculated discrimination scores are nearly identical in the case of consonants, it comes out from our modeling data that measured discrimination is better than calculated discrimination especially in the case of vowels. This is in agreement with result of natural speech perception (Fry et al., 1962; Eimas, 1963) and reflects the typical differences in categorical perception of consonants and vowels.

## 5. Discussion and conclusions

The experimental results presented in this paper indicate that a model of speech production and perception which is shaped with respect to *basic neurophysiological facts* is capable of embedding important features of speech production and speech perception in a straight forward way even if the neurocomputational modeling is relatively basic as is here by using simple standard self-organizing networks. Typical and therefore important features of speech production and perception like production variability of phoneme realizations and categorical speech perception and especially the fact of different degrees of categorical perception for consonants and vowels, occur in a straight-forward way in this production–perception model. Since human speech production and perception easily outperforms speech synthesis and speech recognition systems at least in difficult conditions, it could be useful to include *human-like* speech processing routines into such technical speech processing systems. This may help to increase the quality and the level of performance of technical speech processing systems.

Furthermore this modeling study indicates the close relationship of speech *production* and speech *perception*. Speech perception theories such as the motor theory of speech perception (Lieberman et al., 1967; Liberman and Mattingly, 1985) or the direct-realist theory (Fowler, 1986) have already postulated this close relationship. And recent experimental results provide support for this claim and suggest that the development of an integrative model on speech production and perception is highly desirable. For example perceptual feedback loops (also called self-monitoring processes) are known to activate parts of the speech perception mechanism during overt (external perceptual loop) as well as covert speech production (internal perceptual loop, cf. Indefrey and Levelt, 2004; Postma,

2000; Hartsuiker and Kolk, 2001). In addition imaging studies focusing on speech perception have demonstrated that perception is capable of activating parts of the speech production cortical networks (Fadiga et al., 2002; Wilson et al., 2004; Hickok and Poeppel, 2004, 2007).

*Bidirectional* mappings between phonemic and phonetic and between sensory and phonetic maps are introduced in our neural model in order to illustrate the close relationship between production and perception. The introduction of these bidirectional mappings is the basis for important features of the model like categorical perception. Physiologically a bidirectional mapping comprises two related unidirectional mappings since neurons always forward their firing pulses in one direction (Kandel et al., 2000). Thus physiologically bidirectional mappings are represented by two neural paths connecting the maps in both directions (see the separate arrows in Fig. 1). The phonetic map – which forms the central map for all bidirectional mappings in our model (see Fig. 1) can be interpreted as the central part of the mental syllabary (Levelt and Wheeldon, 1994; Levelt et al., 1999). Neural cortico-cortical connections exist in both directions between this part of the frontal cortex and the sensory areas as well as between this part of the frontal cortex and those temporal regions which process phonemic information (Kandel et al., 2000).

Other computer-implemented models of speech production (Bailly, 1997; Guenther, 1994, 1995, 2006; Guenther et al., 2006) as well as the model introduced here reflect the relationship between perception and production by incorporating *perceptual feedback control loops* or by incorporating production–perception pathways for self-monitoring processes (Indefrey and Levelt, 2004). *Dual stream models of speech perception* have recently been published which introduce a ventral stream for passive auditory processing and a dorsal stream activating auditory-motor networks (e.g. Hickok and Poeppel, 2004, 2007) but passive models of speech perception that do not refer to production processes can also be found (McClelland and Elman, 1986; Gaskell and Marslen-Wilson, 1997; Luce et al., 2000; Norris et al., 2006). The model introduced here reflects the close relationship between speech production and speech perception since on the one hand our model comprises basic features of speech production models (cf. Guenther et al., 2006) and since on the other hand our model is capable of incorporating in addition the dual stream idea (Hickok and Poeppel, 2007) in a straight forward way (see the labels “ventral stream” and “dorsal stream” in Fig. 1).

*Mirror neurons* (visual and audio–visual mirror neuron system) appear to be one of the neural systems that are involved in the association of production and perception processes (Rizzolatti and Arbib, 1998; Studdert-Kennedy, 2002; Kohler et al., 2002; Fadiga and Craighero, 2004; Rizzolatti and Craighero, 2004; Wilson et al., 2004; Iacoboni, 2005; Wilson and Knoblich, 2005; Arbib, 2005). Systems of mirror neurons have been detected which code the abstract

meaning of goal-directed actions (e.g. grasping) and which are capable of co-activating motor *and* sensory (visual and audio–visual) representations of these actions by neural cortico-cortical associations. These visual and audio–visual mirror neuron systems also co-activate abstract concepts (preferably for action words) and thus are capable of associating higher order linguistic representations with goal-directed actions. A *speech* mirror neuron system (“mirror resonant system” after Fadiga and Craighero, 2004, p. 167, “auditory mirror neuron system” or “echo neurons” after Rizzolatti and Craighero, 2004, p. 185f) is postulated which is newer from the viewpoint of evolution in comparison to the mirror neuron system introduced above and which is directly linked with the capacity of humans to learn speech items by imitation. It can be assumed that this *speech mirror neuron system* in parallel co-activates motor representations, sensory representations, and phonemic representations of speech items. Given that from a phonetic viewpoint speech items also are built up by goal-directed actions (called *speech gestures*) which build-up the motor plans for speech items in our model (see Section 2), it can be hypothesized that a mirror neuron layer also exists for the association of motor, sensory, and phonemic representations of speech gestures (see also Westerman and Miranda, 2004).

*Self-organization* is a central principle of learning and self-organizing maps are used for modeling cortical networks (Kohonen, 2001). Within our neurocomputational model artificial self-organizing neural networks are implemented since self-organizing neural networks are biologically plausible and have been used successfully for modeling semantic lexical networks (Ritter and Kohonen, 1989), for (i) modeling semantic and phonological aspects during early lexical development (Li et al., 2004), and for (ii) modeling the generation and recognition of goal-directed movements (Bullock et al., 1993; Tani et al., 2004). A further argument for using self-organizing maps is their success in modeling the mapping between phonemic and phonetic aspects of speech production as demonstrated by the learning experiments for vowels and syllables described in this study.

In our current model different submaps are used for different classes of speech items (V, CV, VC) and separate training procedures were introduced for training these classes of speech items. This separation of the phonetic map in submaps as well as the separation of training procedures for different speech items was done in order to simplify the modeling of the speech acquisition procedure for these three classes of speech items from the computational viewpoint. But in principle all types of speech items (i.e. all types of syllables and words or word components) can be trained simultaneously by introducing just one comprehensive learning task and by using *one single phonetic map*. Recent preliminary experiments indicate that a comprehensive single phonetic map shapes different subregions representing different classes of speech items. The ordering of speech items within these subregions is similar to the pho-

netopic ordering presented in this paper for the different submaps discussed here.

It is unclear whether the training sets used here constitute a representative natural model of babbling and imitation training during early states of human speech acquisition. Our training sets comprise a widespread set of vocalic vocal tract positions and a widespread set of opening and closing movements. At least these sets comprise all vocal tract positions and all opening and closing movements which are physiologically possible. But it is conceivable that toddlers very quickly reduce their set of training items from all physiological possible positions and movements towards a subset of positions and movements which are especially important for speech.

It should be noted that our neural modeling approach does not include modeling of *temporal aspects* of neural functioning. Rather the temporal aspects of production and perception are included in the speech items and thus in the sensory, motor, phonetic, and phonemic states. In our production–perception model sensory and motor states of vowels and syllables are processed as a whole. Our modeling approach thus is sufficient as long as only a description of the training and processing of syllables is wanted. In contrast a detailed temporal organization becomes important if speech items comprise more than one syllable. In this case processing delays must be introduced for all pathways postulated in the model (cf. Guenther et al., 2006) and temporal aspects of neural activity need to be considered (cf. Maass and Schmitt, 1999).

The two training stages identified by our modeling study distinguish between *babbling* (i.e. the build-up stage for sensorimotor representations of pre-linguistic proto-vocalic and proto-consonantal speech gestures) and *imitation* (i.e. the build-up stage for language-specific perceptual, motor, phonetic, and phonemic representations of speech items). A closer modeling of early stages of speech acquisition (Oller et al., 1999) is beyond the scope of this paper. Furthermore in reality the two training stages introduced here overlap in time. This is partially realized in our approach, since babbling and imitation training items are applied in parallel during the imitation training stage after a short babbling training stage.

The next important step would be to introduce processes for building up the *mental lexicon* and for modeling the process of word segmentation and identification (cf. Batchelder, 2002; Werker and Yeung, 2005; Jusczyk, 1999; Brent, 1999). The representation of the mental lexicon of the target language is very important for including top-down processes of speech perception and thus for speech recognition. However consideration of these processes currently goes beyond the scope of the current implementation of our model. But the model in generally is open for integrating a mental lexicon.

Last but not least it has to be stated that the neurocomputational production–perception model developed thus far by no means is an alternative solution for high-performance speech recognition or speech synthesis systems. At

present the model described here is capable of producing and perceiving simple CV- and VC-syllables under ideal conditions. Concerning a further development of the model introduced here two different strategies are imaginable. On the one hand, this model can be further developed in order to handle more complex classes of speech items (words, sentences, or a whole discourse) under ideal and non-ideal conditions (e.g. different speakers, different emotional states, external noise). On the other hand, the organization of the neurocomputational model outlined in this paper could be integrated at least partially into the architecture of current or new speech recognition and speech synthesis systems.

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