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# Neurodynamical model for the multi-stable perception of biological motion

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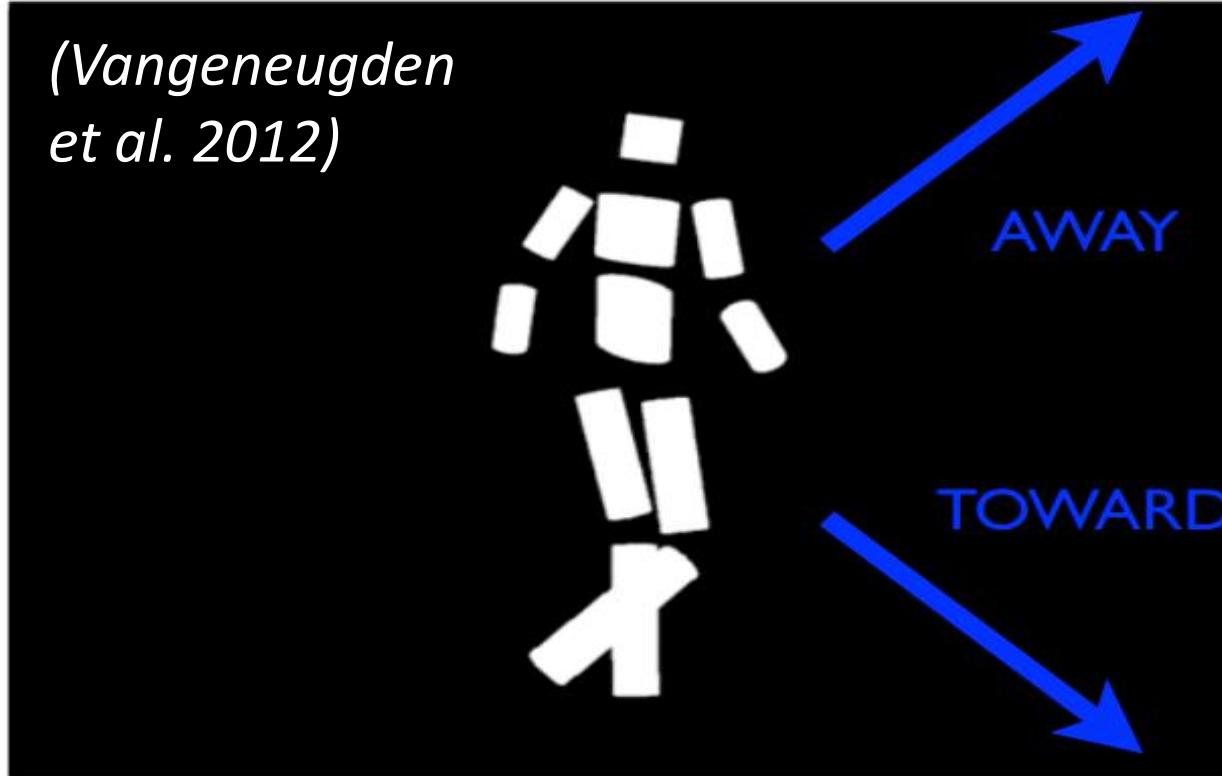


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## Introduction

- Body motion stimulus can induce bistable perception (Vanrie et al. 2004; 2006; Vangeneugden et al. 2012; Schouten et al. 2011).
- The perception of body motion has been modelled using physiologically plausible architectures (Giese & Poggio, 2003; Lange & Lappe, 2006). These models cannot deal with perceptual multi-stability.
- Repetition suppression/enhancement:** response adaptation to repetitive stimuli is important in fMRI paradigms in order to increase selectivity of analysis.
- Ambiguous results from fMRI adaptation paradigms for action stimuli (e.g. Dinstein et al. 2006; Lingnau et al. 2009).
- No or very weak repetition suppression observed at the single cell level for action stimuli (Caggiano et al. 2013; Kilner et al. 2014).

### Bistable body-motion stimulus



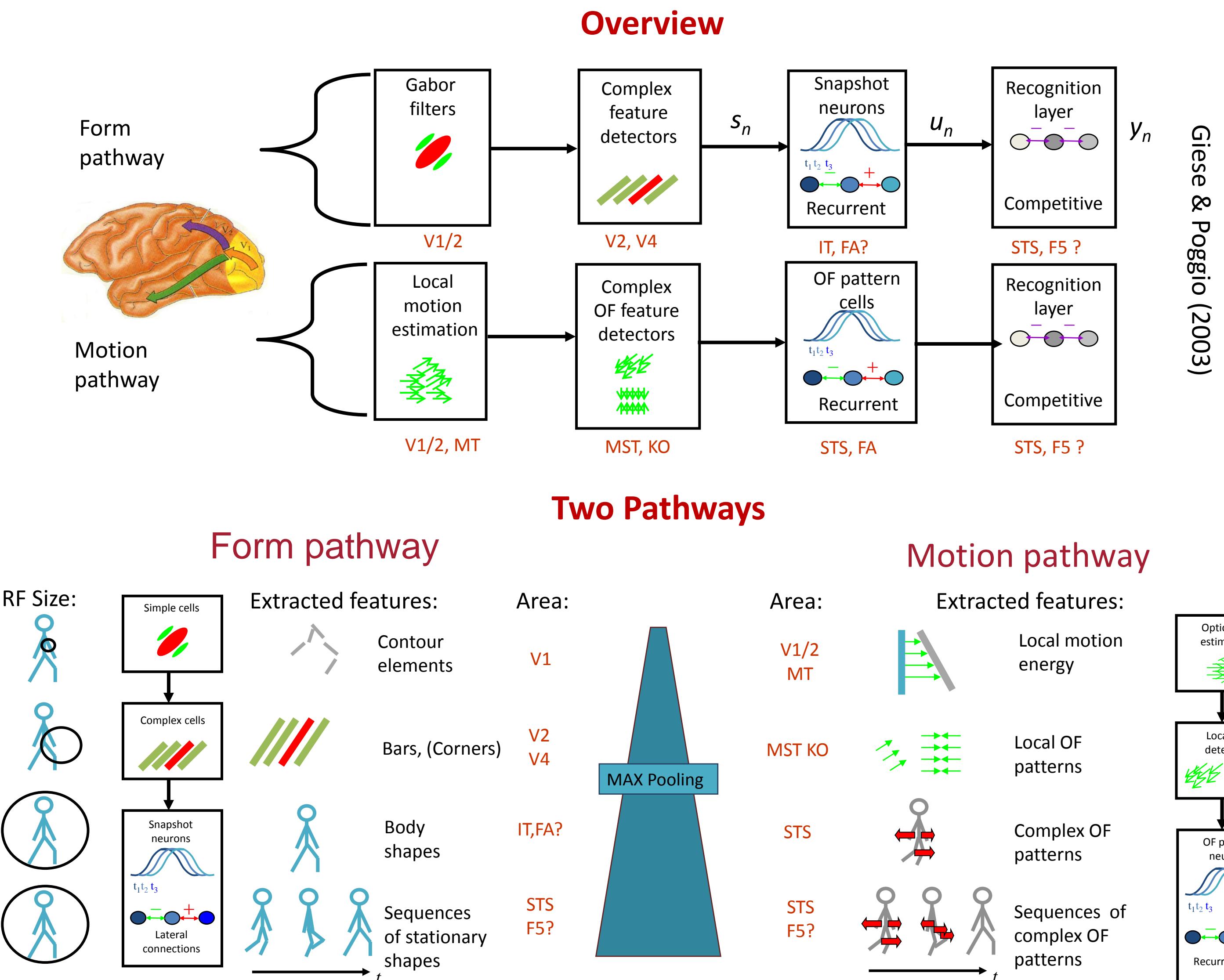
(Vangeneugden et al. 2012)

- No disparity cues.
- Upper and lower body consistent with motion in different directions.
- Two movement directions are perceived in alternation; perceptual switching (Vanrie et al. 2006).
- Similar multi-stability for natural walkers in oblique projection.
- Perceptual multi-stability observed for many other perceptual phenomena (reviews e.g. Blake et al. 2001, Leopold et al. 1999).

## Goal

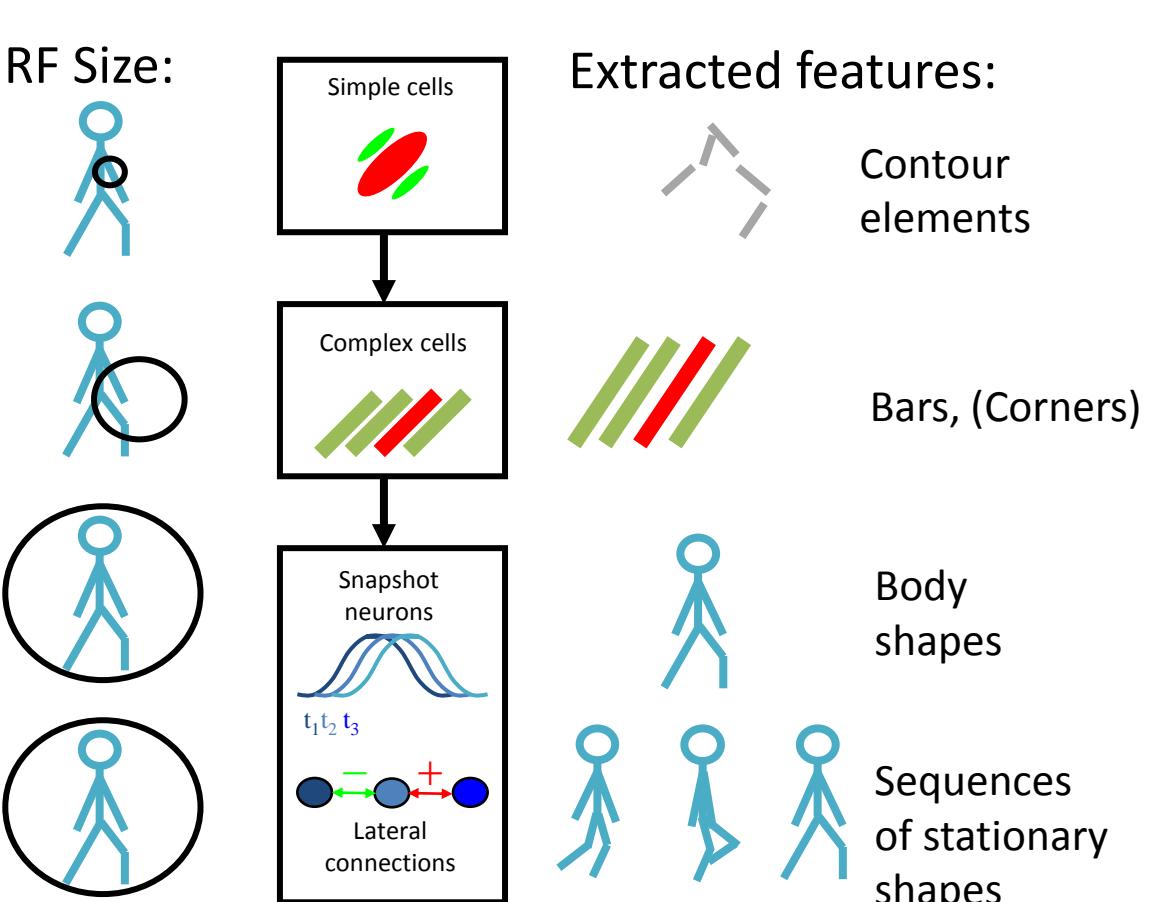
- Development of a model that accounts for these dynamic phenomena in body motion perception.

## Neural model for body motion perception



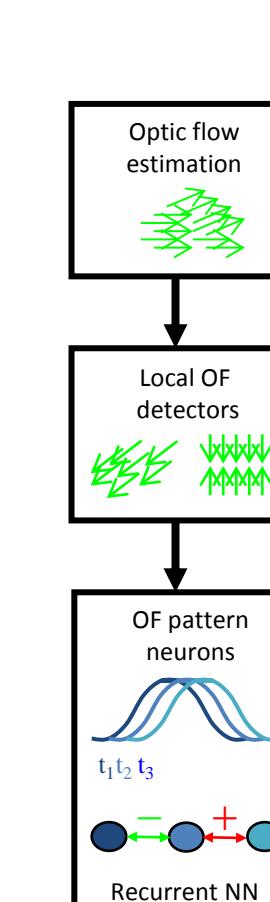
Giese & Poggio (2003)

### Form pathway



### Two Pathways

### Motion pathway



### Mechanism for temporal sequence selectivity: 1D Neural Field

Dynamics of the membrane potential of snapshot neurons: DNF (Amari, 1977)

$$\tau_u \dot{u}(\theta, t) = -u(\theta, t) + w(\theta) * 1(u(\theta, t)) + s(\theta, t) - h$$

- Sequence selectivity emerging from asymmetric lateral connections.
- Stimulus-locked stable travelling pulse solution. (Zhang 1996; Xie & Giese 2002).

u: membrane potential  
s: shape detector output  
θ: snapshot no.  
w: interaction kernel  
h: resting potential  
\*: convolution

- Single stimulus repetition (Caggiano et al. 2012; Kilner et al. 2014)
- New stimulus

## Extension: 2D Neural Field for view selectivity

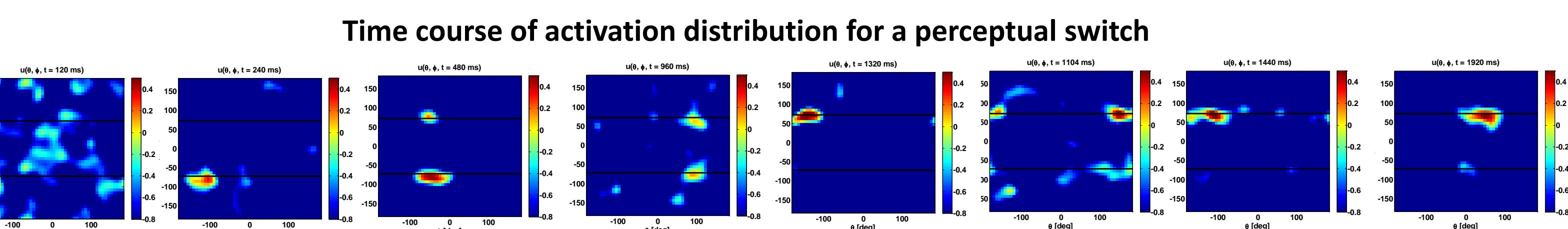
$$\begin{aligned} \tau_u \dot{u}(\varphi, \theta, t) &= -u(\varphi, \theta, t) + w(\varphi, \theta) * 1(u(\varphi, \theta, t)) + s(\varphi, \theta, t) - h \\ &\quad - \alpha a(\varphi, \theta, t) + \varepsilon(\varphi, \theta, t); \\ \tau_a \dot{a}(\varphi, \theta, t) &= -a(\varphi, \theta, t) + 1(u(\varphi, \theta, t)); \end{aligned}$$

Field dimensions:  $\varphi$  (view angle),  $\theta$  (view variable).  
Interaction kernel:  $w(\varphi, \theta)$ .  
Evolution of gait time variable:  $t$ .  
Change of walker view variable:  $\varphi$ .

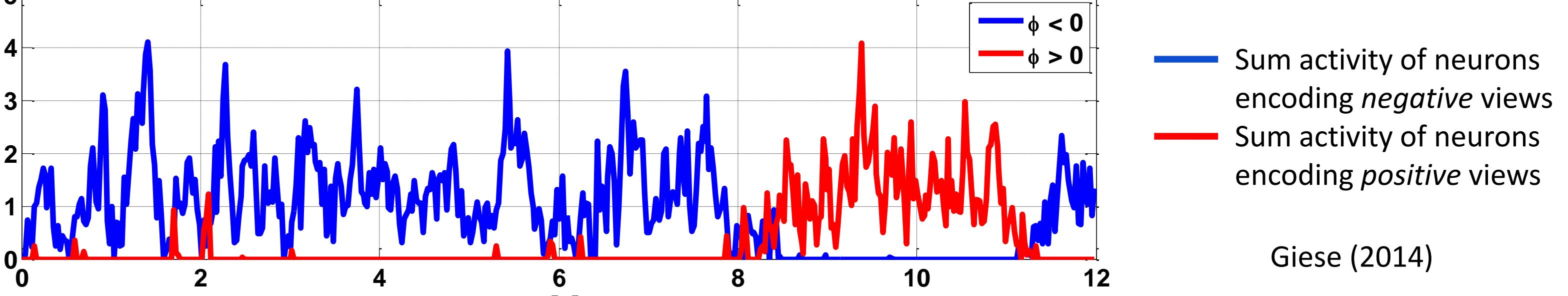
## Simulation results

### 1) Reproduction of perceptual bistability:

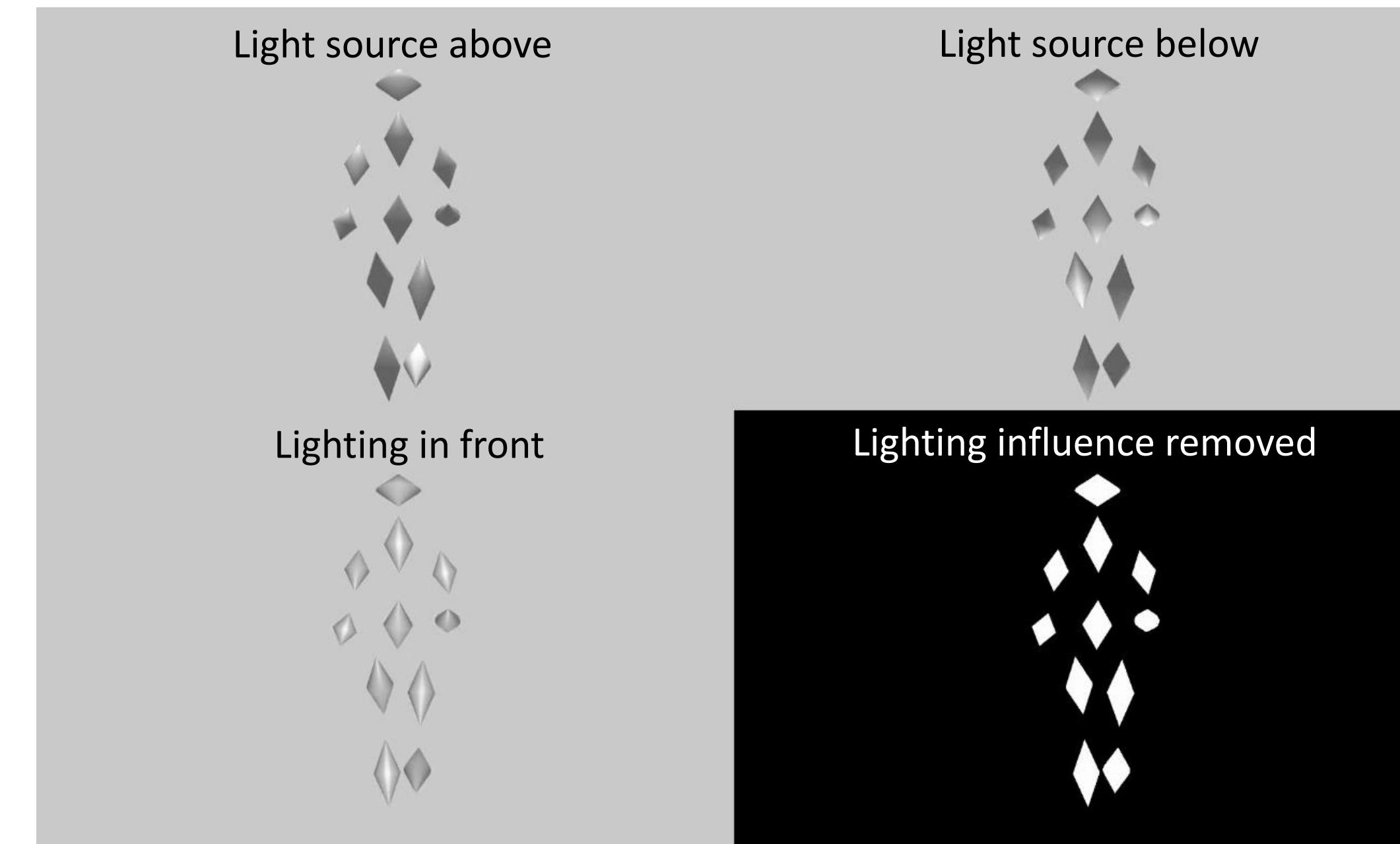
- Ambiguous view results in competition between solutions representing two opposite views.
- Perceptual switches induced mainly by internal noise (adaptation too weak).



### Sum activity as function of time



- Addition of shading removes the bistability with respect to view.
- Model need extension by 'shading pathway'.
- Extraction of internal shading gradients provides discriminative features.
- Extraction by modified filter hierarchy.
- Interesting new illusion shows influence of illumination direction: 'Lighting from above prior'



## Conclusions

- Extension of neurodynamical model for the encoding of body motion can account for multi-stability.
- Perceptual switches likely not driven by adaptation.
- Model accounts for weak adaptation in repetition suppression paradigms with action stimuli.
- Prediction of a new stimulus that should result in stronger adaptation.
- Influence of lighting direction on the perception of biological motion.

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