

Visual development in babies and infants

Marko Nardini

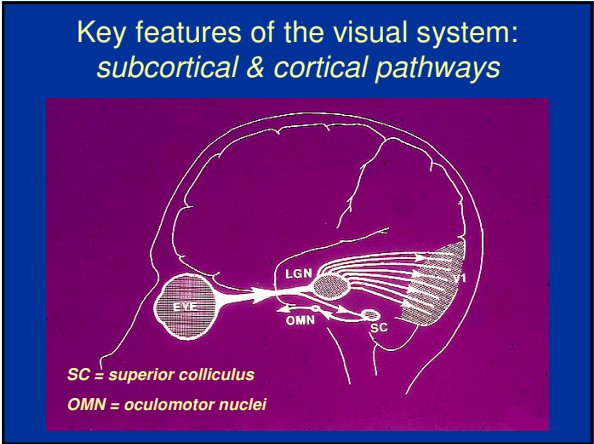
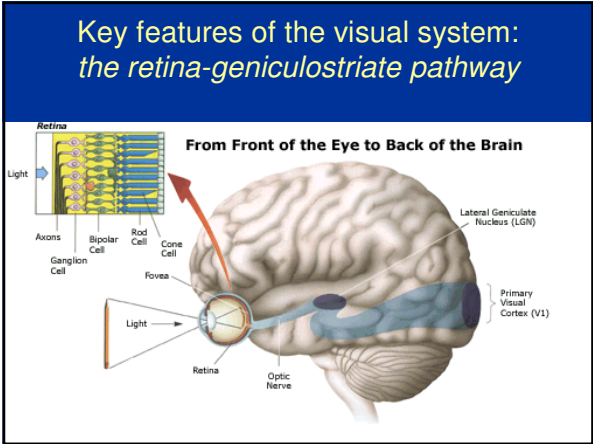
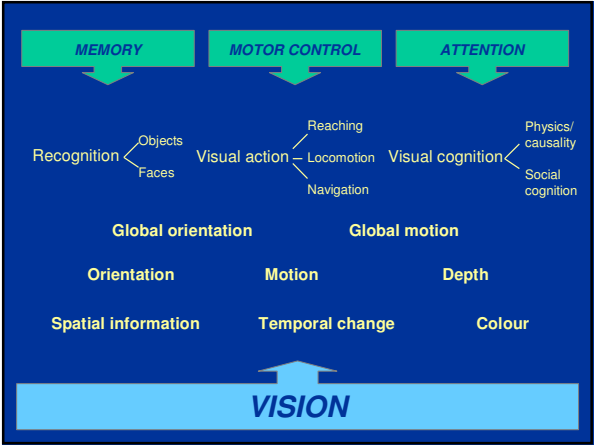
UCL Institute of Ophthalmology

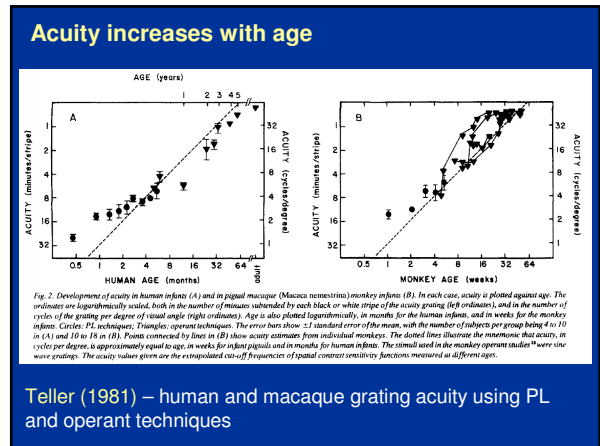
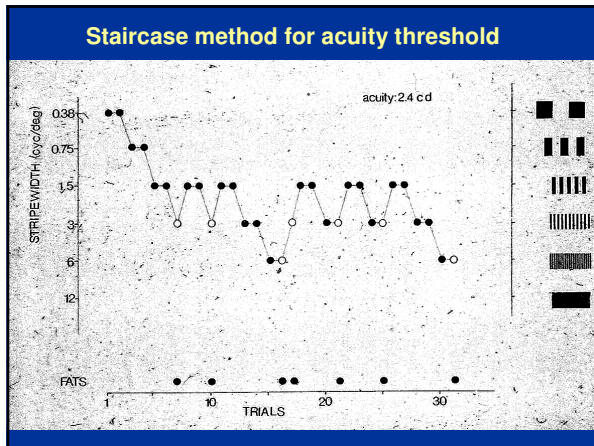


Vision

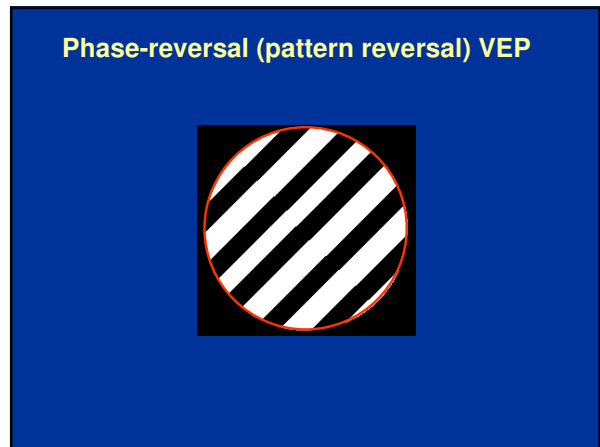
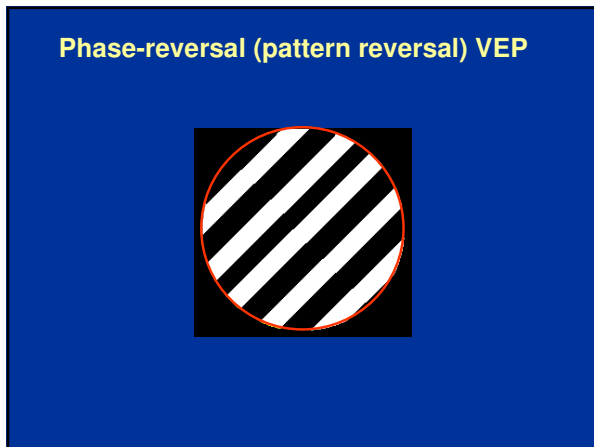
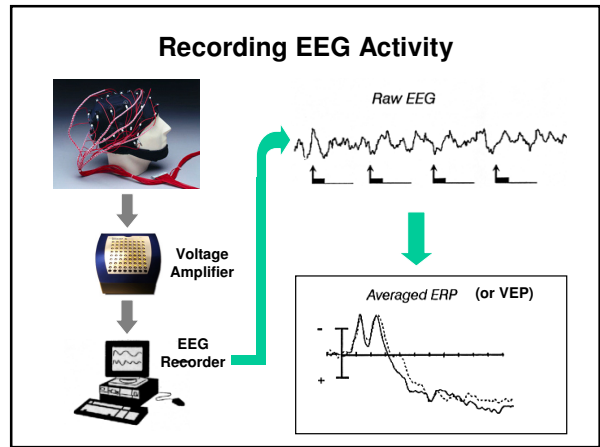
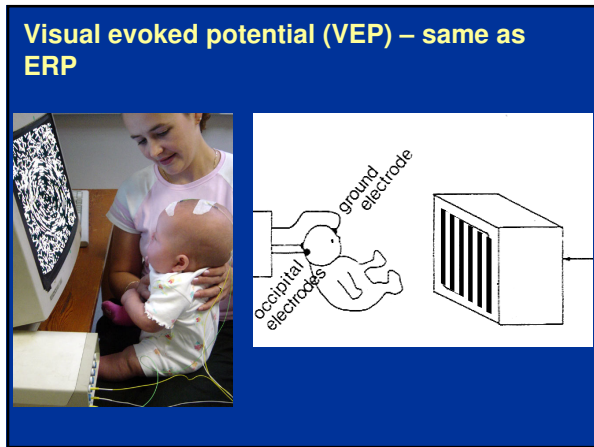
- a major function of the primate brain
- vision develops rapidly in early life and serves as a base for development of action, cognition, communication, social interactions

Figure 19 Much of V1 is located in the calcarine sulci and its relationship to other brain areas is best shown by unfolding the brain and showing it flattened open. The visually responsive areas of the macaque monkey are shown in color. From Van Essen et al. (1992).





Teller (1981) – human and macaque grating acuity using PL and operant techniques



Phase-reversal (pattern-reversal) visual evoked potential (VEP)

amplification
signal averaging, time-locked to phase-reversals

Statistically significant VEP to stripe reversal shows input activity to cortex – though not necessarily cortical function

a

b

a b a b a b a b

Phase-reversing grating (2-10 cycles/sec)

"Sweep VEP"

Regan (1977)

Temporally modulated pattern

5Hz or 10Hz

Frequency is systematically changed (swept) over a large range.

Can measure amplitude of VEP response as a function of frequency, and extrapolate the highest frequency that is processed (i.e. acuity) from this

Fig 7. The mean sweep VEP acuity with 95% confidence bands in 1 month (lunar) increments for the date of Fig. 6

Norcia & Tyler (1985)

Indicates 2-3x better acuity at 1 mo. than PL – but only that there is input to cortex, not necessarily cortical processing or perception

Acuity increases with age –why?

Limits on developing visual acuity

- **Optical blur**
 - clarity of media
 - refraction
- **Receptor density & efficiency**
differentiation of the fovea
- **Neural development**

Limits on developing visual acuity: Optical blur

- clarity of media
- refraction & accommodation

Not generally the limiting factors on infant acuity

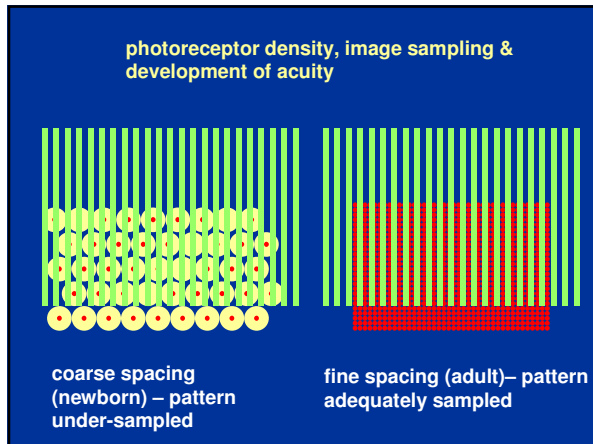
Limits on developing visual acuity: Receptor density & efficiency

- OS = outer segment; contains photosensitive pigment
- short OS = inefficient at detecting light
- fat inner segment (IS) = cones aren't tightly packed = poor spatial sampling of the image
- Development of long fibre = cones displaced to allow dense packing in foveal pit

Limits on developing visual acuity: Receptor density & efficiency

FIGURE 4.0 Central fovea of the human retina. (From Polak, 1957.)

Maximum acuity provided by fovea – displaced cell bodies allow dense packing and minimal obstruction of light to cone outer segments

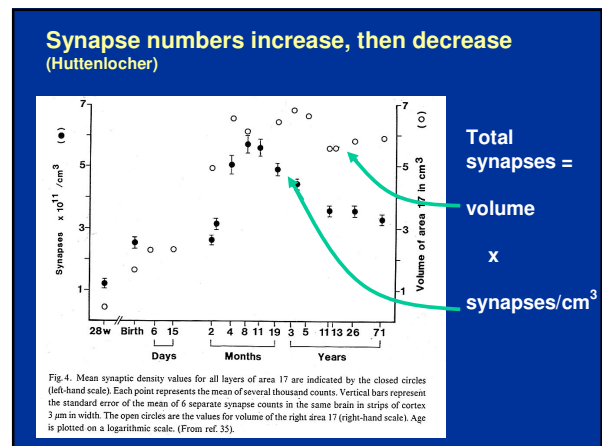


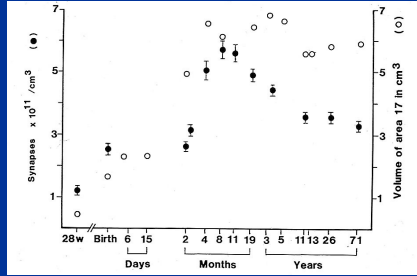
- Banks & Bennett (1988)
- Sampling argument combined with poor efficiency due to short outer segments
 - Calculated in comparison with adult and 'ideal observer' model
 - May account for overall change, but
 - both adult and infant fall far short of ideal observer – little idea of factors
 - poor account of acuity changes during infancy, especially first 3 months
 - Other, more central, changes are going on

- Limits on developing visual acuity: Neural development
- myelination of visual pathways
 - what are functional consequences?
 - development and distribution of cortical neurons
 - developing connectivity in cortex (and elsewhere)
 - increasingly complex dendritic and axonal processes

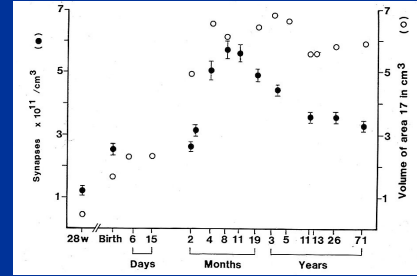
- Development and distribution of cortical neurons
- cell proliferation
 - cell migration
 - cell differentiation into different structural types
- All three processes are complete before birth

- Although all cells are born before birth, the mass of the brain increases postnatally from 350 g – 1350 g (approx x 4)
- This increase must include
- myelin
 - fibres and synapses associated with increased connectivity
- Connectivity determines function





Processes of
 (a) growth of dendrites and synaptic terminals
 (b) selective pruning of connections



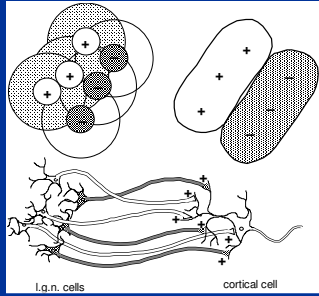
Synaptic increase is seen everywhere in cortex.
 What are its implications for visual function?

What are all those synapses doing?

Connectivity determines receptive field structure – and therefore function

e.g.

orientation selectivity



Hubel & Wiesel 1960's model – probably too simple – intracortical connections are important also!

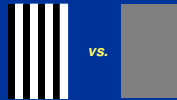
Spatial information: Summary

- Visual acuity shows very rapid development in first few months of life, then slower development towards adult levels by 3-4 years
- Underlying changes in photoreceptor organization, neural connectivity and myelination

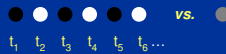
Temporal change

Changes over time provide a basis for detecting **movement** in the visual field

- Change in space (at a single time) – basic measure: *acuity*



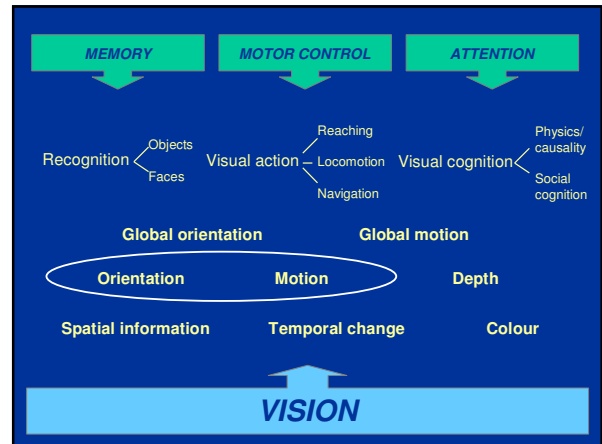
- Change in time (at a single location) – basic measure: *critical fusion frequency (CFF)*



- Behaviourally (pref looking), infants' critical fusion frequency was strikingly mature (Regal 1981): 40Hz (75% of adult) at 4 weeks; indistinguishable from adult at 12 weeks.
- Much more mature than acuity. Makes sense as electroretinograms (ERGs) show newborn cones to respond at up to 75Hz – so this is not a constraint.
- However, EEG measures of flicker information reaching visual cortex show much lower critical frequency. (Apkarian 1993, Morrone et al 1996).
- May be that latencies are long and variable due to incomplete myelination, leading to out-of-phase signals in cortex that (1) do not give coherent EEG, but (2) could still drive a behavioural response - see Atkinson & Braddick OVS 2009

Review

- We have seen how sensitivity to **spatial and temporal changes** in luminance (and wavelength / colour) develops in the first few months of life
- These provide the building blocks for detecting the **orientation, motion** and **depth** of visual patterns
- This requires increasingly sophisticated neural information processing – dependent on cortex (V1)



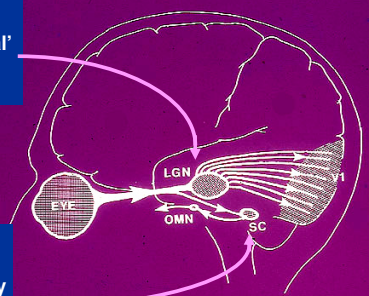
ORIENTATION processing



Two kinds of 'subcortical'..

LGN is 'precortical'
– on the route to
striate cortex

Superior colliculus
is part of a distinct
subcortical pathway
(but also
interconnected with
cortex)



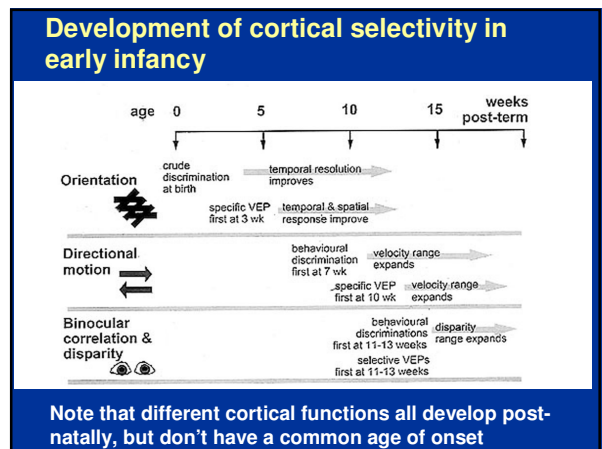
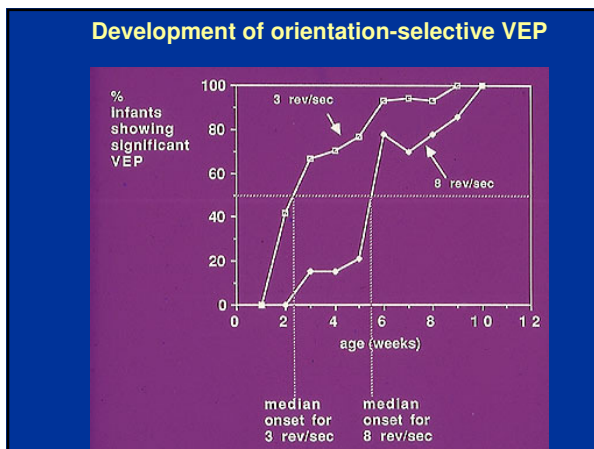
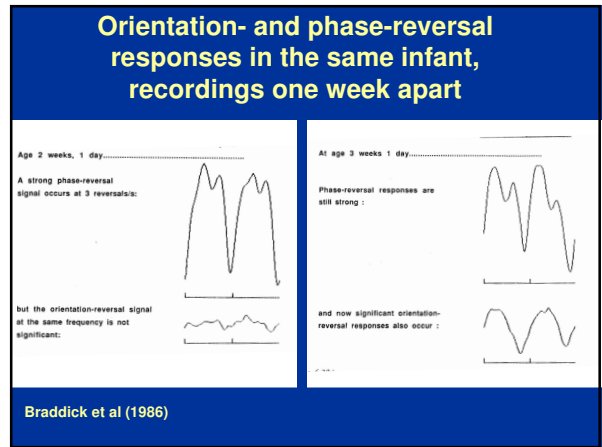
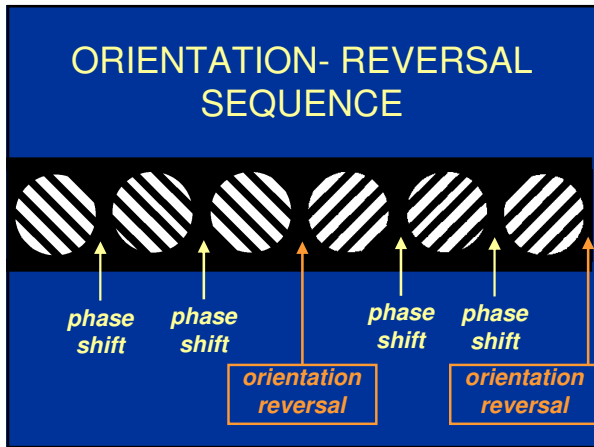
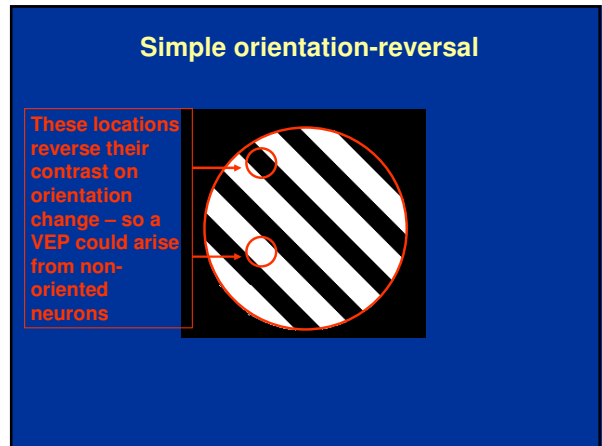
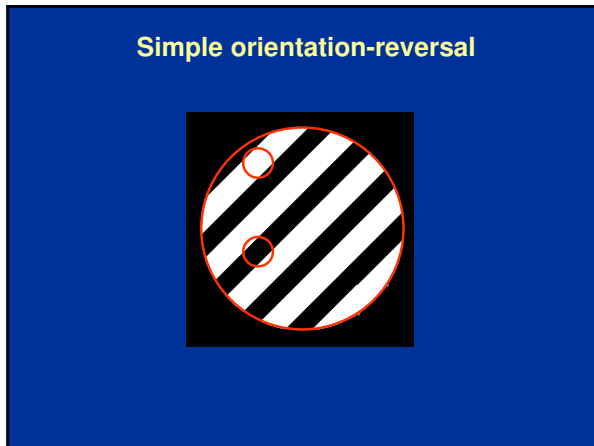
*Neither LGN or SC show orientation
selectivity independent of cortex*

cortical neurons – and not
precortical - show selective
sensitivity to :

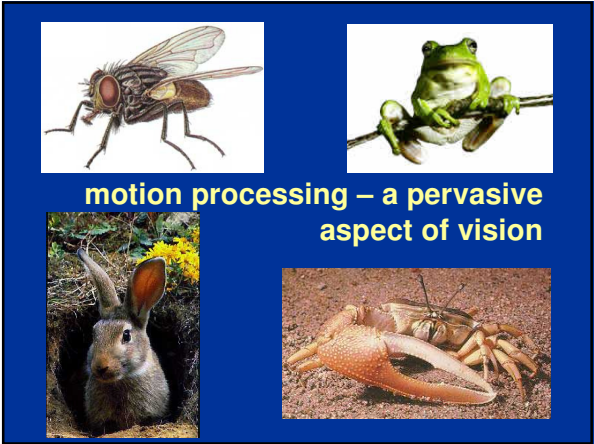
- **orientation**
- **direction of motion**
- **binocular disparity (stereopsis)**

Development of orientation selective cortical
neurons – orientation-reversal VEP



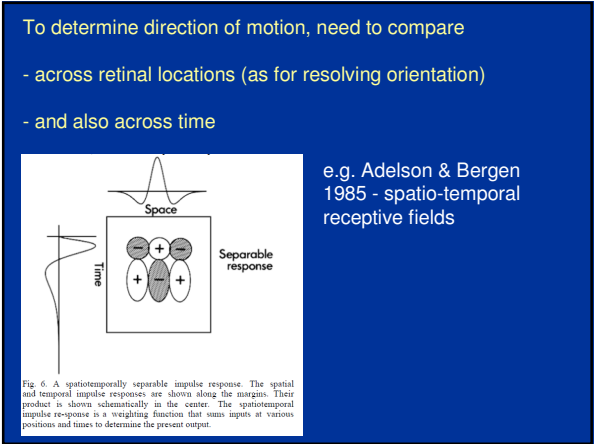
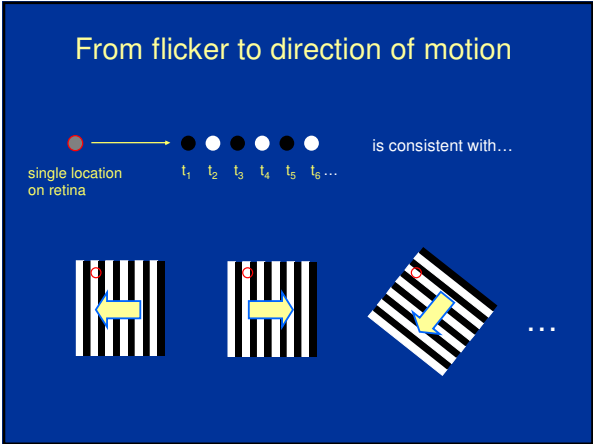


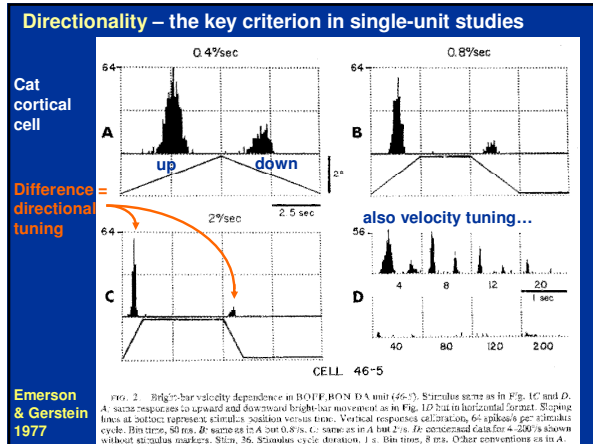
MOTION processing



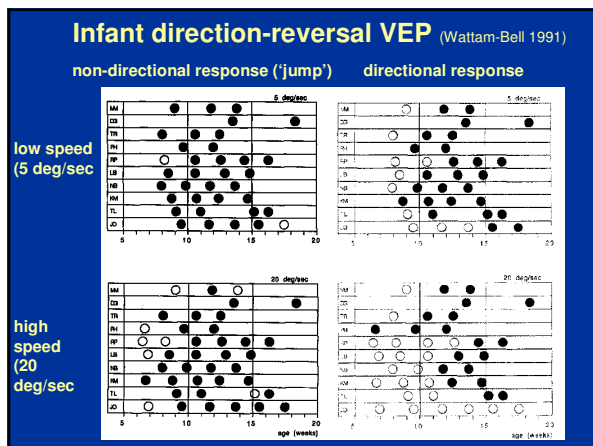
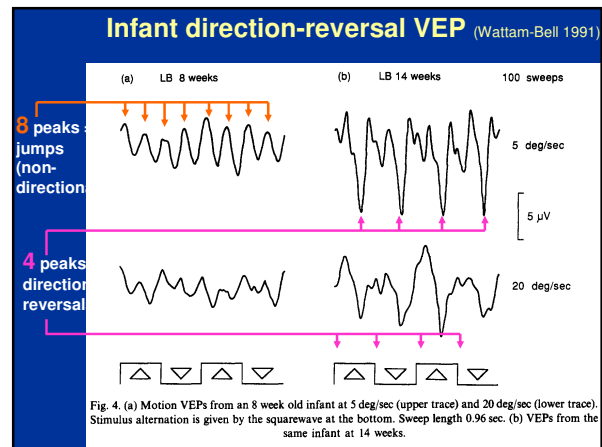
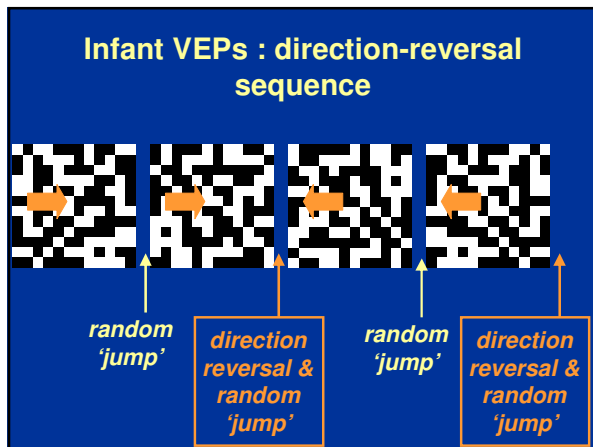
Volkman & Dobson (1976) – 2 month infants preferential looking:
moving > static stimuli

Does this mean that young infants process visual motion?
Consider:
Infants also show high flicker sensitivity in preferential looking (Regal, 1981)
early stages of visual pathway (e.g. retinal ganglion cells) respond to temporal change but not to direction of motion



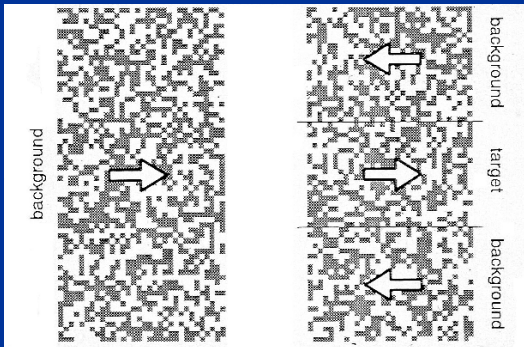


- ### Directional sensitivity in infants
- evoked potential approach
 - preferential looking approach

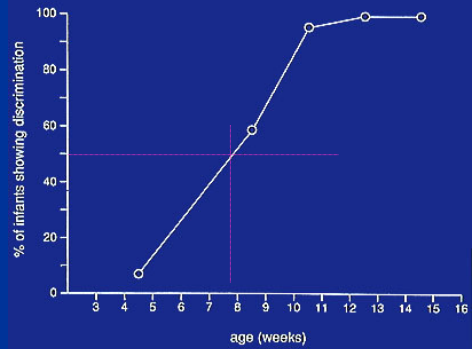


- ### Infant direction-reversal VEP
- first seen about 10-11 weeks on average
 - develops later for higher speeds
 - direct comparison shows that it develops later than orientation-specific responses (Braddick et al., 2005)

Preferential looking for directional motion

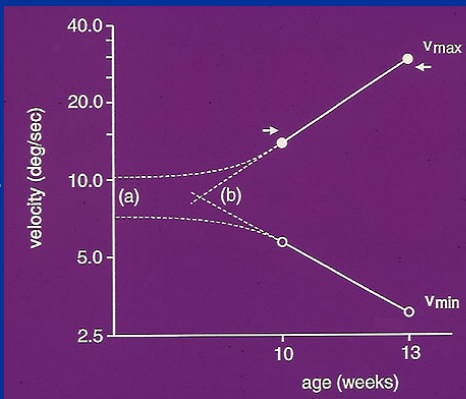


Preferential looking for directional motion



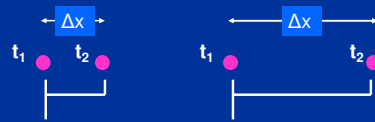
Preferential looking for directional motion

with age, sensitivity extends to both higher and lower speeds



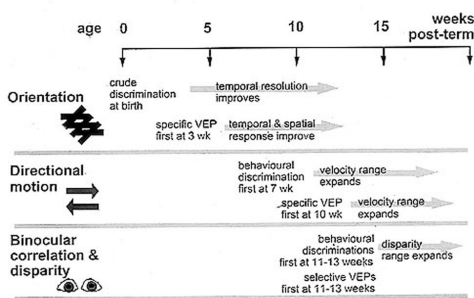
extension to higher velocities

- higher velocities with age - primarily a spatial rather than temporal change
- a 'fine to coarse progression' (not expected from acuity changes)



- extension of horizontal connectivity in cortex?
- compare with extending disparity range for stereopsis

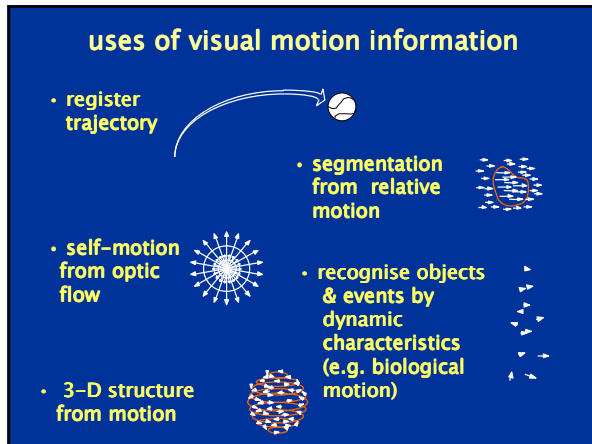
Development of selective filters in early visual processing



All specific functions of primary visual cortex – but they don't have a common onset. Specific aspects of cortical connectivity each have to develop.

Plasticity of motion processing

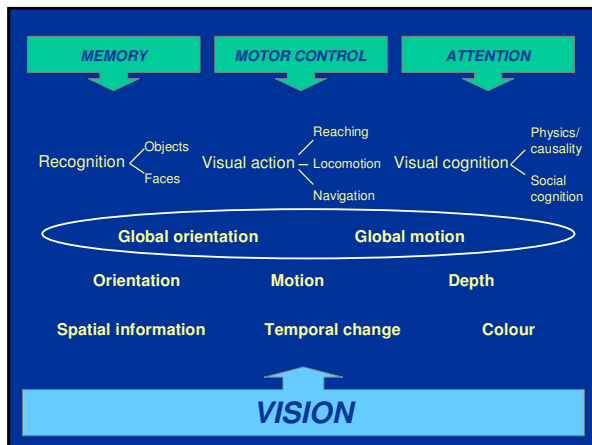
- kittens reared in stroboscopic illumination – absence of directional cells in visual cortex (Cynader, Berman & Hein, 1973; Pasternak et al, 1981)
- kittens reared with directional bias show biased distribution of directional selectivity in cortical cells (Daw & Wyatt, 1976)
- separate periods of directional bias and monocular deprivation – show distinct critical periods for motion sensitivity (1st) and binocularity (2nd) (Daw, Berman & Ariel, 1978)



Use of motion for perceptual tasks by 3-5 month-old infants

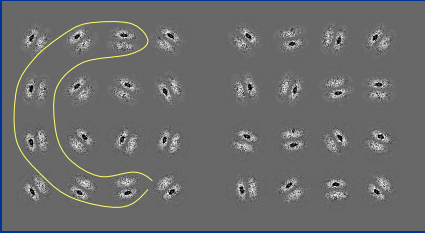
• 3-D structure from motion	Arterberry & Yonas (1988)
• distinguish rigid motion from non-rigid deformation	E J Gibson et al (1979)
• group parts of occluded object by common motion	Kellman & Spelke (1983)
• discriminate motion-defined forms	Kaufman-Hayoz et al (1986)
• discriminate biological motion	Berthenthal et al, 1985

...so directional information can be used for sophisticated perceptual analysis, soon after it first becomes available to the infant



GLOBAL form and motion processing


Coherent organization Random organization




To perceive shapes, we need to look for useful changes in orientation over a large area – e.g. those indicating a **contour**.

Similarly with motion...

Expanding optic flow
(moving forwards)



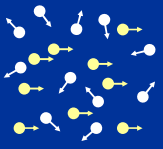
Contracting optic flow
(moving backwards)




The same local motions are present in both cases, but the global organization is very different

To perceive global motion we need to integrate local motions over a large area

macaque V5/MT (dorsal)
 response to coherence level of random dot motion
 (Britten et al, *J Neurosci*, 1992; *Vis Neurosci*, 1993)



macaque V4 (ventral)
 response to concentric or radial configurations
 (Gallant Braun & Van Essen, *Science*, 1993):



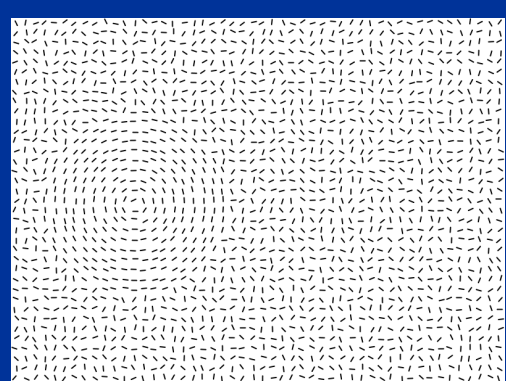
Measure of sensitivity to global form or motion: **coherence threshold**

What % of elements needs to be coherently organised in order for the global organisation to be perceived?

i.e.

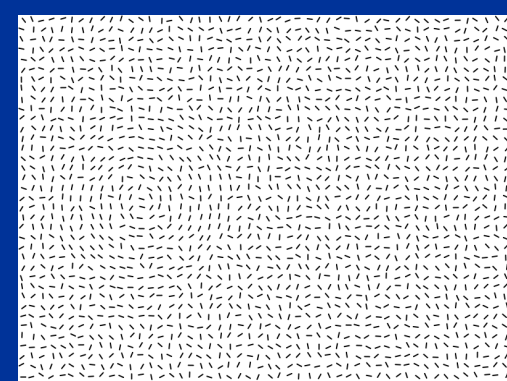
Lowest % at which global organisation is detected = observer's **coherence threshold**

Form coherence



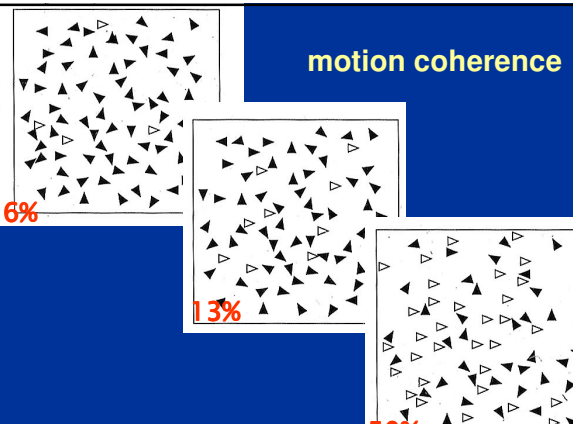
Coherence = 100%

Form coherence



coherence = 60%

motion coherence



6%

13%

50%

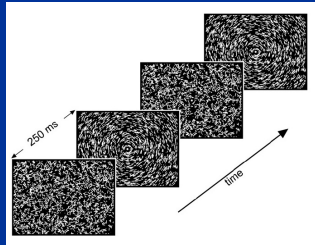
Development of global form and motion processing

Preferential looking and VEP measures:

- *Local* form emerges earlier than local motion
- Evidence for sensitivity to global motion and global form by 4-6 months (Braddick & Atkinson, 2007)
- Global form thresholds reach adult levels *later* than global motion (Gunn et al, 2002)

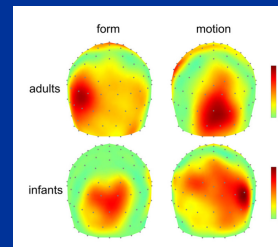
Extra-striate visual areas in development of global form and motion processing

Wattam-Bell et al (2010) – high density ERP recording with global form and motion stimuli in 5-month-olds and adults



Form stimulus.

Motion stimulus is identical, except that line segments represent motion of dots



Like adults, infants show distinct patterns of activity for global form vs. motion

However, topography of responses is very different for infants and adults

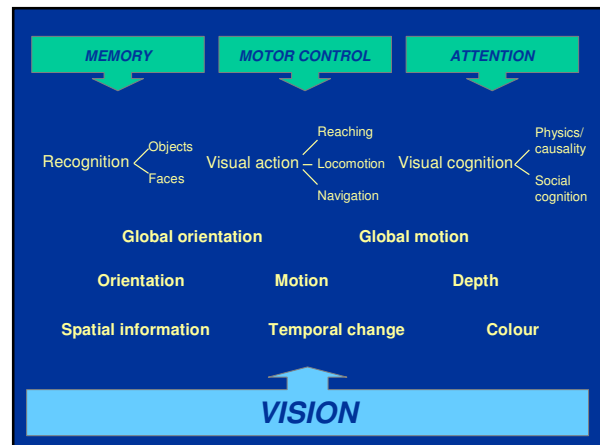
Implies major re-organization of extra-striate visual processing in development.

Global motion from V5 in infants, but dominated by V3/V3A and V6 in adults?

Wattam-Bell et al (2010)

SUMMARY:

Visual functions



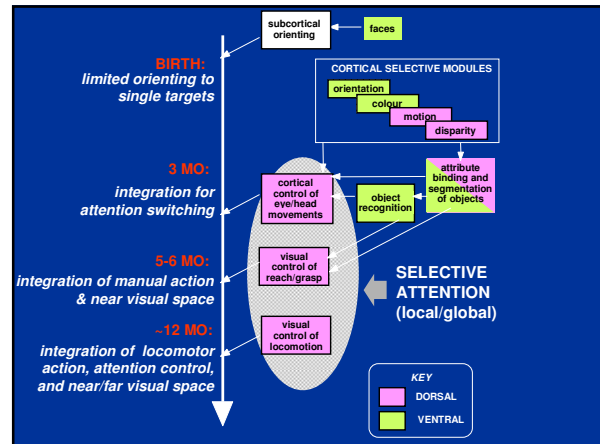
SUMMARY:

Research methods

- Forced-choice preferential looking
- Visual evoked potentials (VEP), aka Event-related potentials (ERP)

MODEL

- Atkinson & Braddick



END

Reading list (p. 1 of 4)

Overview:

Atkinson, J & Braddick, O (in press). Visual development (Chapter 12). In Zelazo, P.D. (Ed.) *Oxford Handbook of Developmental Psychology*. OUP

Specific studies:

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Wattam-Bell J. (1991) The development of motion-specific cortical responses in infants. *Vision Res* 31:287-297.

Braddick, O, Birtles, D, Wattam-Bell, J & Atkinson, J (2005). Motion- and orientation-specific cortical responses in infancy. *Vision Research* 45: 3169-3179.

Braddick, O, & Atkinson, J (2007). Development of brain mechanisms for visual global processing and object segmentation. In C. von Hofsten & K. Rosander (Eds.), *From action to cognition (Progress in Brain Research, Vol. 164)* Amsterdam: Elsevier.

Gunn, A et al (2002). Dorsal and ventral stream sensitivity in normal development and hemiplegia. *Neuroreport* 13(6): 843-847.

Wattam-Bell, J et al (2010). Reorganization of Global Form and Motion Processing during Human Visual Development. *Current Biology* 20(5): 411-415.