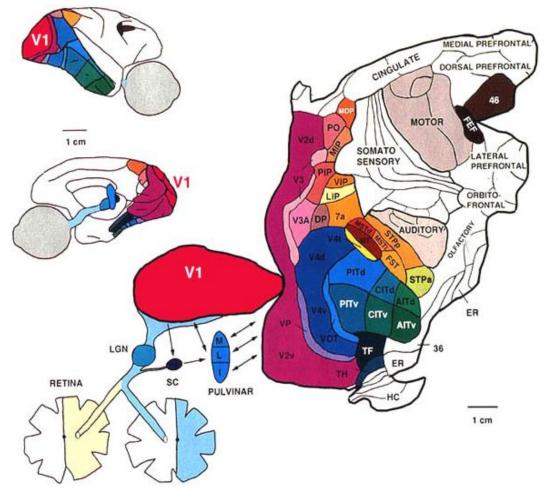
Modeling Extrastriate Areas

Dr. James A. Bednar

jbednar@inf.ed.ac.uk http://homepages.inf.ed.ac.uk/jbednar

Higher areas

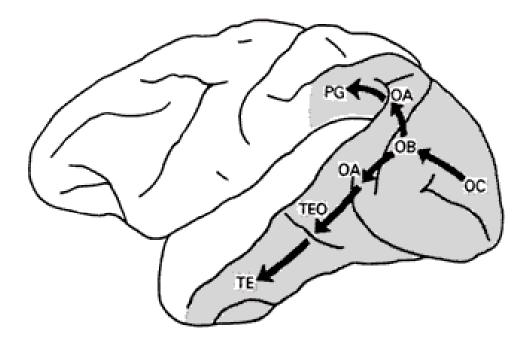


Macaque visual areas (Van Essen et al. 1992)

- Many higher areas beyond V1
- Selective for faces, self-motion, etc.
 Not as well
 - understood

What/Where streams

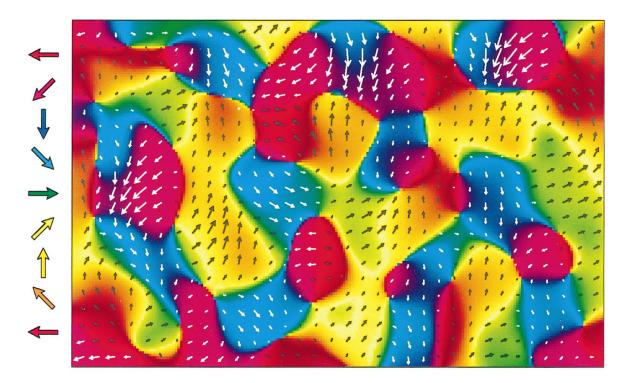
Typical division:



Ventral stream:

"What" pathway from V1 to temporal cortex (IT) **Dorsal stream:** "Where" pathway from V1 to parietal cortex (e.g. MT)

V2 OR/DR map

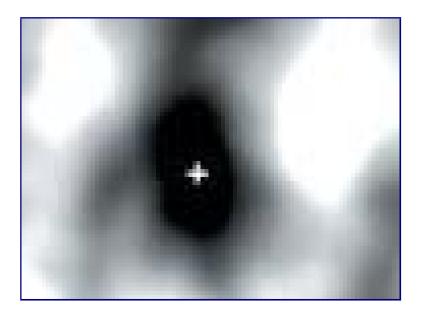


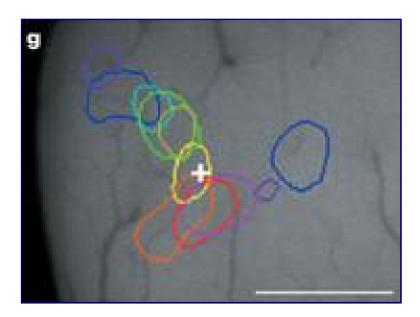
V2 cat direction map (Shmuel & Grinvald 1996)

Maps found in V1 are usually also found in V2 (except OD)

RFs are larger, probably more complex (not really clear)

V2 Color map

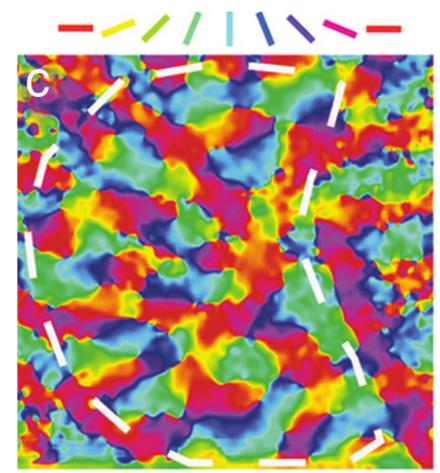




Xiao et al. $2003 - Macaque; 1.4 \times 1.0$ mm

- Like V1, color preferences organized into blobs
- Rainbow of colors per blob (Xiao et al. 2007: in V1 too?))
- Arranged in order of human perceptual color charts (CIE/DIN)
- Feeds to V4, which is also color selective

MT/V5



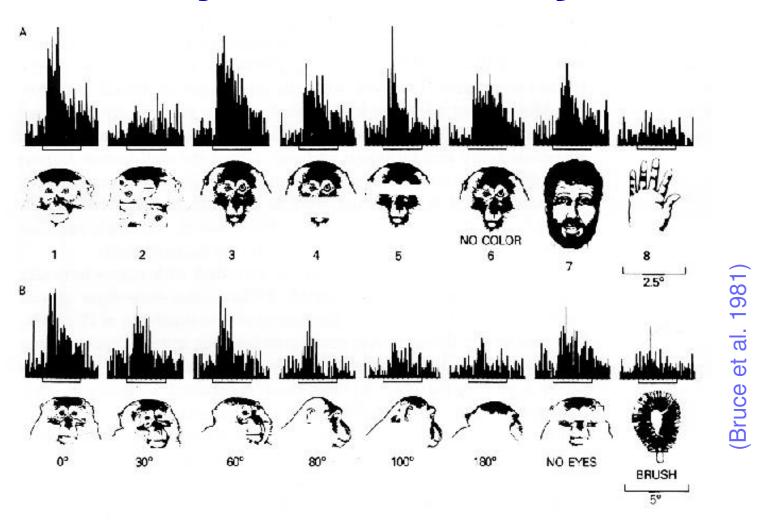
MT has orientation maps, but the neurons are more motion and direction selective

Involved in estimating optic flow

Neural responses in MT have been shown to directly reflect and determine perception of motion direction

(Britten et al. 1992; Salzman et al. 1990)

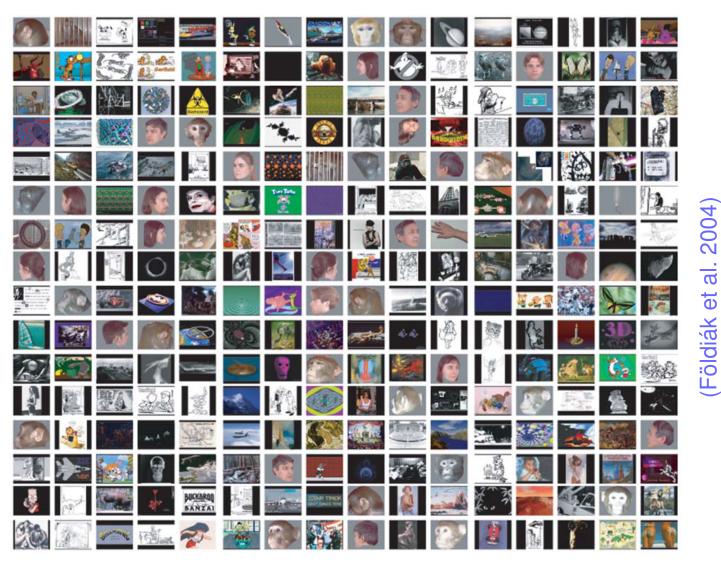
Object selectivity in IT



Some cells show greater responses to faces than to other classes; others to hands, buildings, etc.

CNV Spring 2008: Extrastriate models

Rapid Serial Visual Presentation



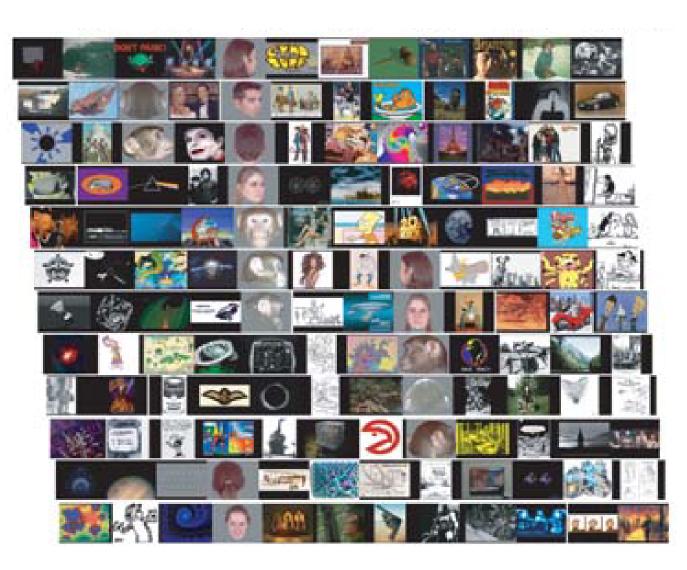
1000s of images (> 15% faces) presented to neuron for 55 or 110ms

RSVP: Face-selective neurons



- Some monkey STSa neurons show clear preferences
 top 50 faces are images
- Response low to remaining patterns

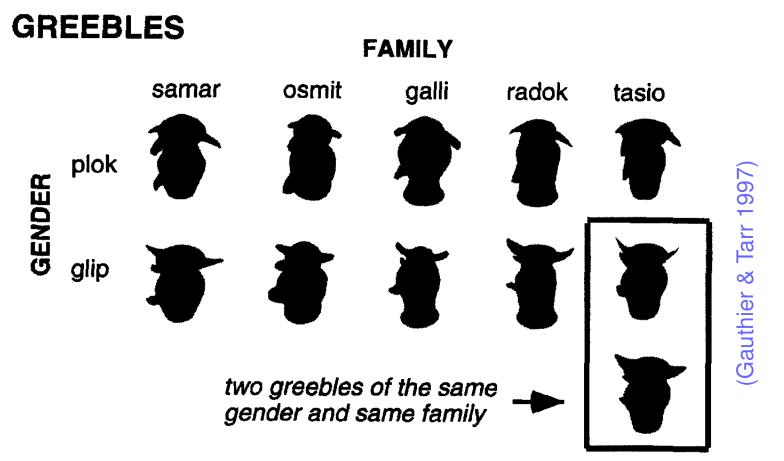
RSVP: Non-face-selective neurons



Other neurons don't make much sense at all

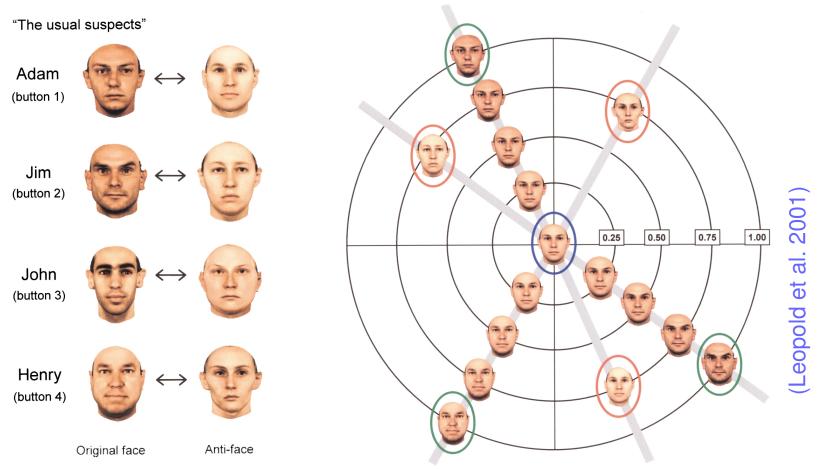
CNV Spring 2008: Extrastriate models

Form expertise



Most of the "specialness" of faces appears to be shared by other object categories requiring configural distinctions between similar examples.

Face aftereffects



Aftereffects are seemingly universal. E.g. face aftereffects: changes in identity judgments; blur/sharpness aftereffects, contrast aftereffects...

Invariant tuning

Higher level ventral stream cells have response properties invariant to size, viewpoint, orientation, etc.

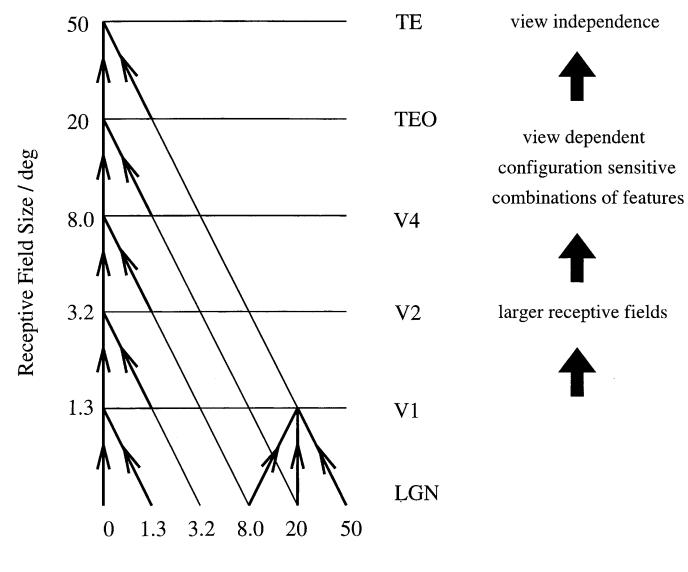
Similar to complex cells, but higher-order. E.g. can respond to face regardless of its location and across a wide range of sizes and viewpoints.

Why is invariance hard?



Simple template-based models won't provide much invariance, but could build out of many such cells.

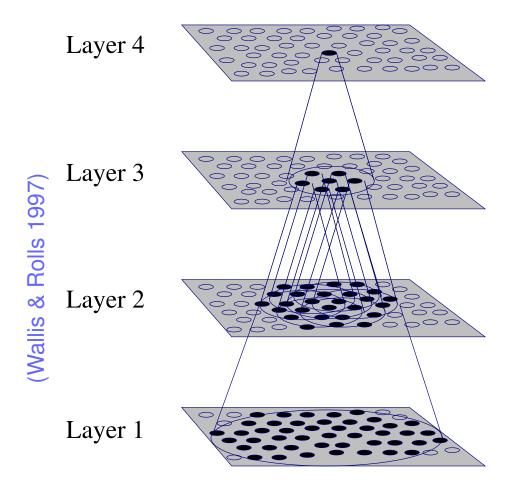
RF sizes



Eccentricity / deg

(Rolls 1992)

VisNet



Develops neurons with invariant tuning

Assumes fixed V1 area

Ignores topographic organization

Trace learning rule

VisNet uses the trace learning rule proposed by Földiák (1991). Based on Hebbian rule for activity y^{τ} and input x_j^{τ} :

$$\Delta w_j = \alpha y^{\tau} x_j^{\tau} \tag{1}$$

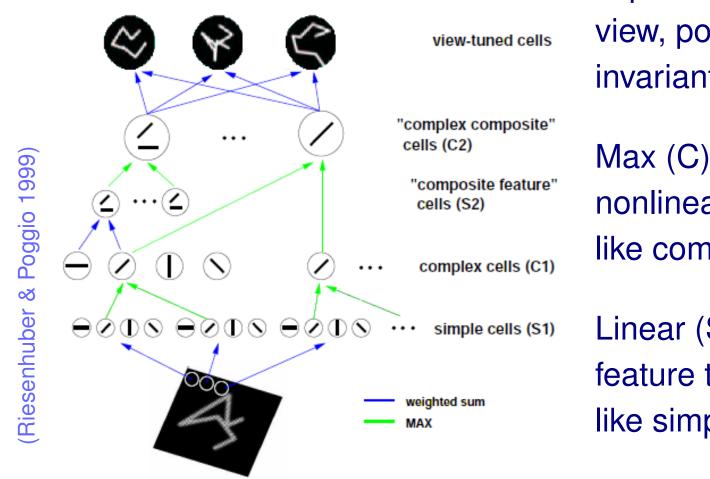
but modified to use recent history ("trace") of activity:

$$\Delta w_j = \alpha \bar{y}^{\tau} x_j^{\tau} \tag{2}$$

$$\bar{y} = (1 - \eta)y^{\tau} + \eta \bar{y}^{\tau - 1}$$
 (3)

General technique for invariant responses?

HMAX



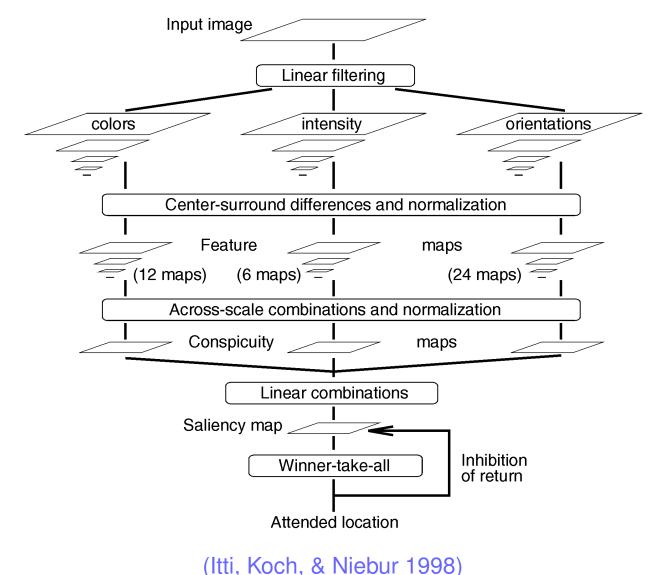
Top level (only) learns view, position, size invariant recognition

Max (C) units: nonlinear pooling, like complex cells

Linear (S) units: feature templates, like simple cells

No clear topography

Koch and Itti saliency maps



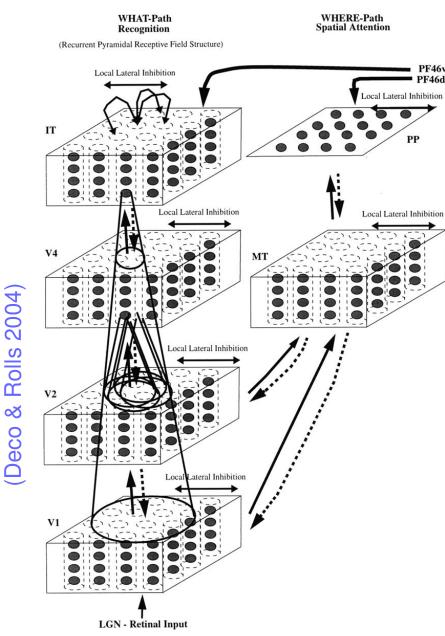
Attention model: most salient feature attended

Various feature maps pooled at different scales

Single winner: attended location

Inhibition of return: enables scanning

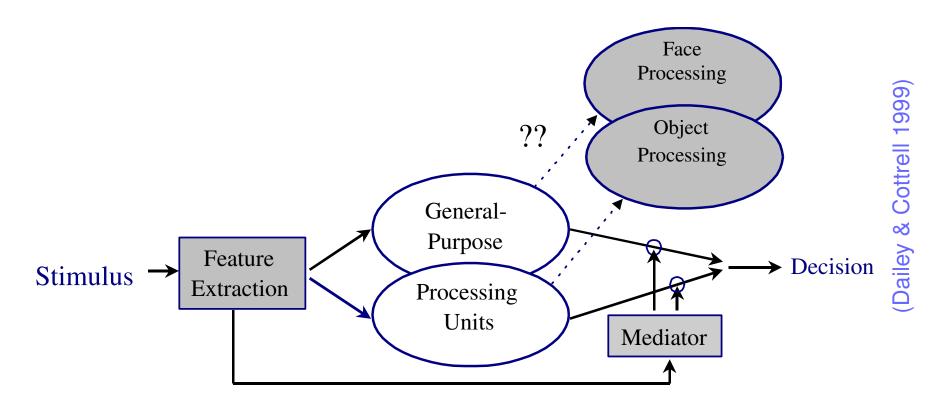
Other attention models



There are a number of other models of behavior like attention, most quite complex Hard to tie individual model areas to specific experimental results from those areas

Also need to include superior colliculus

Modeling separate streams



Slight biases are sufficient to make one stream end up selective for faces, the other for objects

More complexities

Need to include eye movements, fovea/periphery.

At higher levels, neurons become multisensory.

Eventually, realistic models will need to include auditory areas, touch areas, etc.

Feedback from motor areas is also more important at higher levels.

Training data for such models will likely be harder to make than building a robot – will need embodied models.

Summary

- Need to include many areas besides V1
- Complexity and lack of data are serious problems
- Eventually: situated, embodied models
- May be useful to focus on species with just V1 or a few areas before trying to tackle whole visual hierarchy
- Lots of work to do

References

- Britten, K. H., Shadlen, M. N., Newsome, W. T., & Movshon, J. A. (1992).
 The analysis of visual motion: A comparison of neuronal and psychophysical performance. *The Journal of Neuroscience*, *12*, 4745– 4765.
- Bruce, C., Desimone, R., & Gross, C. G. (1981). Visual properties of neurons in a polysensory area in superior temporal sulcus of the macaque. *Journal of Neurophysiology*, *46* (2), 369–384.
- Dailey, M. N., & Cottrell, G. W. (1999). Organization of face and object recognition in modular neural network models. *Neural Networks*, *12* (7), 1053–1074.

Deco, G., & Rolls, E. T. (2004). A neurodynamical cortical model of visual attention and invariant object recognition. *Vision Research*, *44* (6), 621–642.

Földiák, P. (1991). Learning invariance from transformation sequences. *Neural Computation*, *3*, 194–200.

Földiák, P., Xiao, D., Keysers, C., Edwards, R., & Perrett, D. I. (2004).
Rapid serial visual presentation for the determination of neural selectivity in area STSa. *Progress in Brain Research*, *144*, 107–116.

Gauthier, I., & Tarr, M. J. (1997). Becoming a 'Greeble' expert: Exploring mechanisms for face recognition. *Vision Research*, *37* (12), 1673– 1682. Itti, L., Koch, C., & Niebur, E. (1998). A model of saliency-based visual attention for rapid scene analysis. *IEEE Transactions on Pattern Analysis and Machine Intelligence*, *20* (11), 1254–1259.

Leopold, D. A., O'Toole, A. J., Vetter, T., & Blanz, V. (2001). Prototypereferenced shape encoding revealed by high-level aftereffects. *Nature Neuroscience*, *4* (1), 89–94.

Riesenhuber, M., & Poggio, T. (1999). Hierarchical models of object recognition in cortex. *Nature Neuroscience*, *2* (11), 1019–1025.

Rolls, E. T. (1992). Neurophysiological mechanisms underlying face processing within and beyond the temporal cortical visual areas.

Philosophical Transactions: Biological Sciences, *335* (1273), 11–21.

Salzman, C. D., Britten, K. H., & Newsome, W. T. (1990). Cortical microstimulation influences perceptual judgements of motion direction. *Nature*, *346*, 174–177, Erratum 346:589.

Shmuel, A., & Grinvald, A. (1996). Functional organization for direction of motion and its relationship to orientation maps in cat area 18. *The Journal of Neuroscience*, *16*, 6945–6964.

Ungerleider, L. G., & Mishkin, M. (1982). Two cortical visual systems. In Ingle, D. J., Goodale, M. A., & Mansfield, R. J. W. (Eds.), *Analysis* of Visual Behavior (pp. 549–586). Cambridge, MA: MIT Press. Van Essen, D. C., Anderson, C. H., & Felleman, D. J. (1992). Information processing in the primate visual system: An integrated systems perspective. *Science*, 255, 419–423.

- Wallis, G. M., & Rolls, E. T. (1997). Invariant face and object recognition in the visual system. *Progress in Neurobiology*, *51* (2), 167–194.
- Xiao, Y., Casti, A., Xiao, J., & Kaplan, E. (2007). Hue maps in primate striate cortex. *Neuroimage*, *35* (2), 771–786.
- Xiao, Y., Wang, Y., & Felleman, D. J. (2003). A spatially organized representation of color in macaque cortical area V2. *Nature*, *421*, 535–539.

 Xu, X., Collins, C. E., Khaytin, I., Kaas, J. H., & Casagrande, V. A. (2006). Unequal representation of cardinal vs. oblique orientations in the middle temporal visual area. *Proceedings of the National Academy* of Sciences of the USA, 103 (46), 17490–17495.