

Modeling

**pre-attentive stereo grouping
by intracortical interactions
in early visual cortex.**

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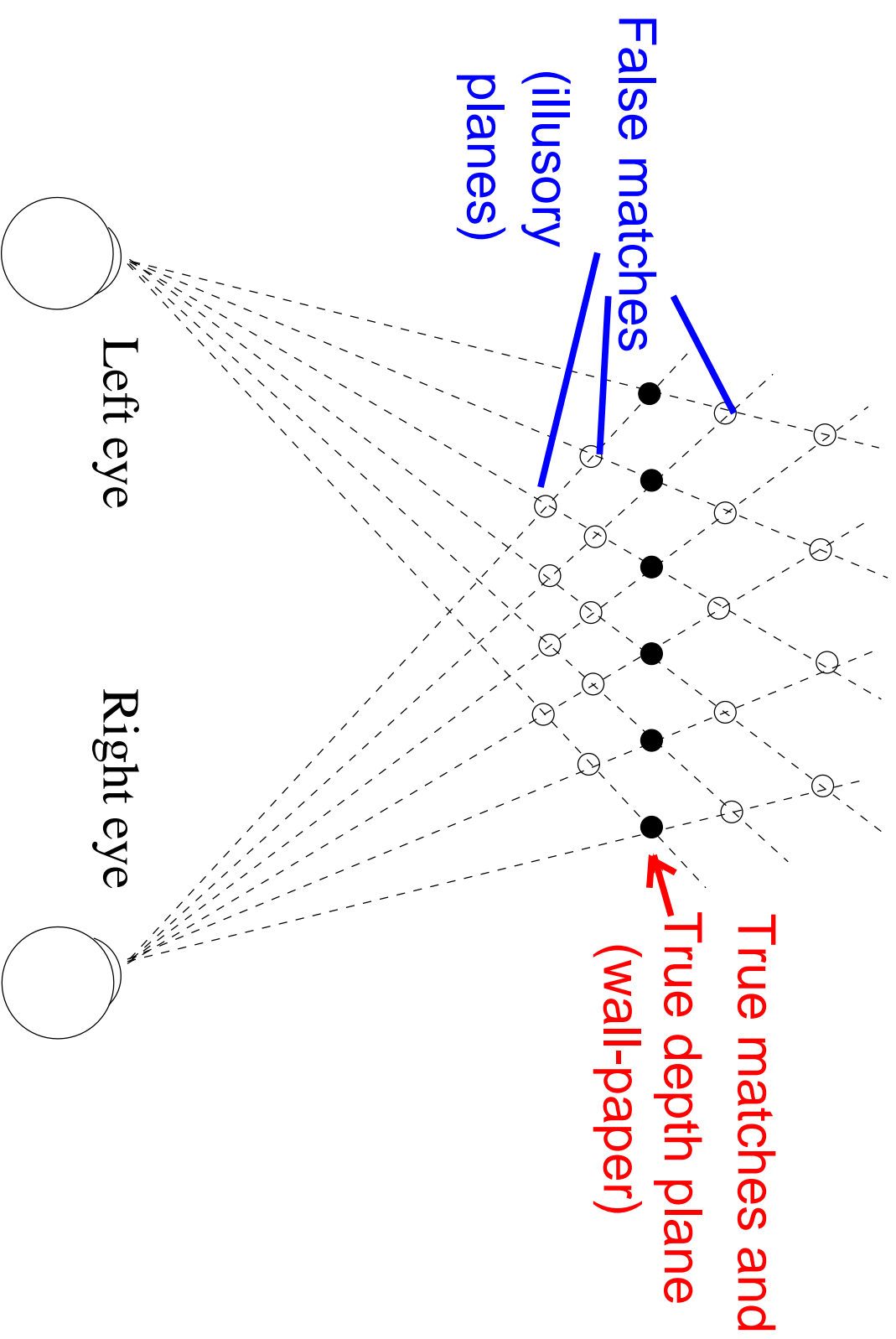
Purpose: to see if intracortical interactions can account for some stereo grouping phenomena in physiology and psychophysics

Stereo Grouping Phenomena modelled here:

- Enhanced V2 responses to stereo edges. (von der Heydt et al, 2000).
- Disparity capture (wall-paper effects) manifested by V2 responses (Bakin et al, 2000).
- Pop-out of a target of a unique depth from distractors of a different depth.
- Transparency.

The stereo matching problem

Example: the wall-paper effect



the visual input samples both the true and the false matches

L neurons respond to both the true and false matches.
(Summing and Parker, 2000)

but the cortex has to compute the true matches only

L2 neurons respond to only the true matches in the wall-paper defect (Bakin, Nakayama, and Gilbert, 2000 observed that a V2 neuron's response is tuned to the disparity value outside its receptive field, at the boundary of the depth plane (of the wall-paper-like matching.).

The stereo segmentation (grouping) problem

detecting or highlighting discontinuities in depth, or depth

edges, can serve segmentation, e.g., to segment two nearby planes of different depth, or to detect a target of a different depth — **pop-out**.

In der Heydt, Zhou, and Friedman, (2000) observed that **V2 cells respond more vigorously when their receptive fields are near a depth edge**.

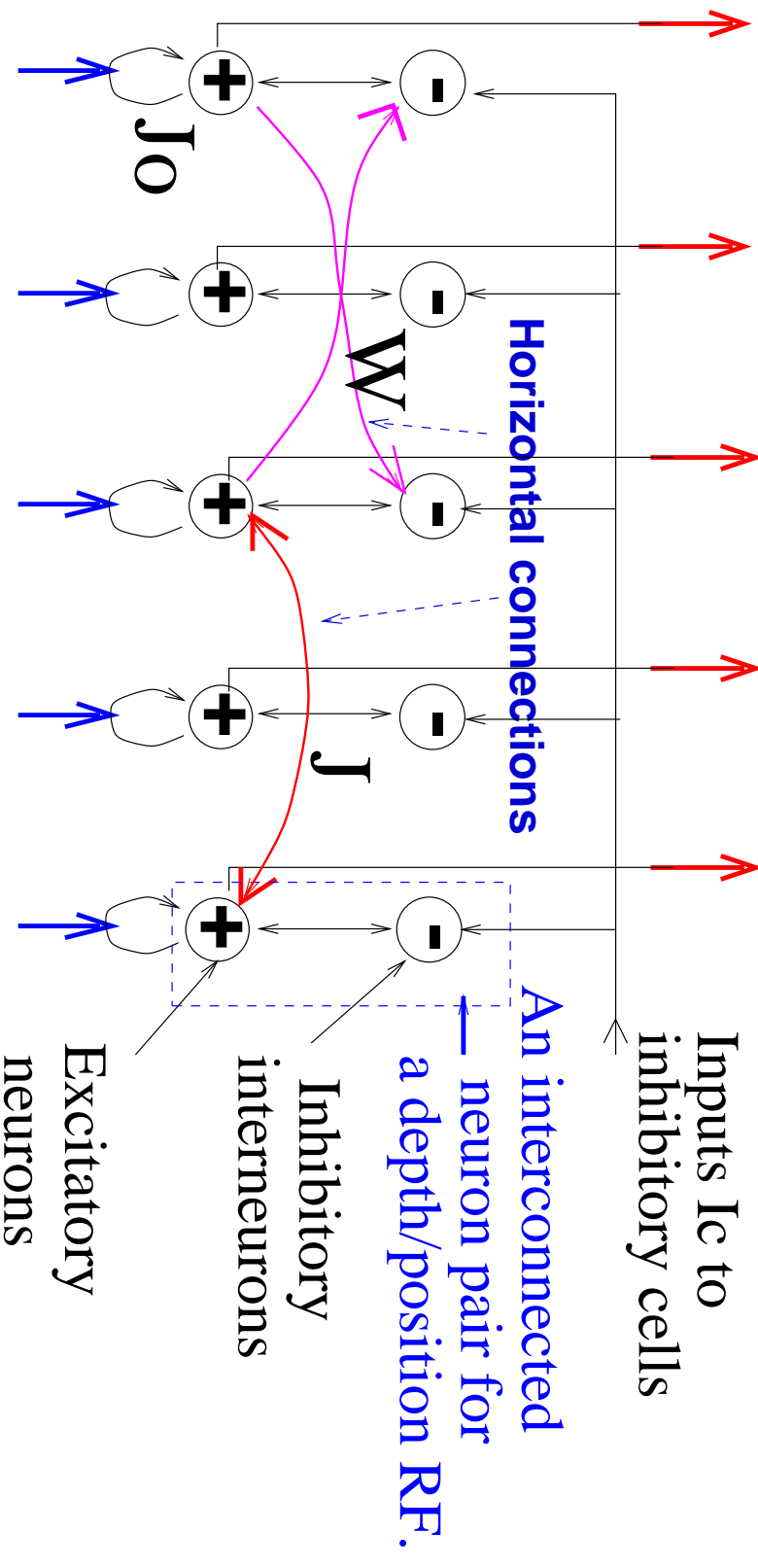
transparency: perceptually segregating two superimposed depth planes.

The model

Cortical outputs to higher visual areas

higher responses at depth discontinuity

outputs include mainly true matches only



Visual inputs, filtered through the disparity tuned RFs, to the excitatory cells.

Inputs include both **TRUE and FALSE** matches between monocular inputs

Model features and elements

• The model aims to emulate intracortical computations in **V2**.

• **Each model unit is binocular and disparity tuned.** The excitatory cells model cortical pyramidal cells. Each excitatory cell couples with a local interneuron to form a disparity tuned unit with a finite (small) size receptive field. **Other input dimensions (e.g., orientation, color, motion) are omitted.**

• Different cells are tuned to different depths, the receptive fields (RFs) of all model cells **sample 3D visual space** (2D fronto-parallel and 1D depth).

Both true and false matches provide input to the model (pyramidal) cells.

Long but finite range horizontal connections mediate monosynaptic excitation and disynaptic inhibition between nearby pyramidal cells. Horizontal connections tend to link cells tuned to similar depth.

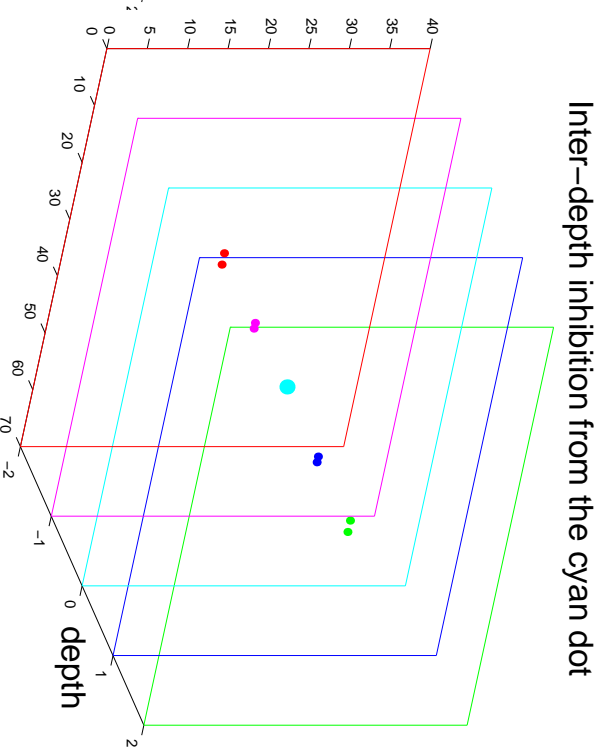
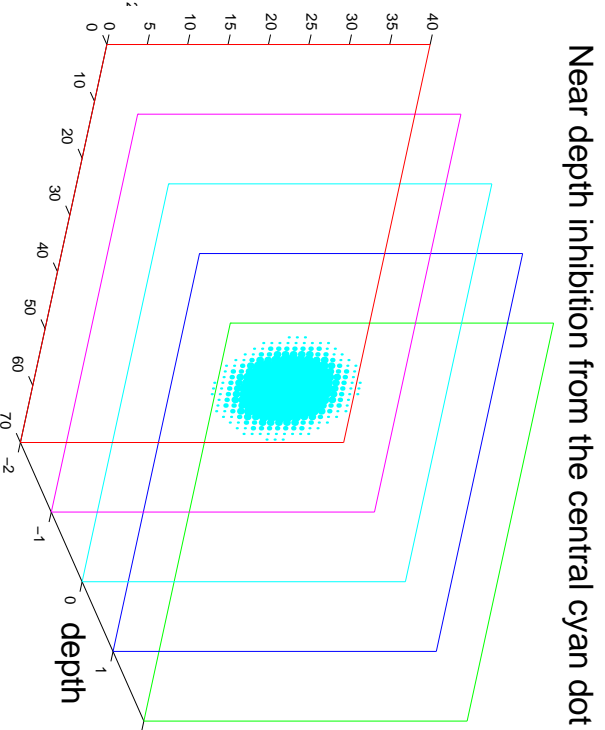
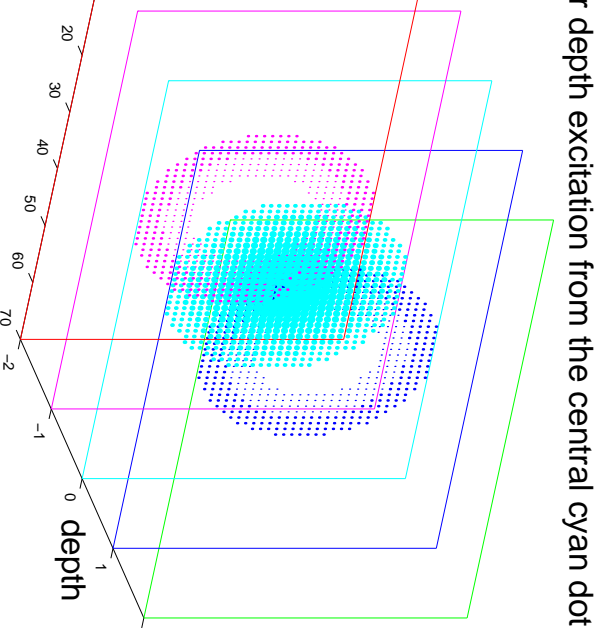
Cells responding to the same monocular location but different depths inhibit each other.

Horizontal connections mediate contextual influences such that, after initial transients, (1) the model cells respond significantly only to the true matches, and, (2) responses to depth discontinuities (depth edges or pop-out targets) are relatively higher.

The model horizontal connection pattern

There are 3 components in the connections

Near depth excitation Near depth inhibition Inter-depth suppression



Model cells sample the visual space of 5 depth planes (5x70x40 3-d locations).
Different depth planes are color coded

Each dot on a depth plane represents an RF center.
Dot size codes interaction (or response) strength

The equations of motion:

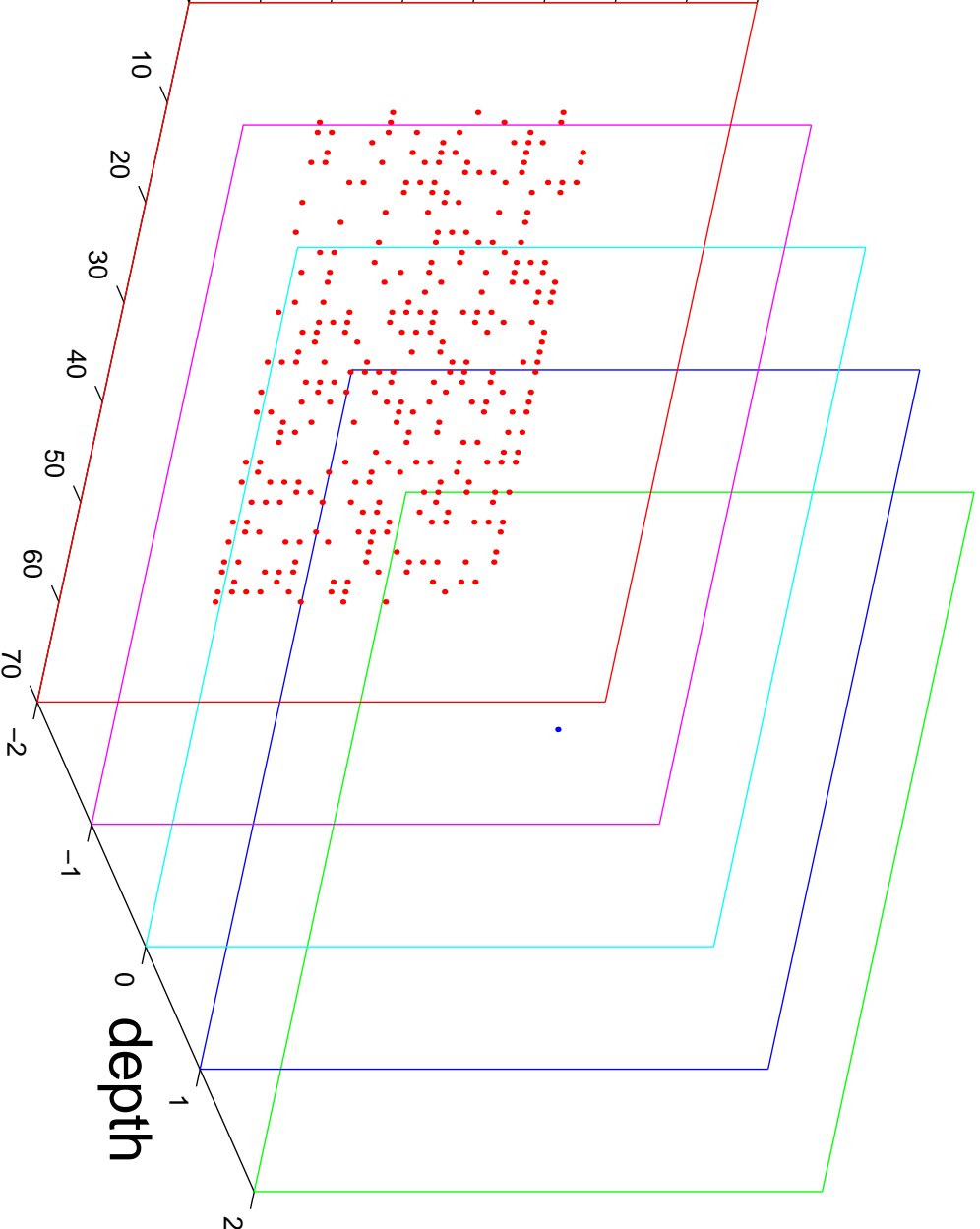
$$\dot{x}_{id} = -\alpha_x x_{id} - g_y(y_{id}) + J_0 g_x(x_{id}) + \sum_{j,d' \neq i,d} J_{id,jd'} g_x(x_{jd'}) + I_{id} + I_0$$

$$\dot{y}_{id} = -\alpha_y y_{id} + g_x(x_{id}) + \sum_{j,d' \neq i,d} W_{id,jd'} g_x(x_{jd'}) + I_c$$

for y : membrane potential for excitatory or inhibitory cells, i, d index of frontal parallel location i and depth d , J , W , horizontal connection matrix, g sigmoid-like activation functions, I_{id} visual inputs, etc.

Model computation illustrated by Popout

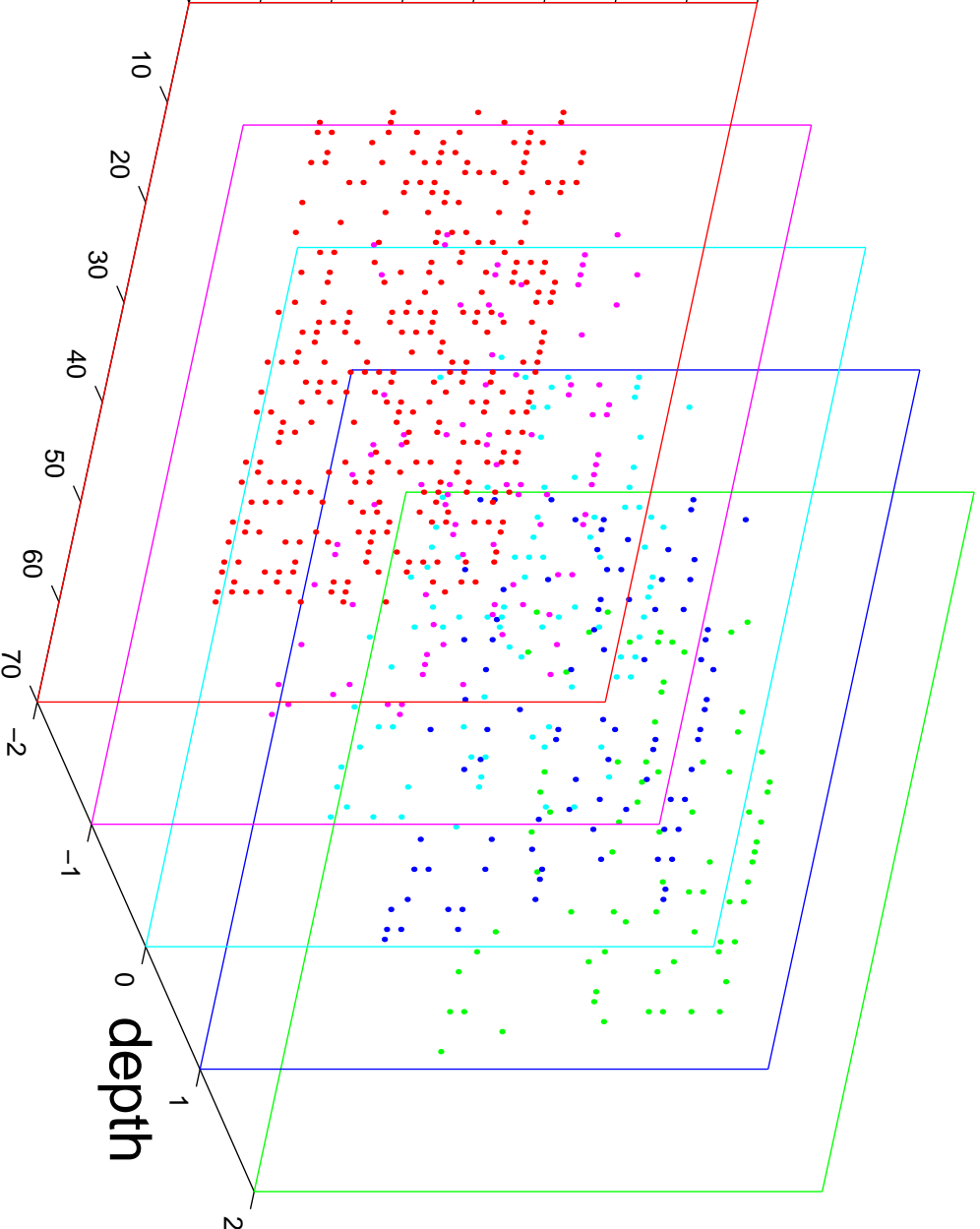
Scene Input



The scene consists of one
depth plane and a single
lone target dot at a
different (blue) depth

Input to the model

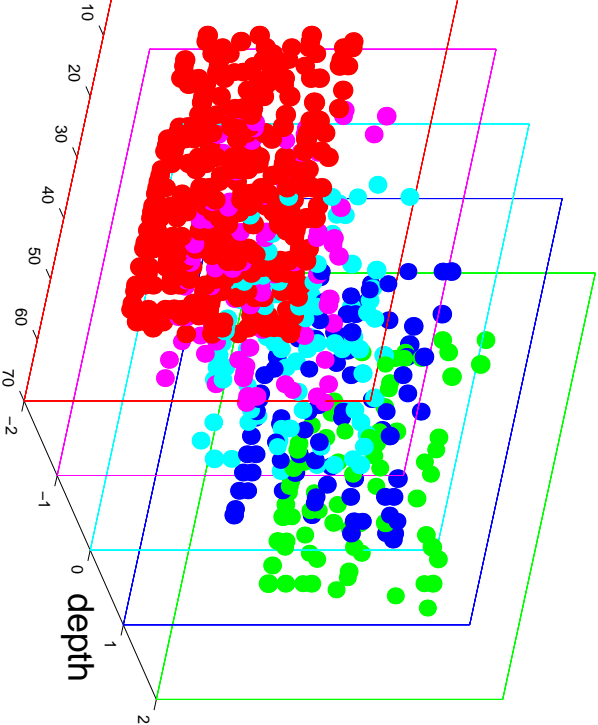
Input to cortex



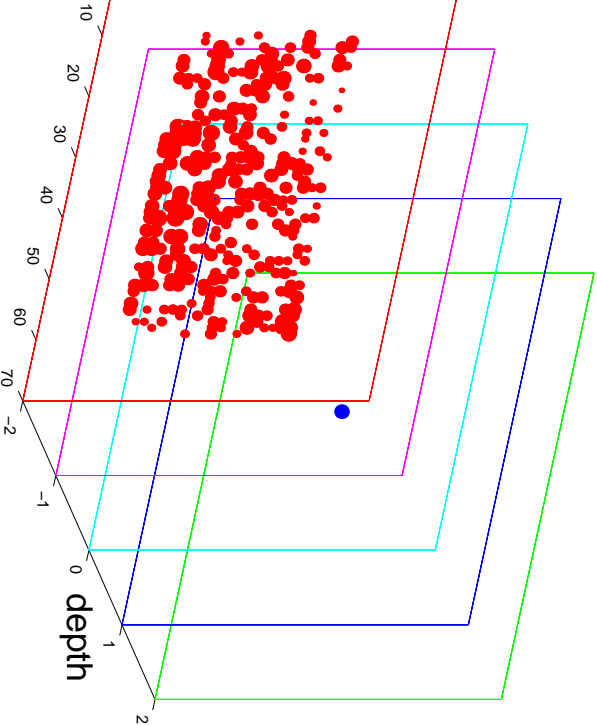
contains true and false
matches

Evolution of activity with time

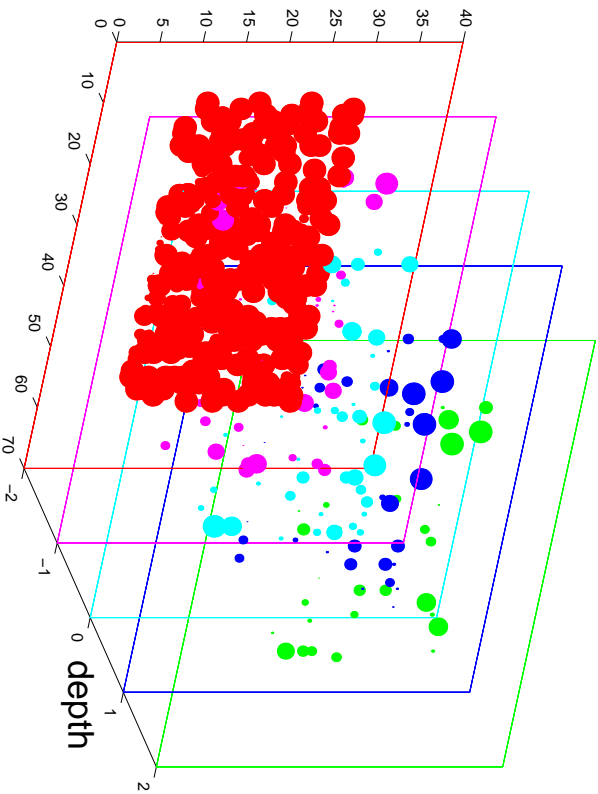
cortical activity at 0.60 membrane time constants



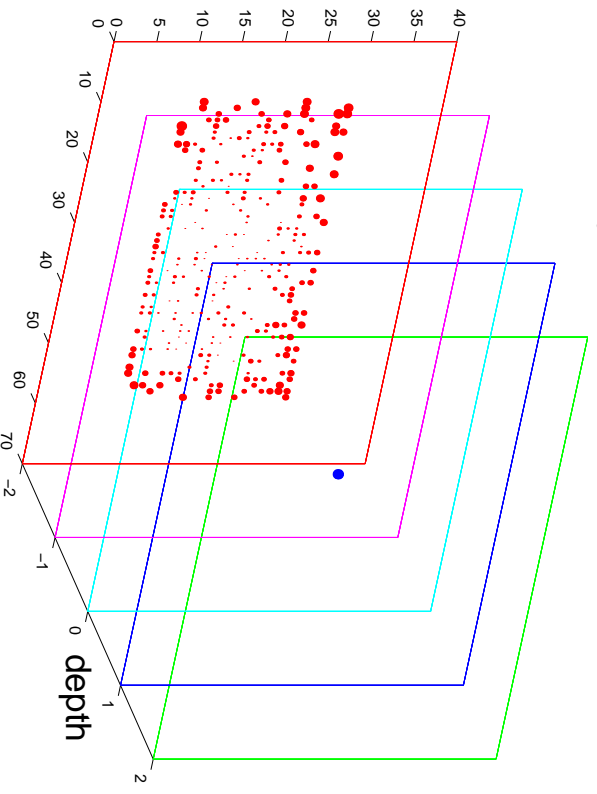
cortical activity at 6.00 membrane time constants



cortical activity at 1.40 membrane time constants

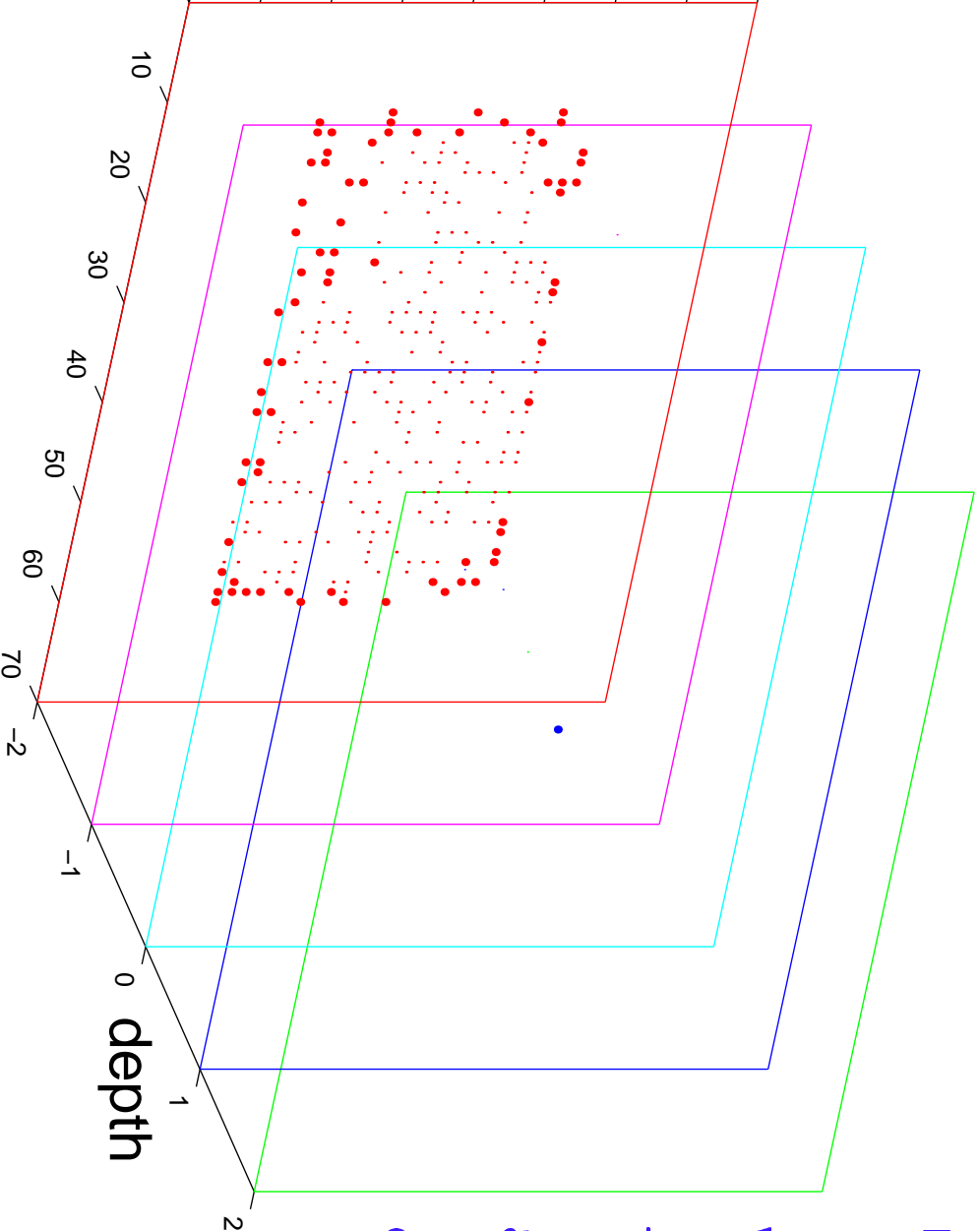


cortical activity at 12.80 membrane time constants



After initial transients, time averaged model response

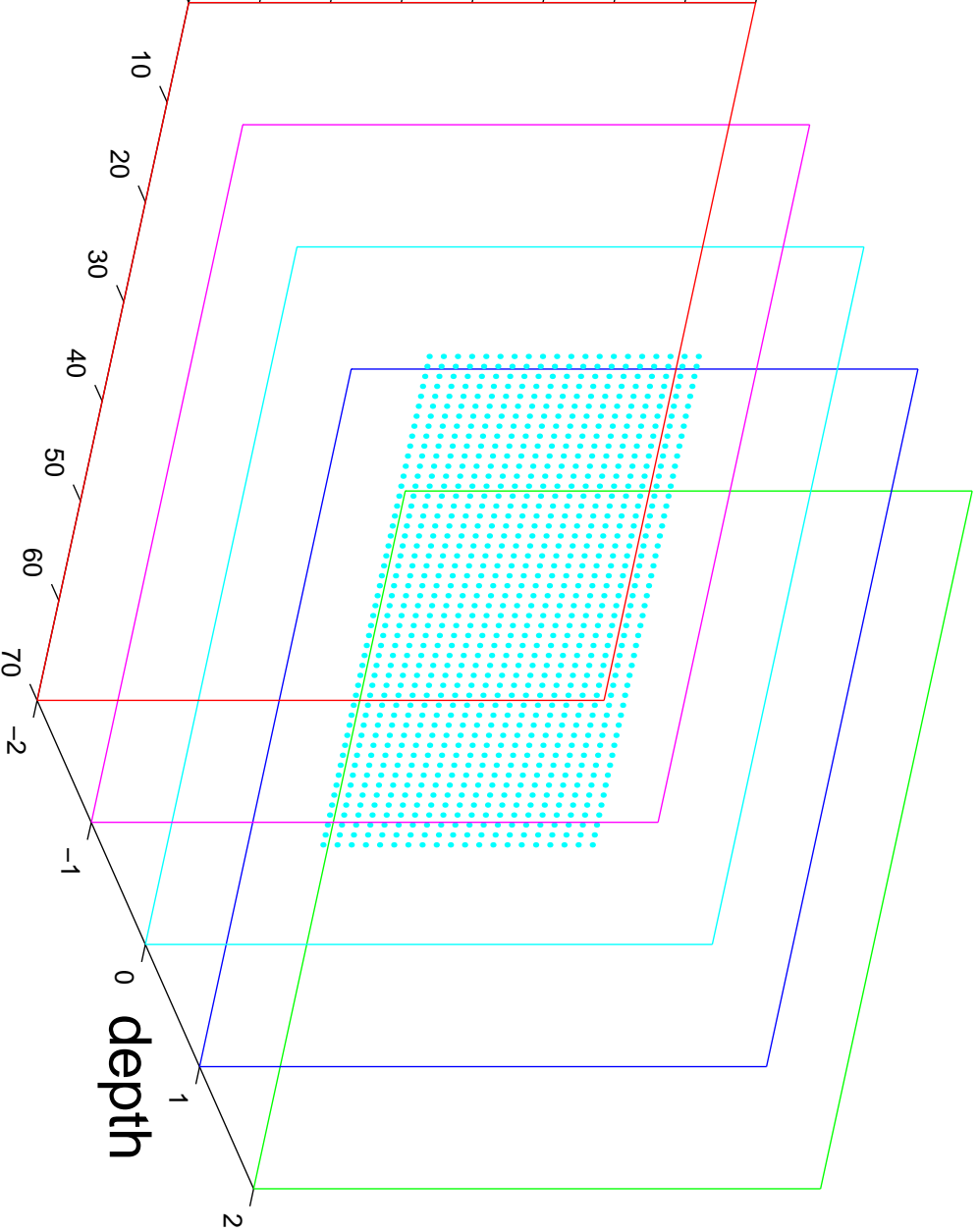
Time averaged cortex activity



contains mainly true
matches
with higher responses to
the lone target — **popout**
and to the borders
of the depth plane

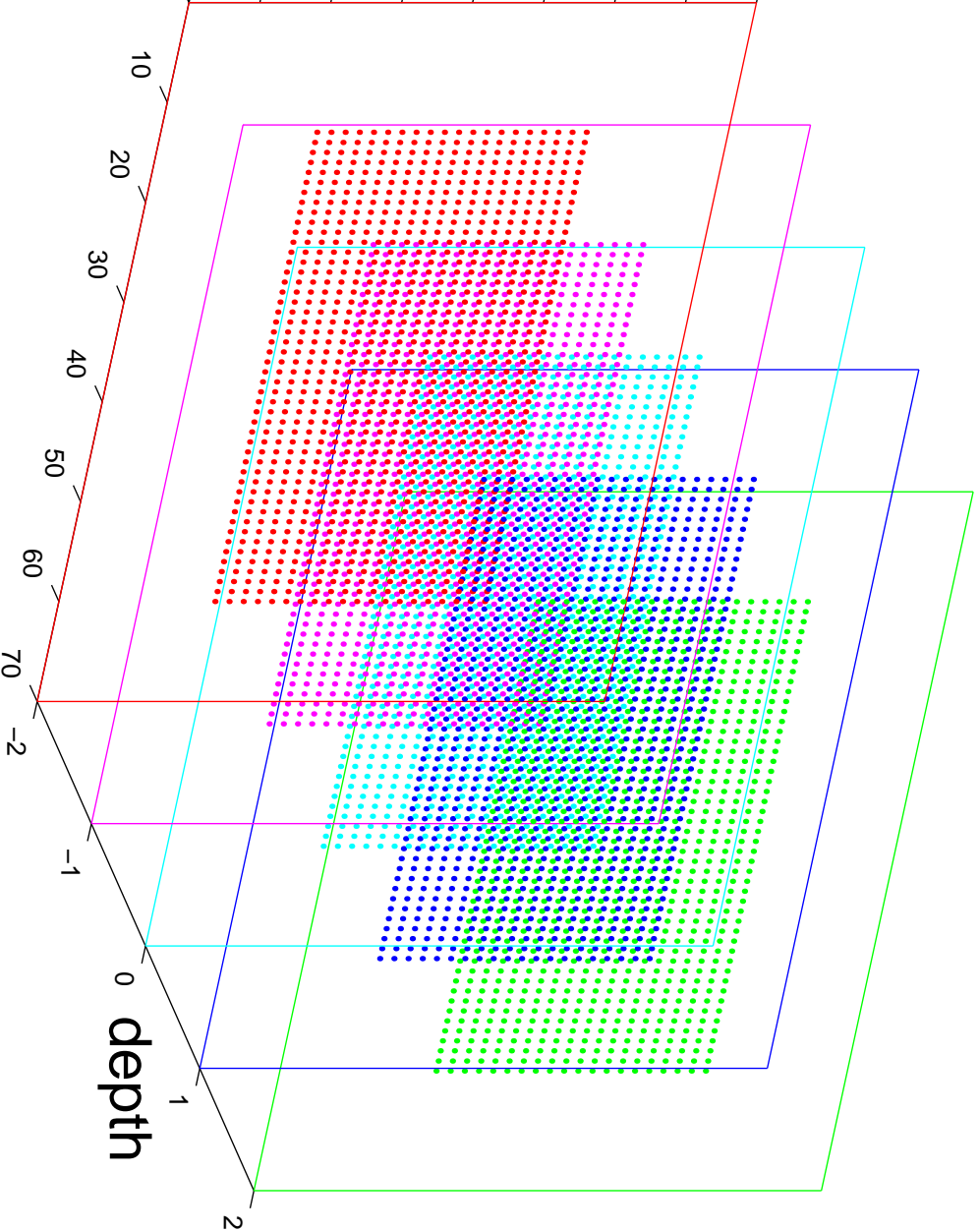
Model computation for Disparity Capture

Scene Input



Input to the model

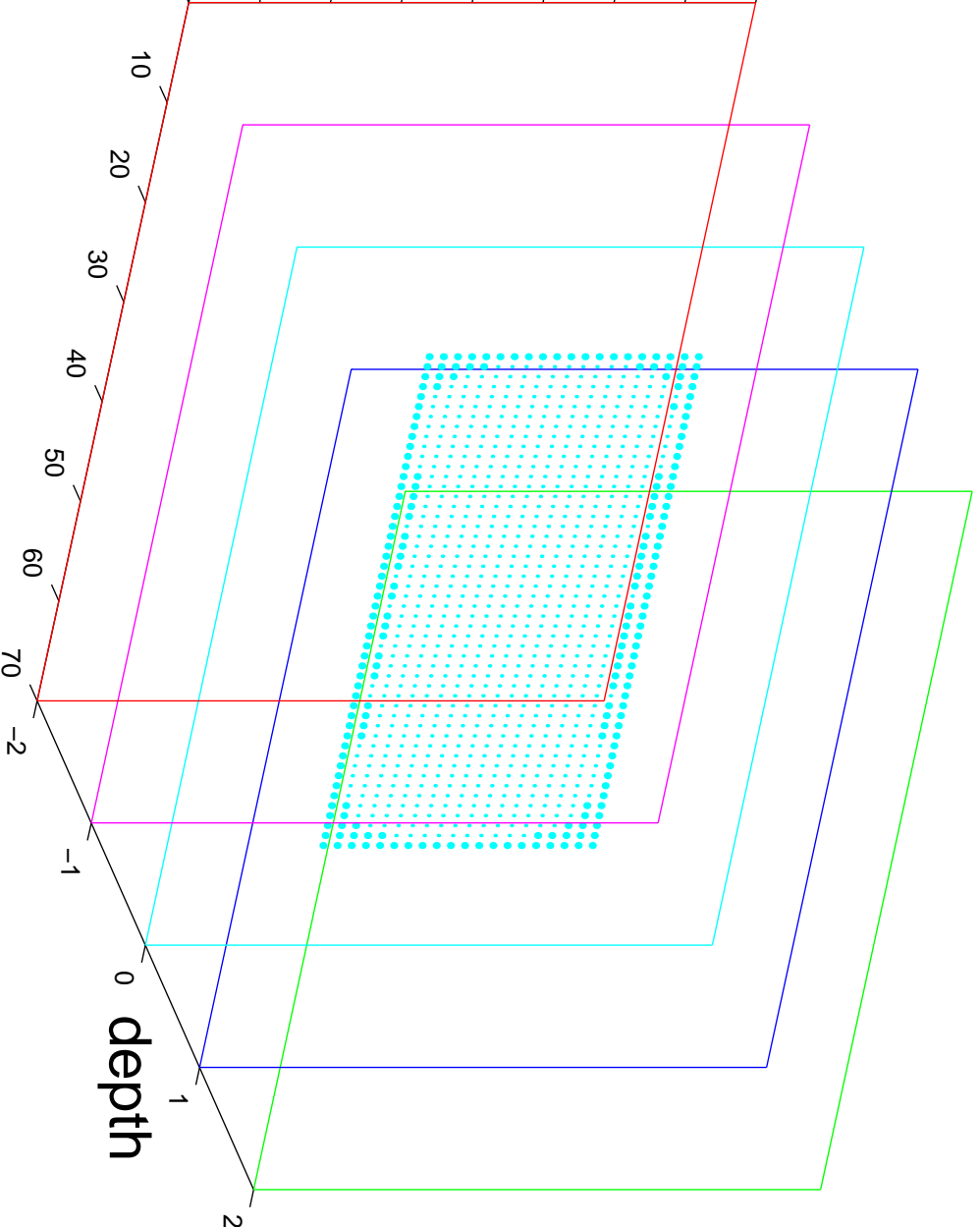
Input to cortex



contains true and
illusory depth planes
from true and
false matches

After initial transients, time averaged model response

Time averaged cortex activity

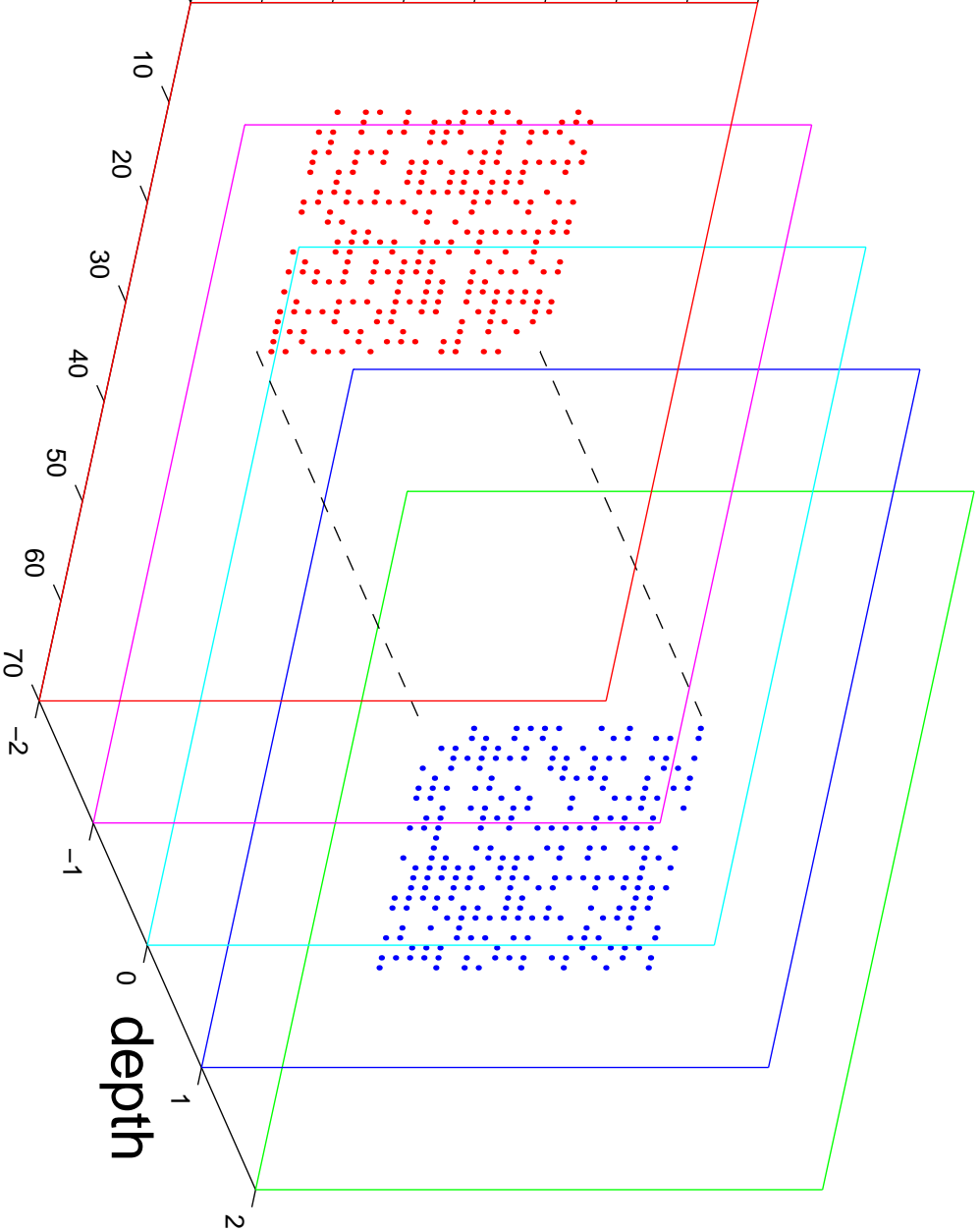


**Higher responses to
the borders of the plane
enable disparity
information at the
boundaries to propagate
into the center
of the plane**

**Contextual influences
enable disparity
information at the
boundaries to propagate
into the center
of the plane**

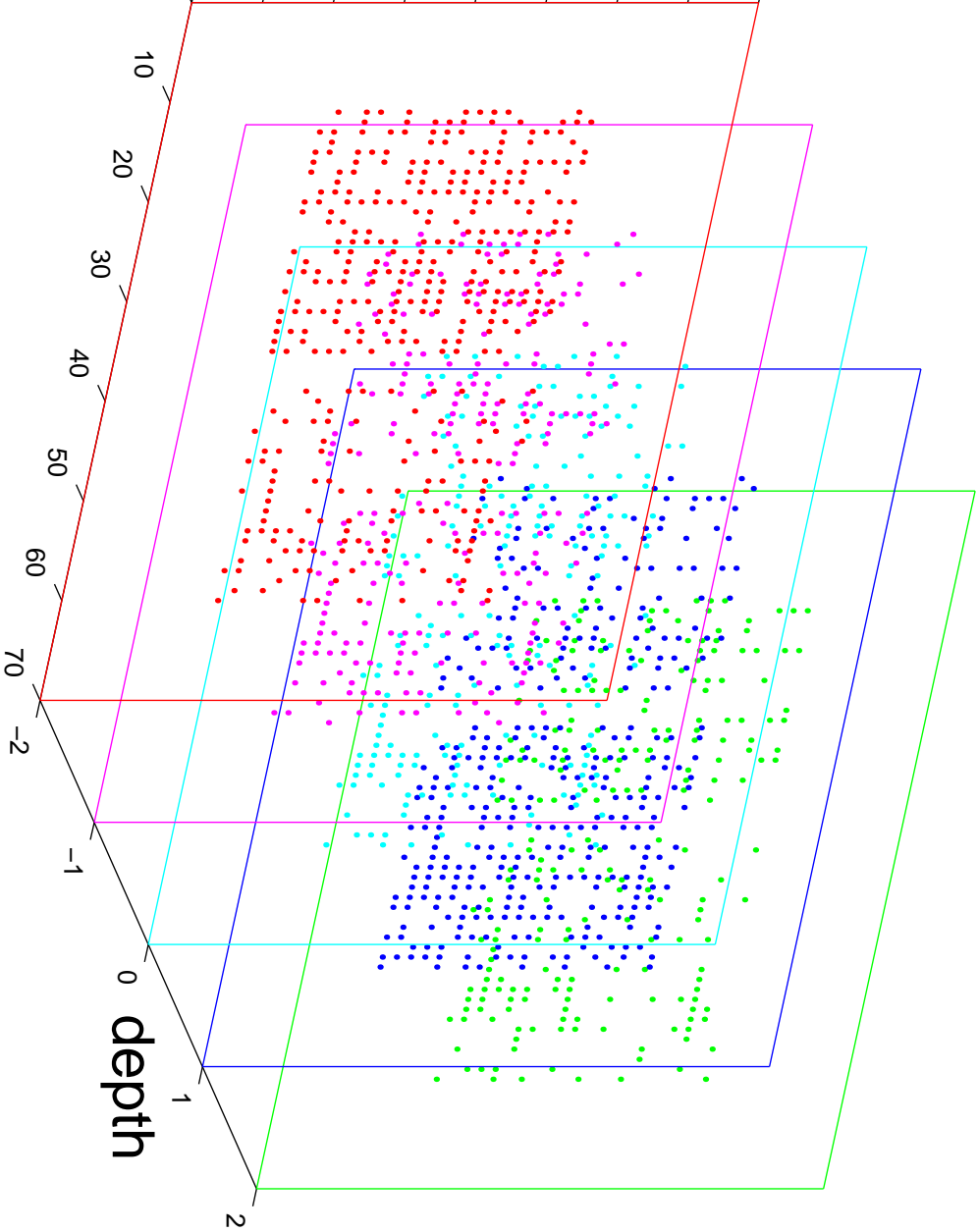
Model computation for Depth Discontinuity

Scene Input



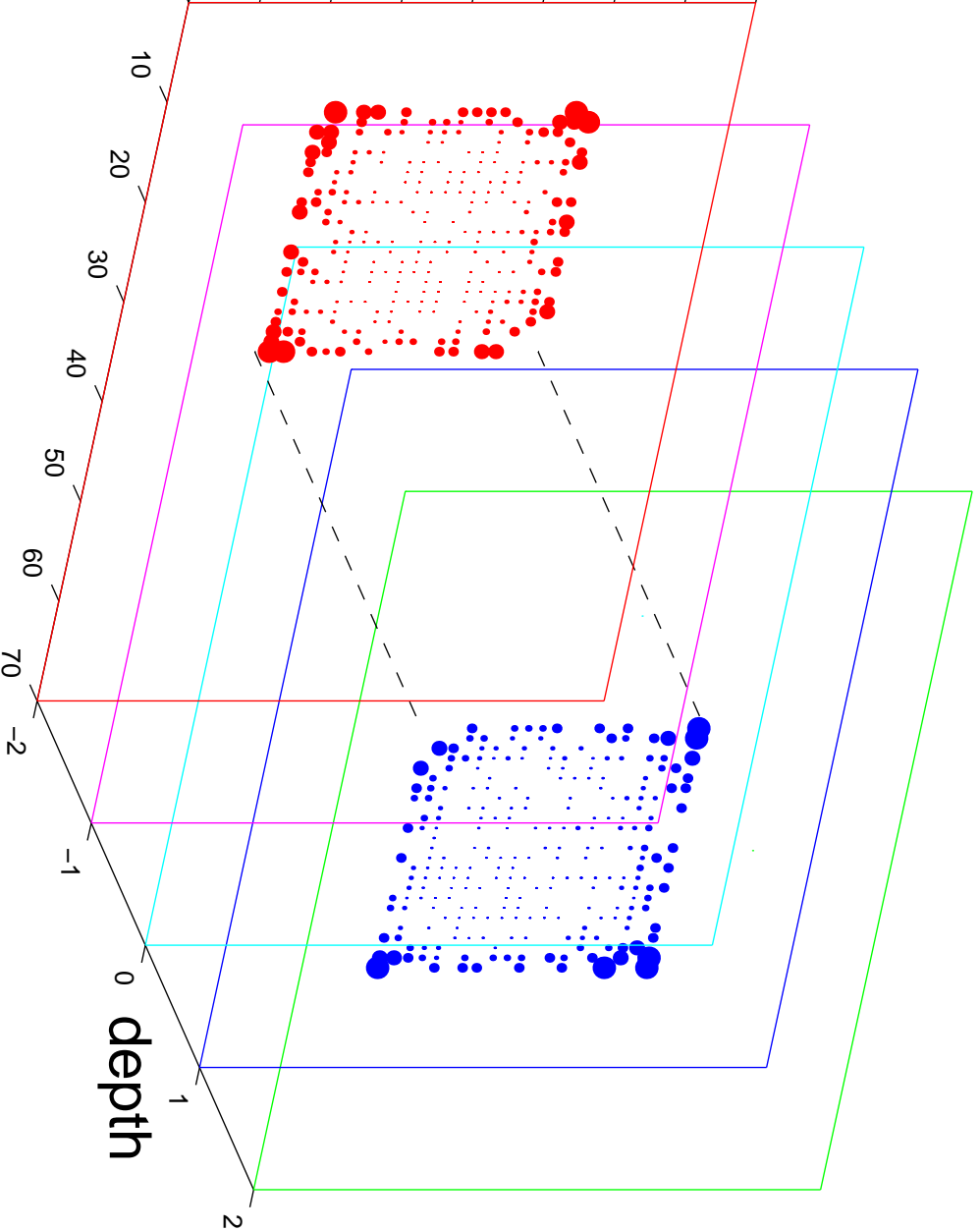
Input to the model

Input to cortex



After initial transients, time averaged model response

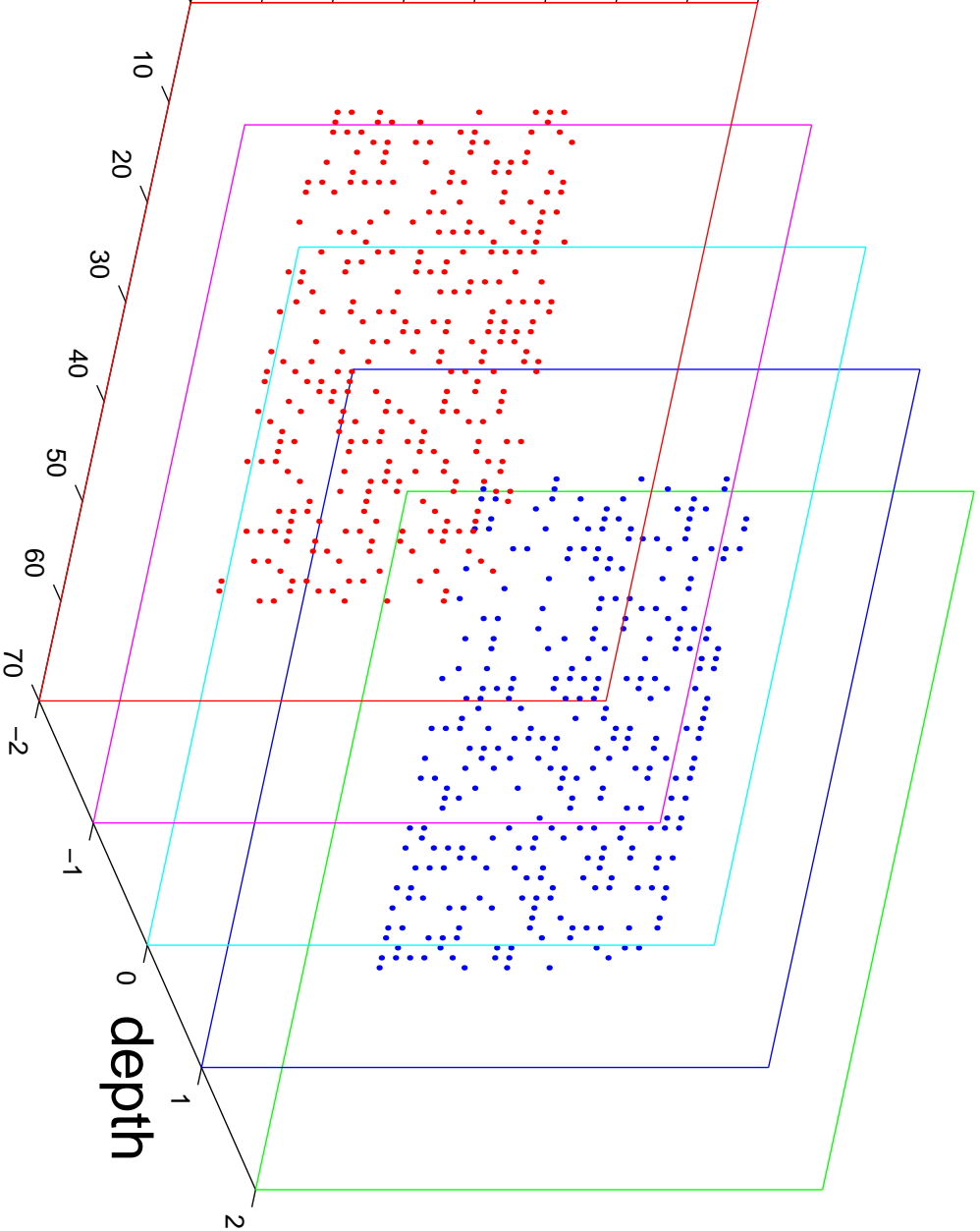
Time averaged cortex activity



Higher responses to
the depth discontinuity
and boundaries

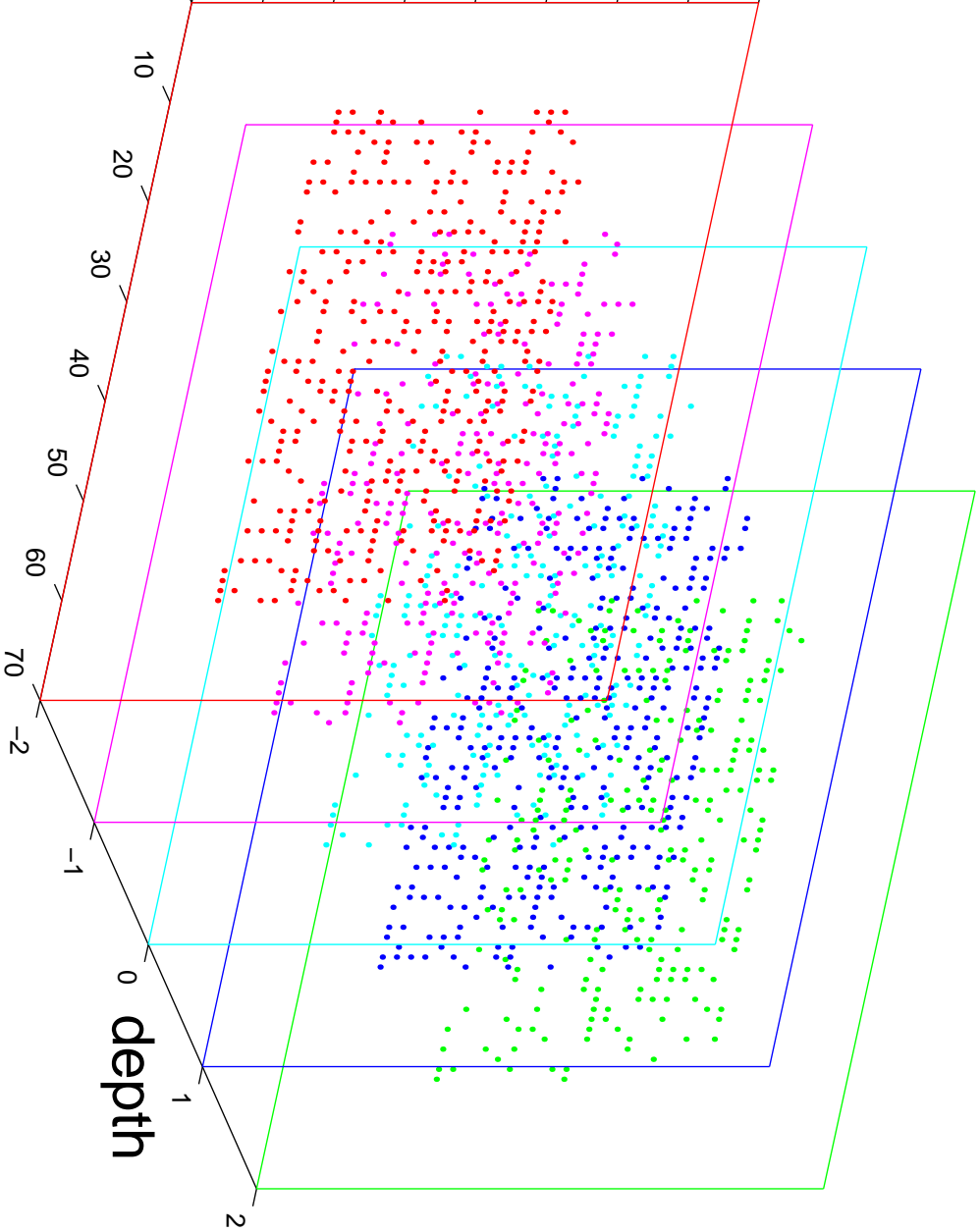
Model computation on Transparency

Scene Input



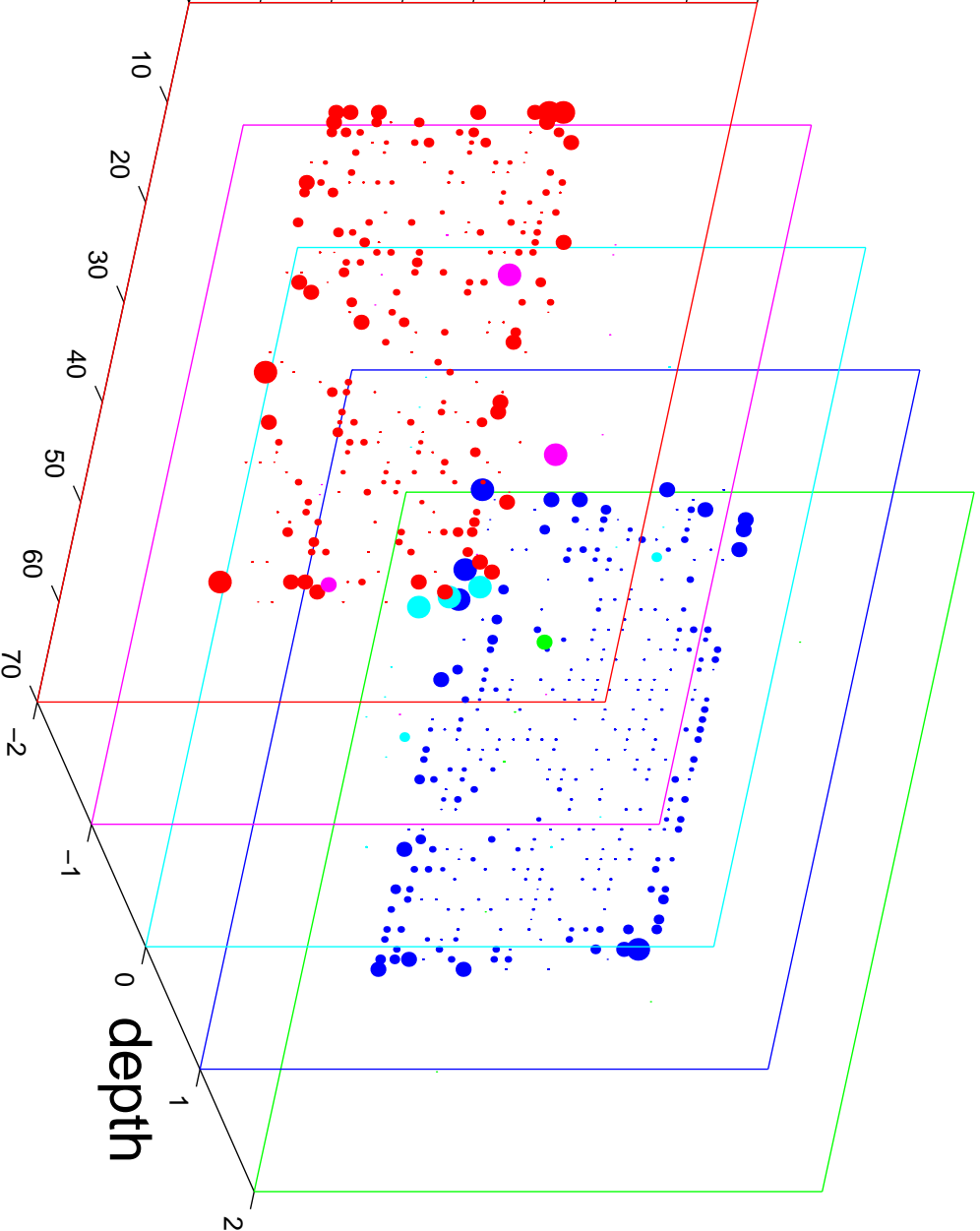
Input to the model

Input to cortex



**After initial transients,
time averaged model response**

Time averaged cortex activity



Depth segregation

Higher responses to
the plane boundaries
plus ghost dots

Summary and Discussion

- › Aim to capture both physiology and psychophysics of stereo grouping.
- › Suggest contextual influences in early cortex play important roles.
- › Relating to previous models: (1) Use cooperative algorithms like previous models (e.g., Marr and Poggio); (2) popout and depth edge highlights modelled for the first time; (3) more physiologically realistic mechanisms for transparency than previous models (e.g., Prazdny, 1985, Pollard et al 1985, Nishihara, 1987, Qian and Sejnowski, 1989, Marshall et al 1996).

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Due to unexpected difficulties to obtain a U.S. visa for travel, the author is very sorry not to be able to attend this VSS meeting. Many thanks to Ariella Poppo for her help to present this poster.