

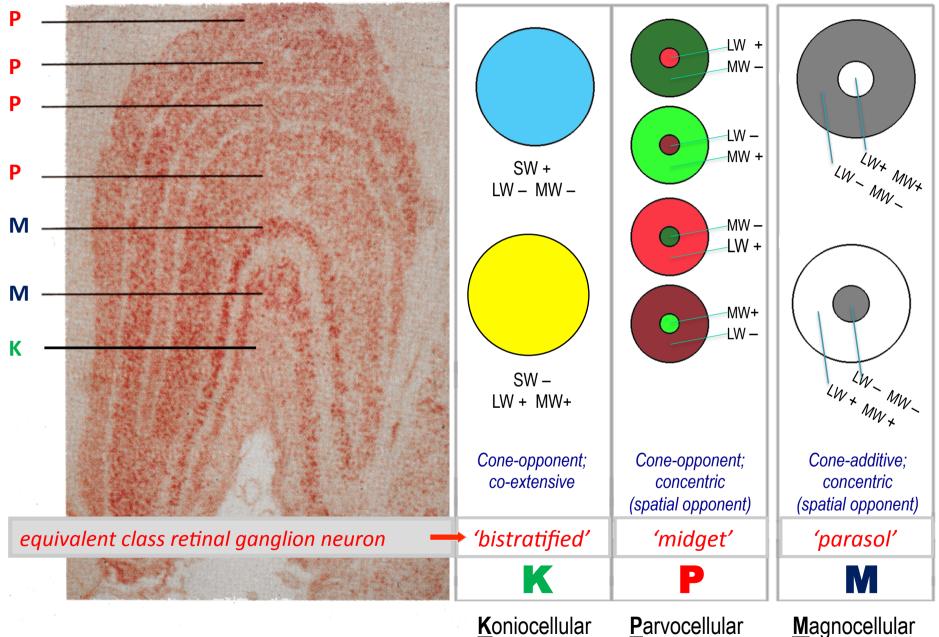
How similar are different cultures' concepts of colour?

Do different languages share equivalent basic colour terms?

# Basic Color Terms: Their Universality and Evolution Berlin & Kay 1969.

- 1 & 2 Light and dark (i.e. 'black' & 'white')
- 3 Red
- 4 & 5 Green & Yellow
- 6 Blue
- 7 Brown
- 8-11 Purple Pink Orange Grey

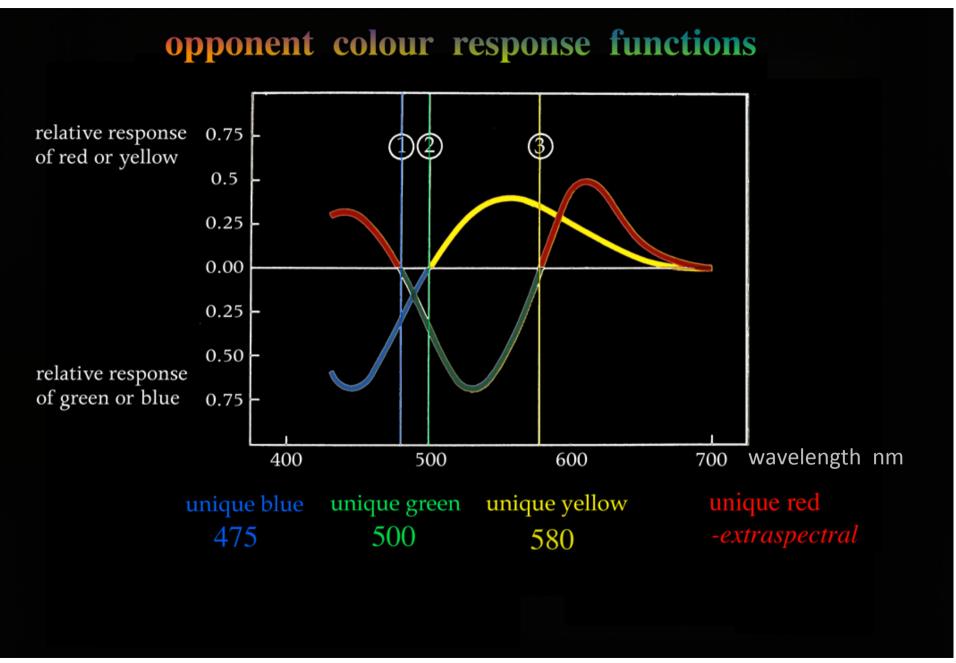
# LGN – receptive field properties of 3 different channels

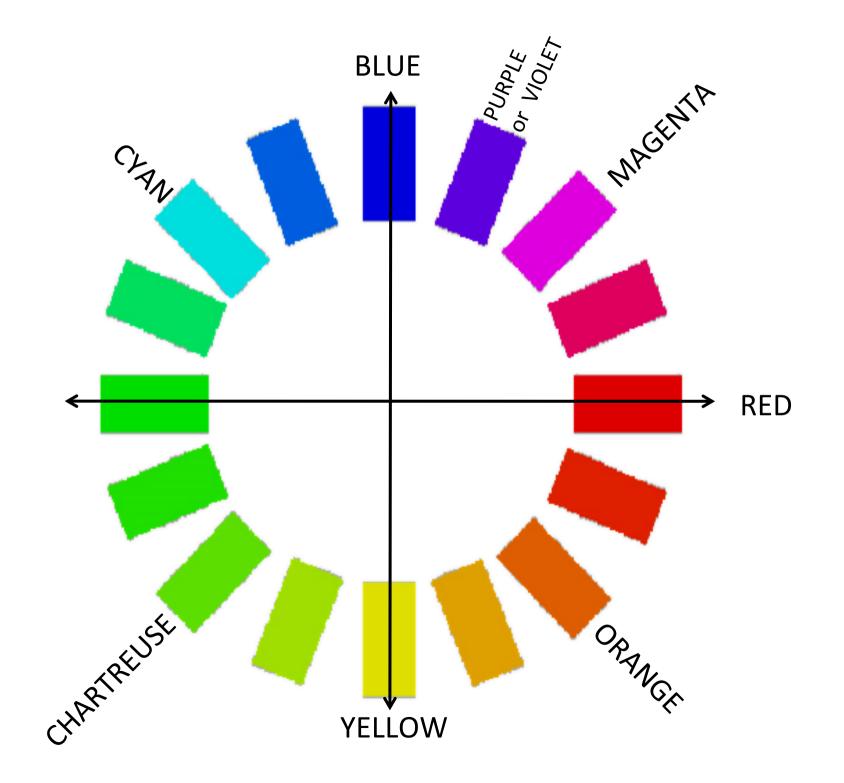


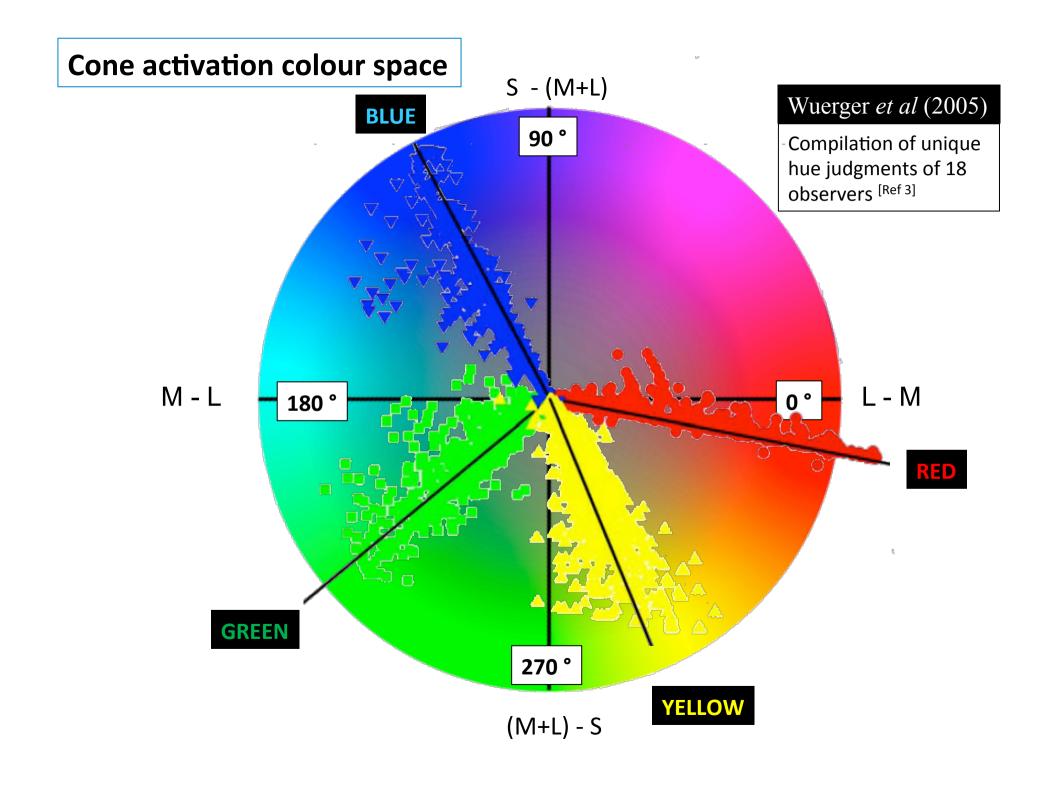
**P**arvocellular

**M**agnocellular

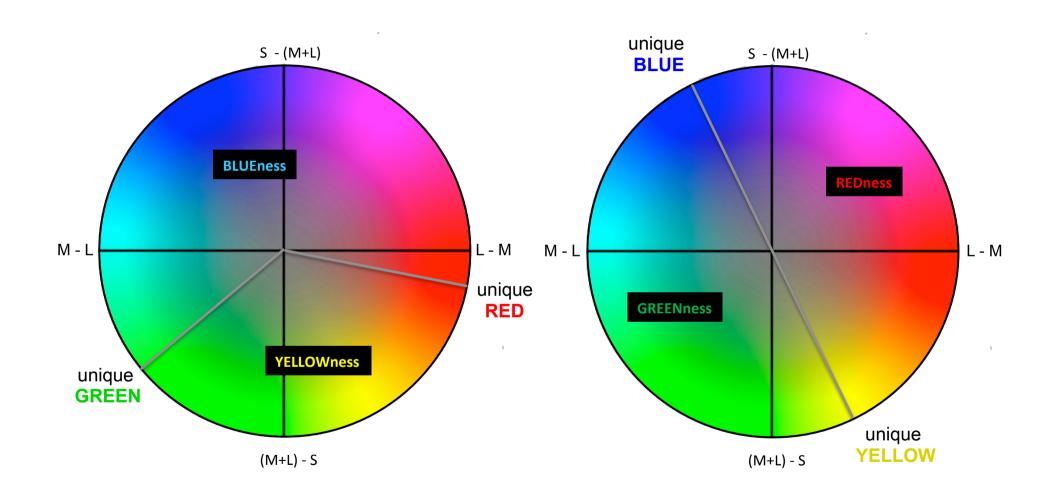
NAMING SEQUENCE		WAVELENGTH	TRADITIONAL SPECTRUM	
Red	R	668	Red	
Reddish-yellow	R Y		Orange	
Yellow Yellowish-green	Y G Y	580	Yellow	- 09 - 1-1-1-1-1-1-1-1-1-1-1-1-1-1-1-1-1-1-1
Green	G	520	Green	
Greenish-blue	G B	500	Green	
Blue	В	464		<u>- 8</u>
Bluish-red	R B	440	Blue	
			Indigo	
			Violet	- 05



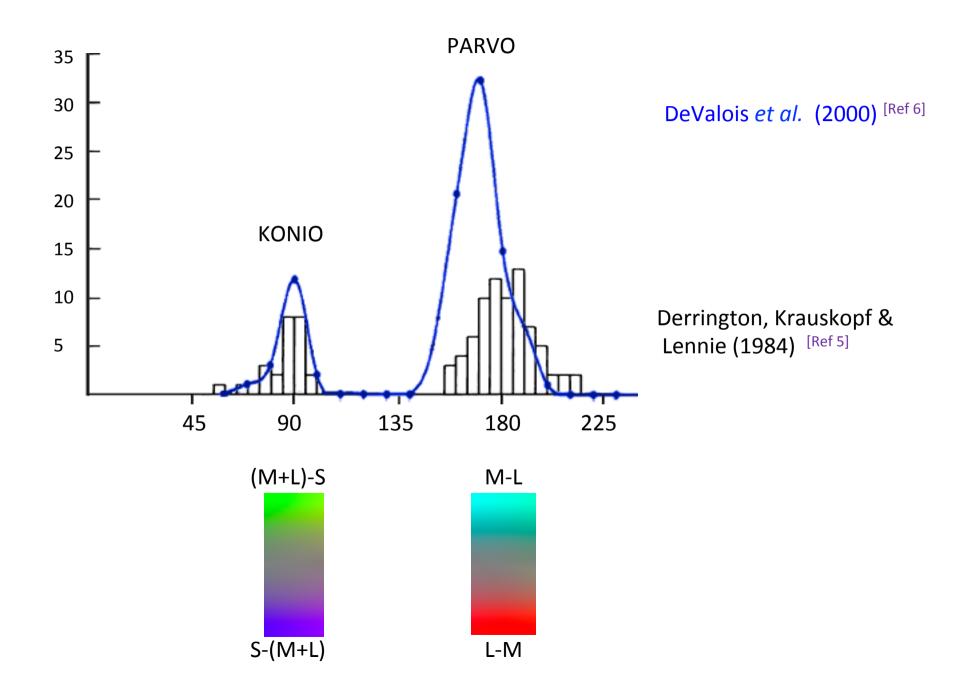




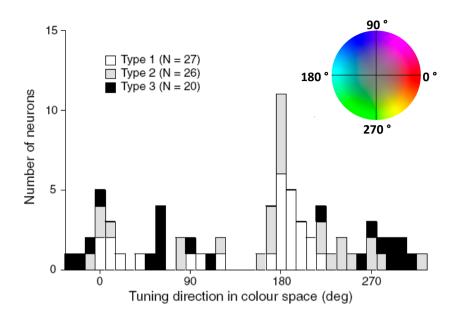
# **Cone activation colour space**

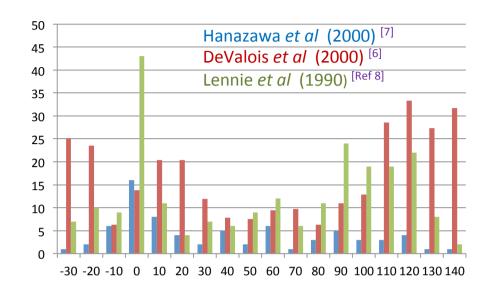


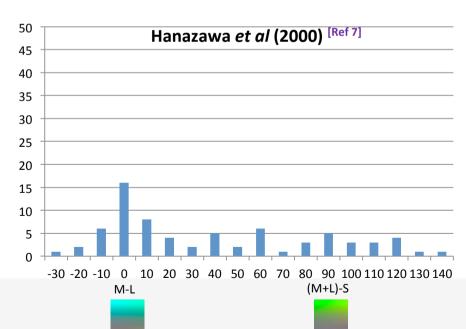
# **LGN** - frequency distribution of chromatic tuning



## **Neural colour axes in V1**

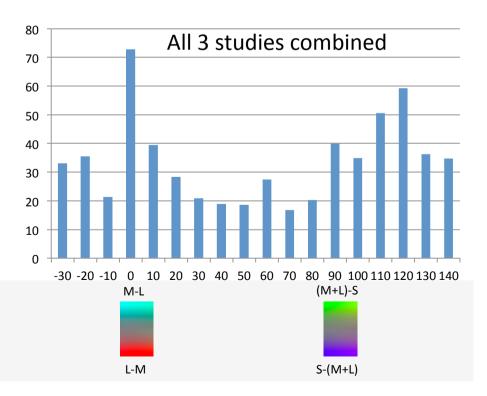




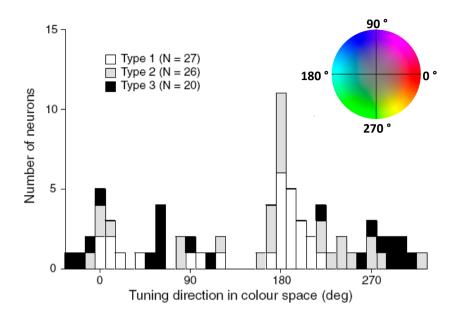


S-(M+L)

L-M

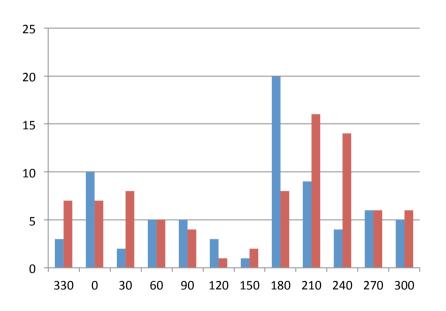


## Neural colour axes in V1 & V2



V1: Hanazawa et al (2000) [Ref 7]

10 deg. histogram (360 scale)



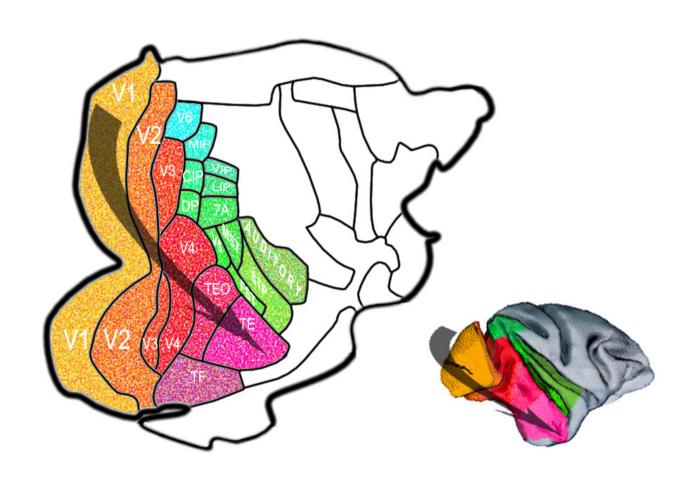
V1: Hanazawa *et al* (2000) [Ref 7] V2: Kiper *et al* (1997) [Ref 9]

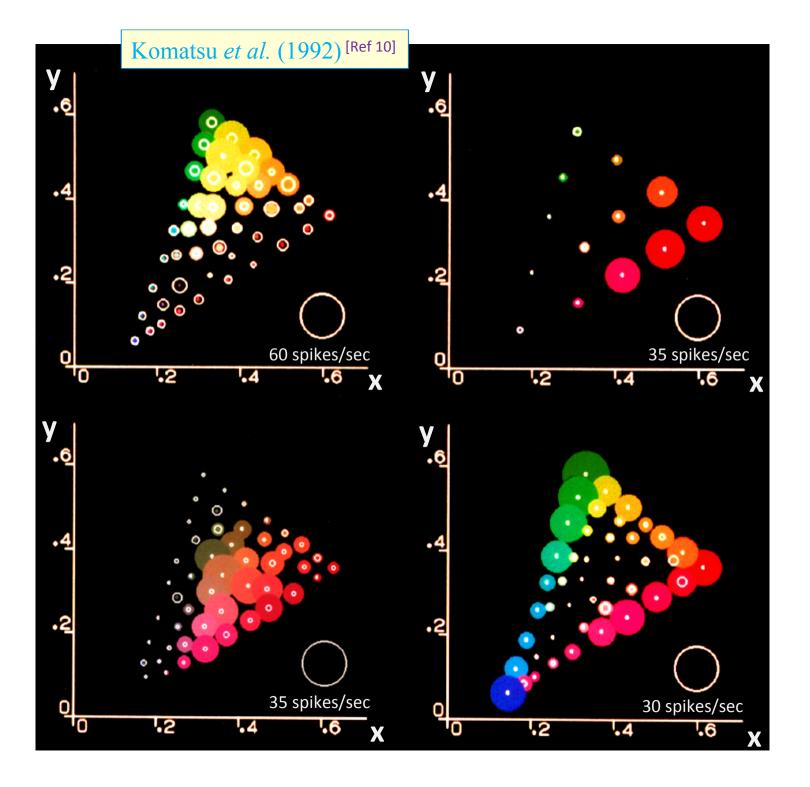
Similar looking?

- but correlation coefficient = 0.29, not significant

30 deg. histogram (360 scale)

And, at the far end of the 'ventral' visual pathway...

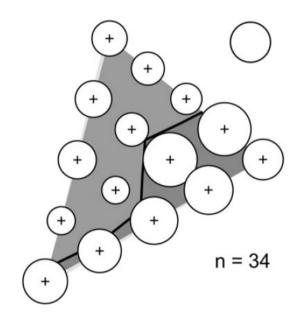




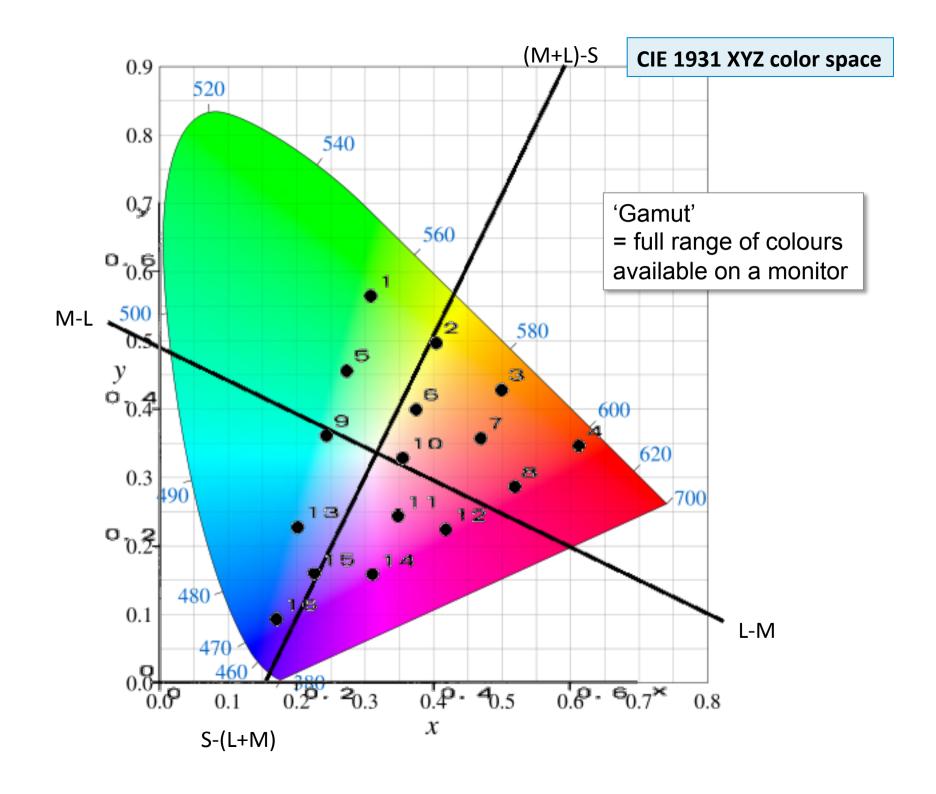
# Komatsu et al (1992)

# .4-

# Yasuda et al (2010)

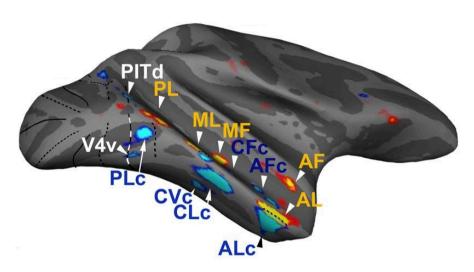


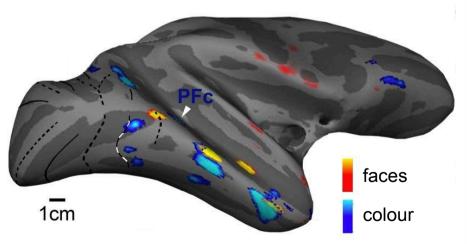
Pooled colour response fields of 19 IT neurons

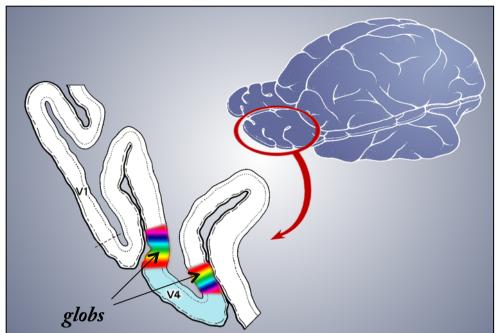


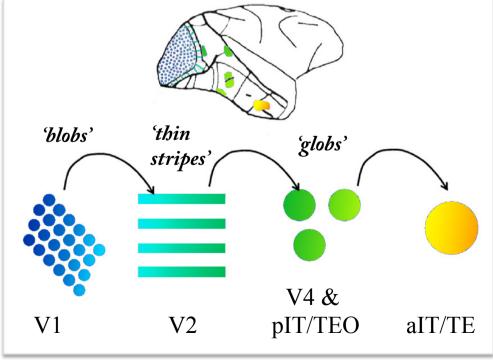
Conway *et al.* (2007) [Ref 12] Lafer-Sousa & Conway (2013) [Ref 13]

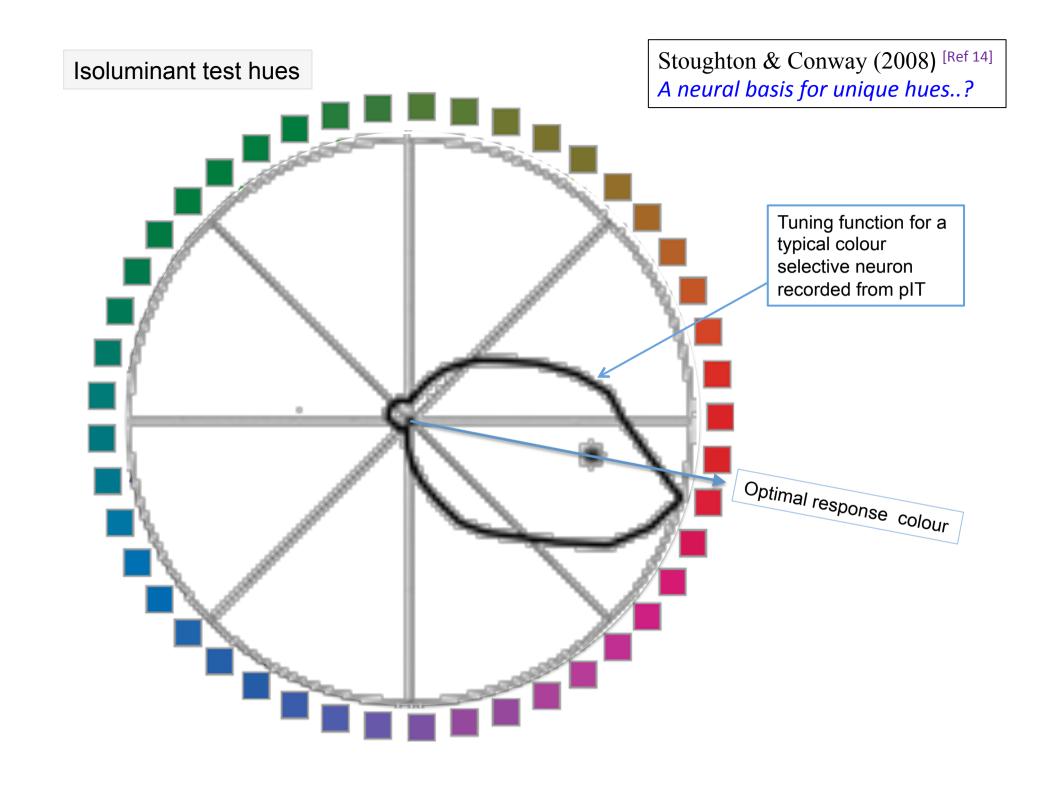
Colour pathway 'globs' revealed by fMRI

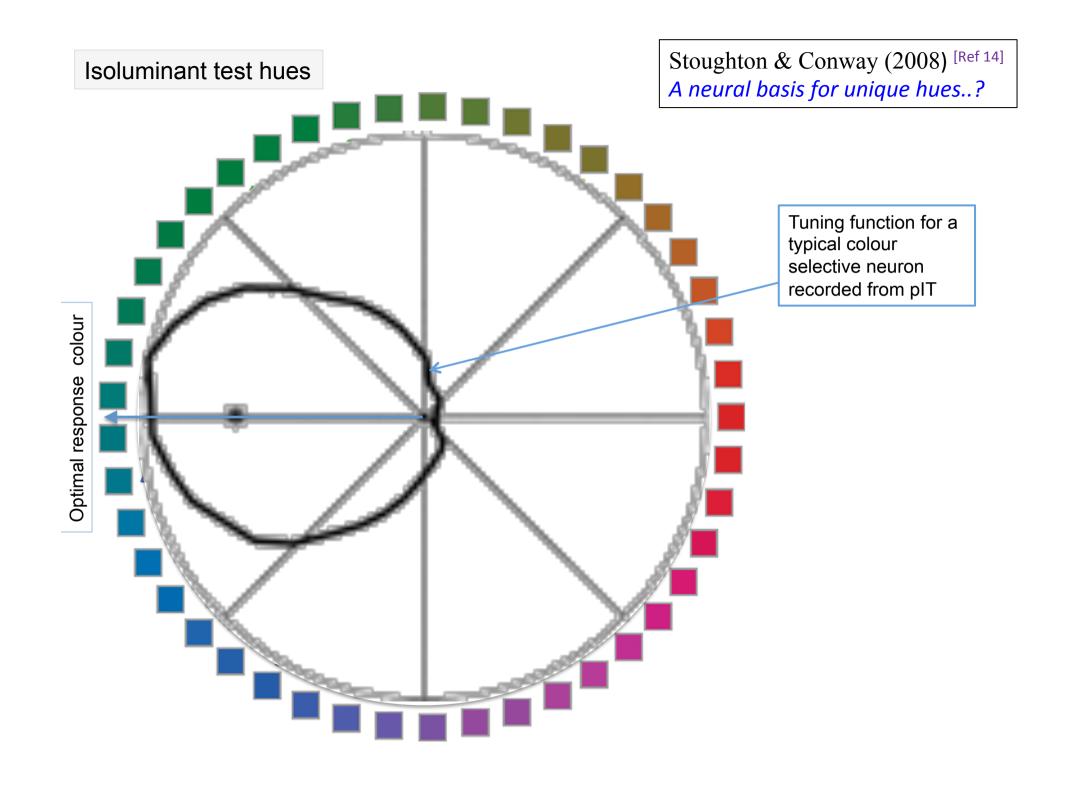


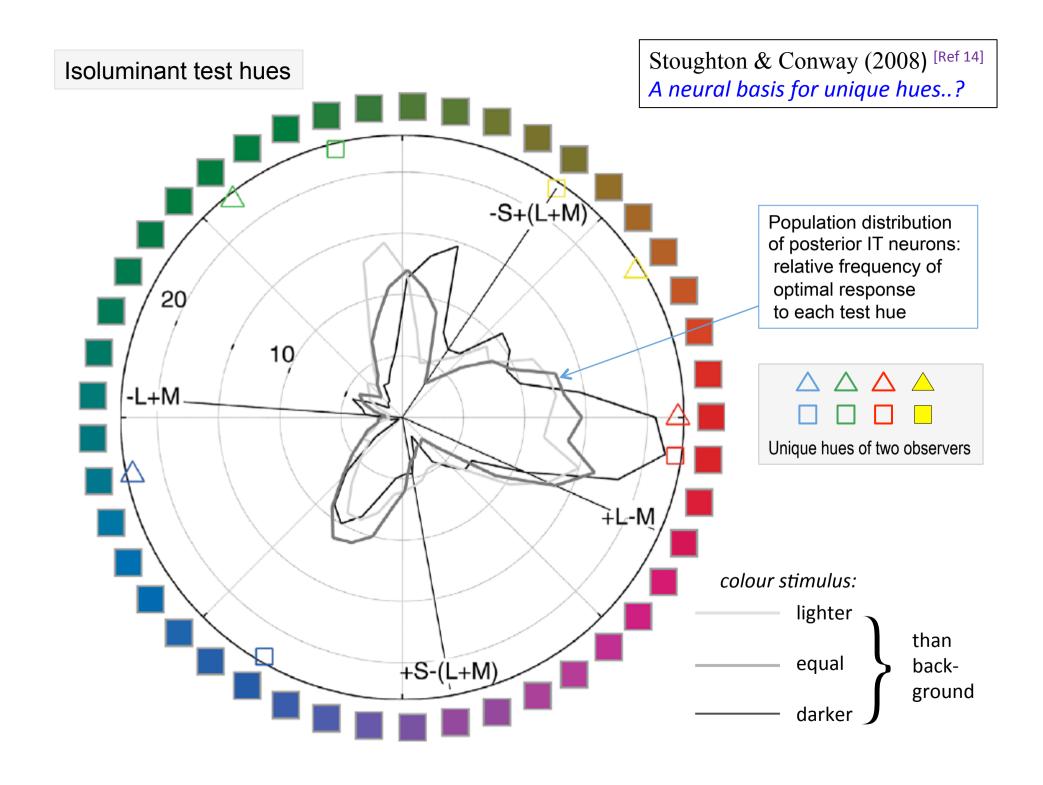








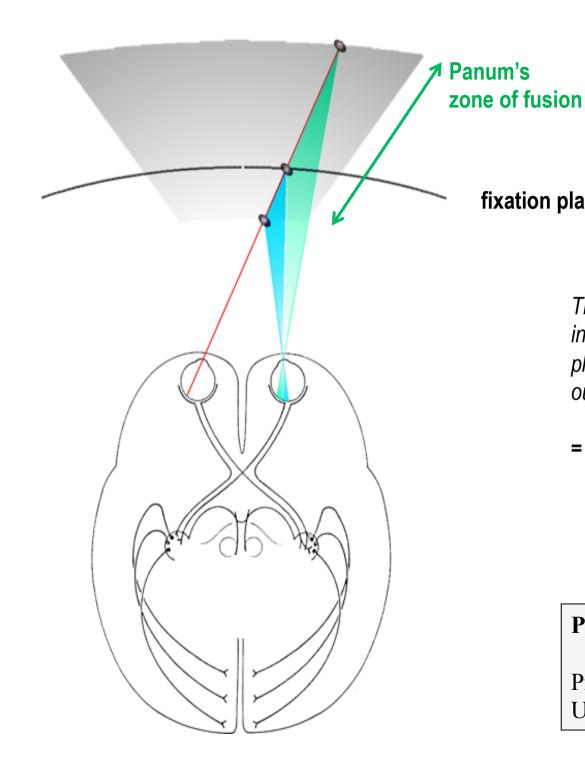




# Stereoscopic Vision







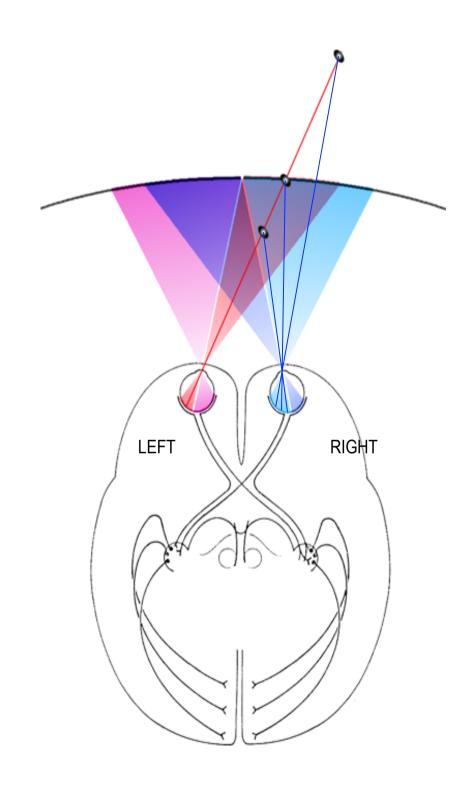
fixation plane

There is a limited range of depths in front, and behind, of the fixation plane in which a <u>single image</u> is seen; outside this range there is double vision.

= Panum's zone of fusion (c. 1860)

## Peter Panum

Professor of Physiology University of Kiel 1853-1864



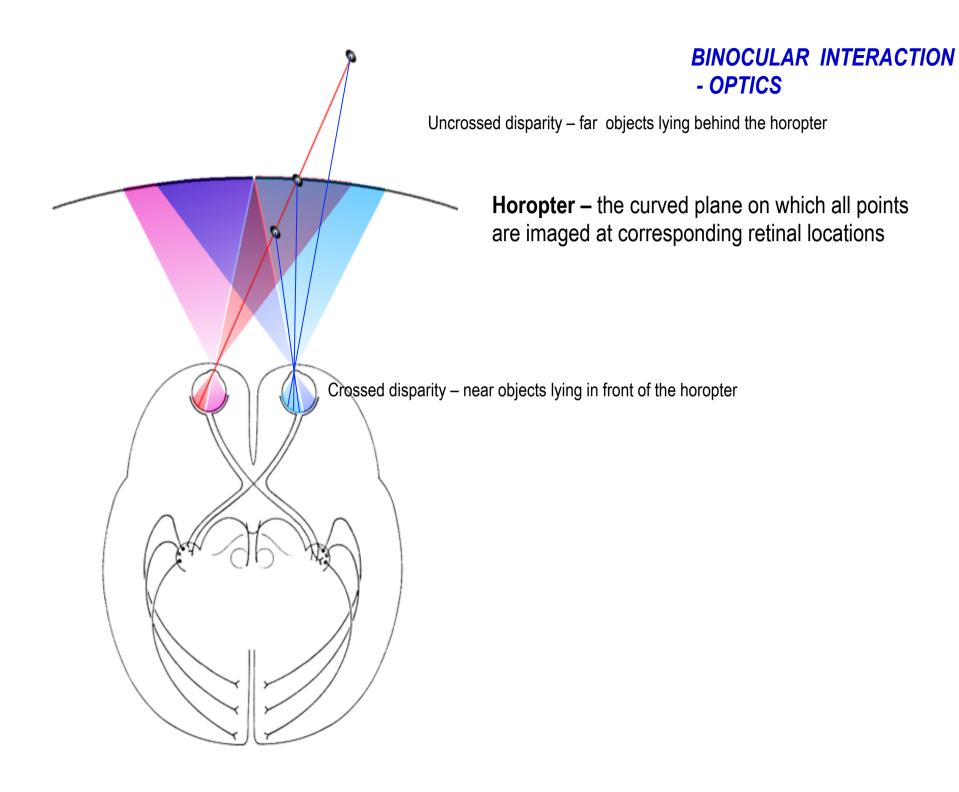
# BINOCULAR INTERACTION - OPTICS

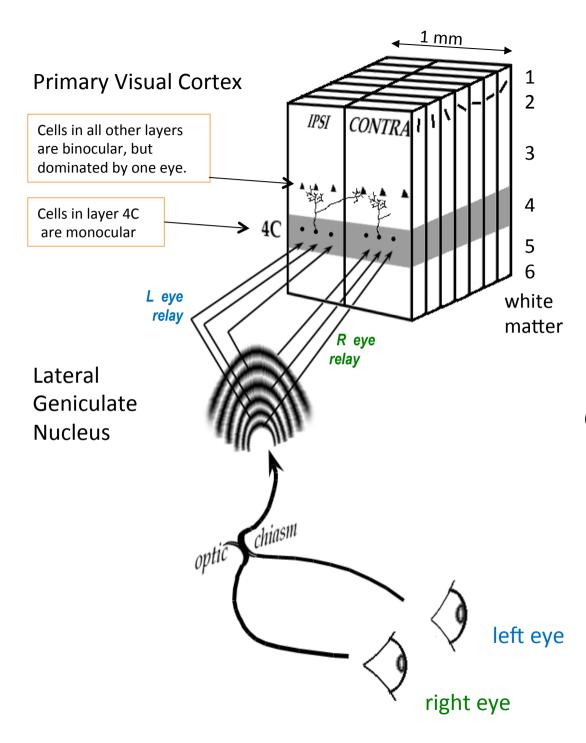
**fixation plane** - images of objects located on the plane of fixation fall at corresponding locations in each retina.

Objects at locations in front, or behind the fixation plane produce images at *noncorresponding* locations in each retina.

e.g.

3 items along the same line of sight in the left eye are imaged at 3 separate locations in the right eye.





# FUNCTIONAL ARCHITECTURE OF PRIMARY VISUAL CORTEX

David **Hubel** & Torsten **Wiesel** 

## V1 (left)

two independent modular subsystems:

- ocular dominance columns
- orientation columns

## LGN (left)

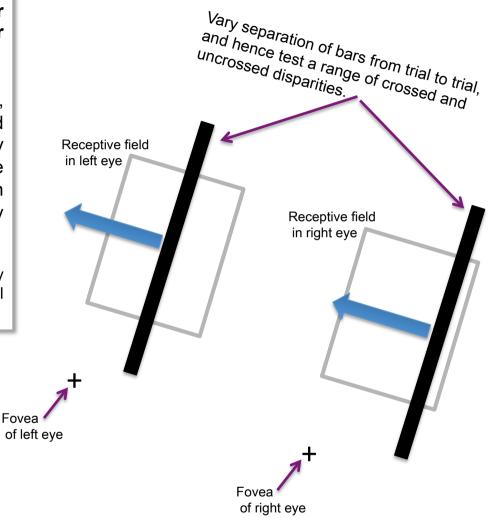
6 monocular layers;

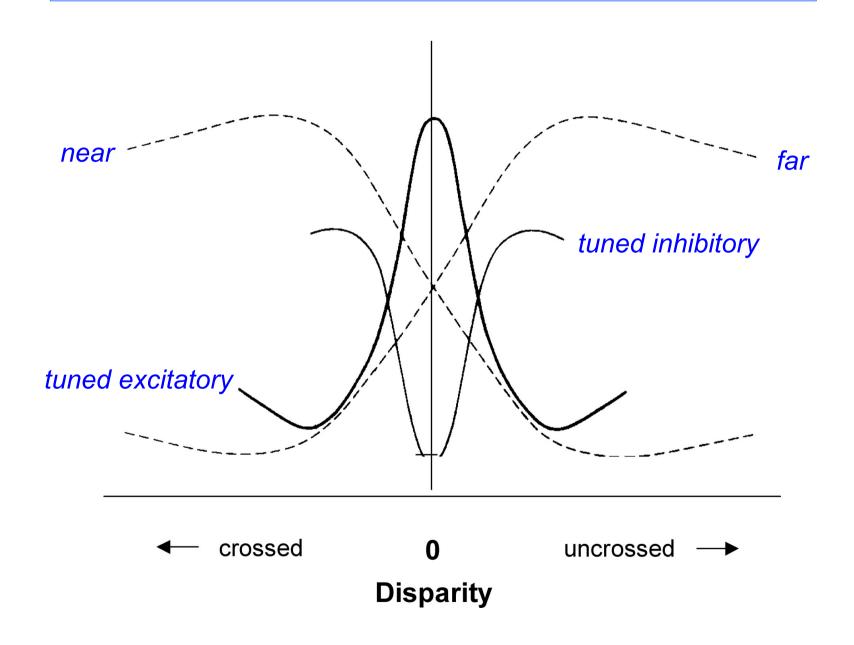
- each layer maps a right, or a left eye half-retina

Typical methodology for testing binocular disparity tuning under conditions of ocular paralysis and anaesthesia.

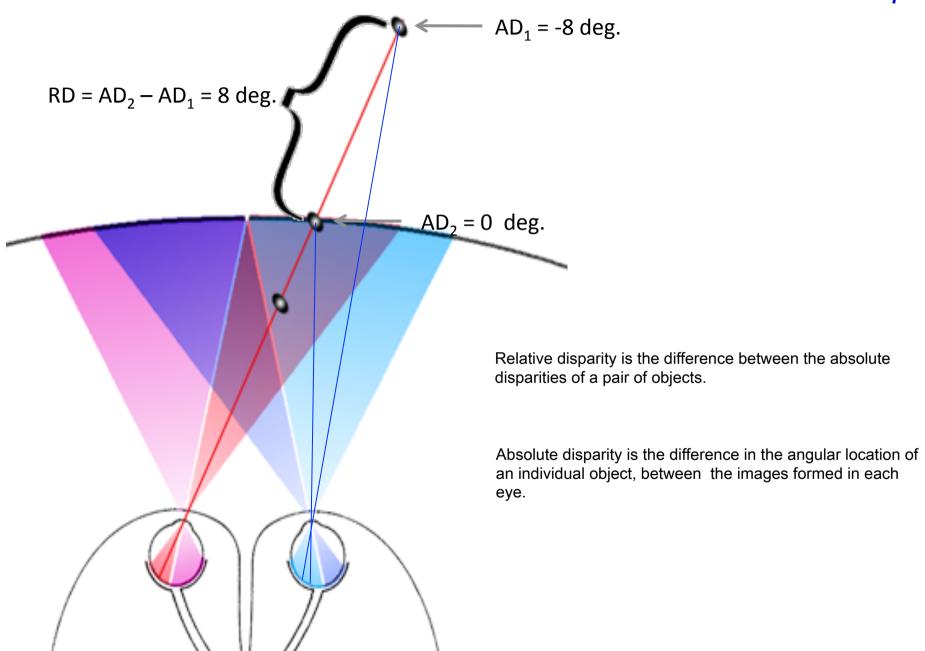
Due to paralysis, the ocular axes diverge, separating the receptive fields in either eye, and allowing independent (dichoptic) stimulation by two separate bar stimuli. As there is only one stimulus (per receptive field) against a plain background, this is a test of absolute disparity tuning.

The following slide shows the four varieties of disparity tuning that have typically been observed in several separate studies of area V1, V2 & V3.

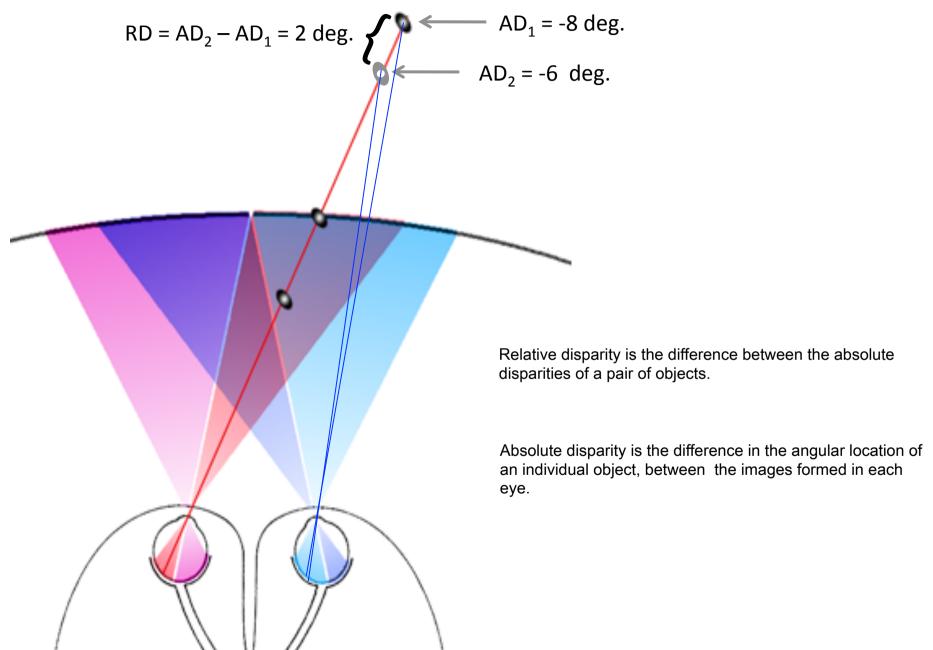




# BINOCULAR INTERACTION Absolute & Relative disparity



# BINOCULAR INTERACTION Absolute & Relative disparity



# Absolute Disparity v. Relative Disparity

#### **ABSOLUTE DISPARITY**

Value is dependent on eye vergence

Gives depth with respect to fixation plane (horopter)

Not very sensitive:

 threshold for detecting depth in front/behind fixation plane is 6 min, i.e. ± 8 cm at 1.0 m.

#### **RELATIVE DISPARITY**

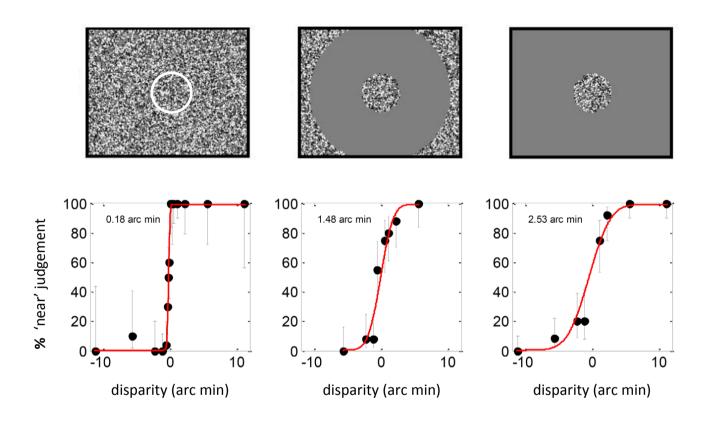
Value is independent of eye vergence

Gives relative depth irrespective of fixation plane (horopter)

Very sensitive:

 threshold for detecting depth in front/behind fixation plane is 6 sec, i.e. ± 3 mm at 1.0 m.

# Relative disparity allows greater stereoacuity



# Absolute Disparity v. Relative Disparity

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Characterises 'DORSAL SYSTEM' function (human fMRI)

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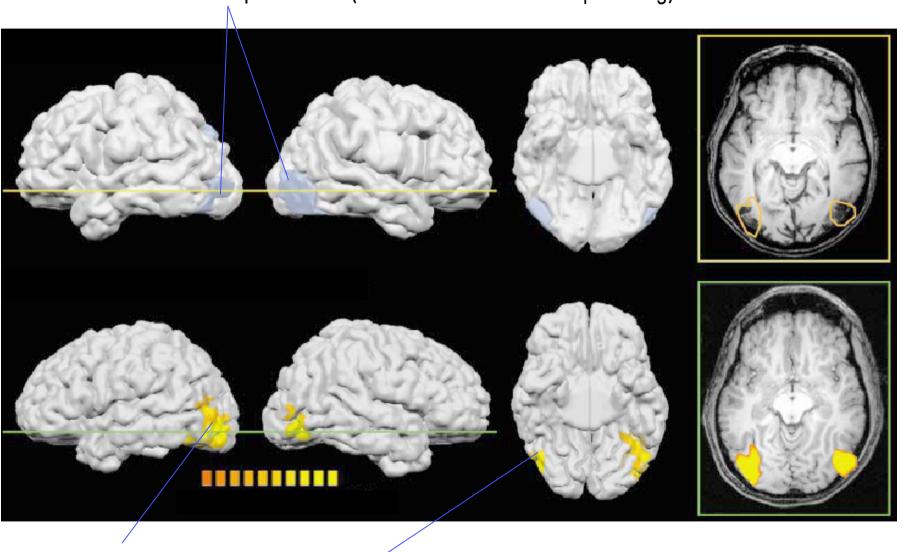
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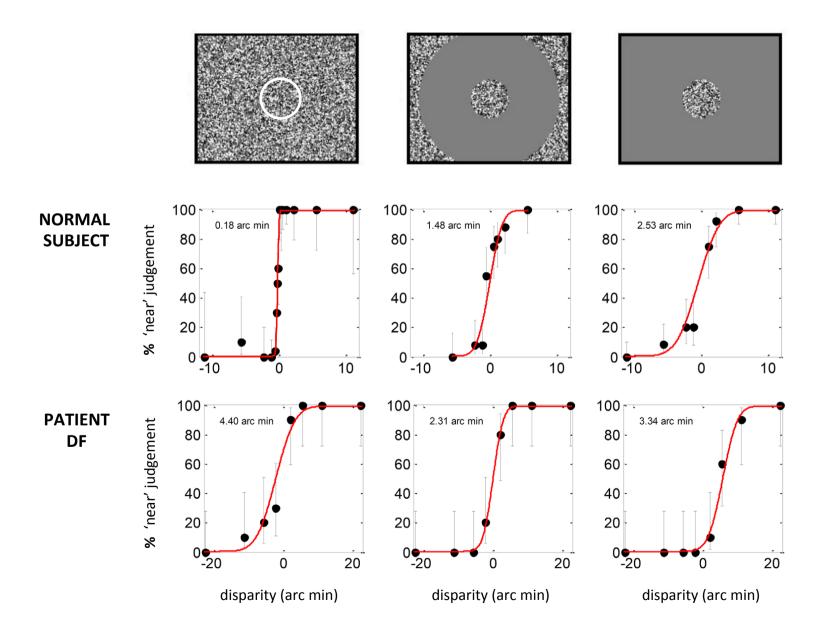
Characterises 'VENTRAL SYSTEM' function (human fMRI)

# Patient DF (bilateral area LO lesions & agnosic)

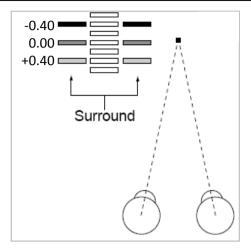
brain lesions in patient DF (case of carbon monoxide poisoning)



area LO in a normal subject



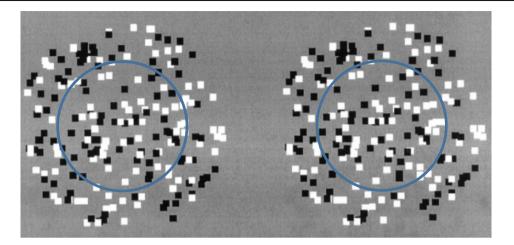
# Comparison of neural selectivity for absolute & relative disparity in areas V1 V2 V3 V3A V4 & V5 [Refs 17-21]

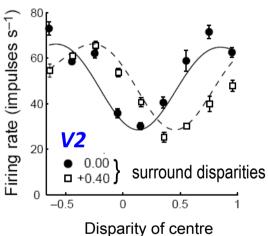


# Stimulus set-up

Dot stimulus is composed of centre & surround annulus differing in disparity

Receptive field of test neuron fits inside the centre region.

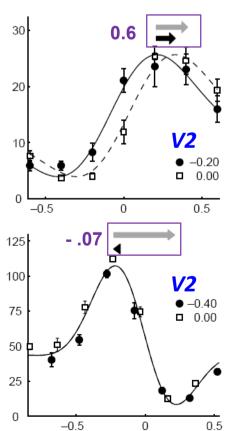


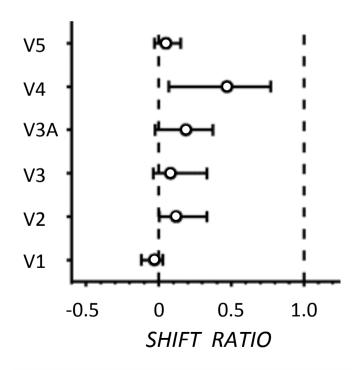


predicted shift (0.40)

observed shift (0.32)

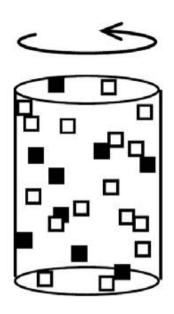
**shift ratio** =  $\frac{0.32}{0.40}$  = **0.8** 



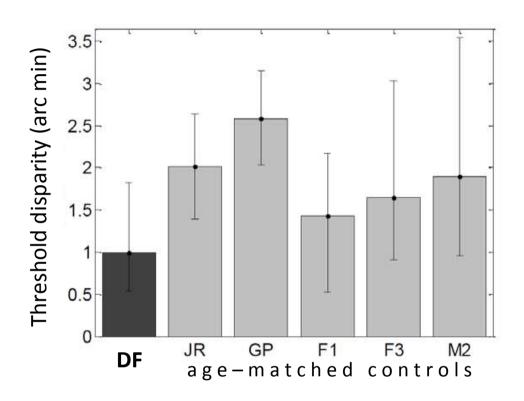


[shift ratio = 1 implies full selectivity for relative disparity; shift ratio = 0 implies selectivity for absolute disparity]

# Read et al. (2010) [16] Stereovision of patient DF



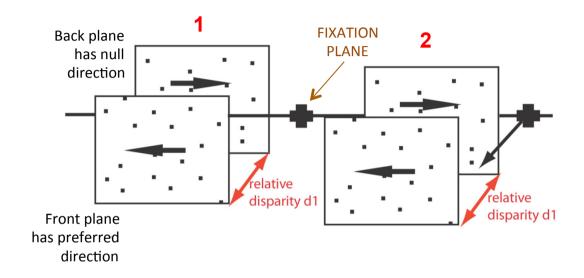
Simulated transparent, revolving cylinder. Task: identify direction (L v R) of front surface

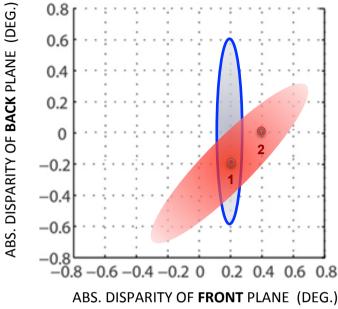


DF outperforms most normal subjects...!

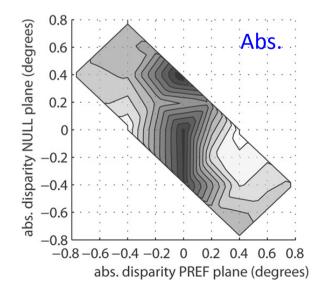
# Krug & Parker (2011) [22] Neurons in area V5 signalling relative disparity

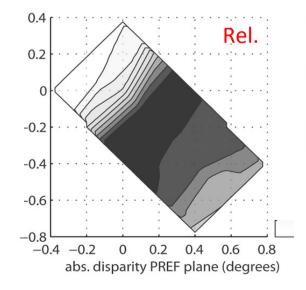
Stimulus: superimposed transparent planes of dots with opposite directions of motion & differing disparity

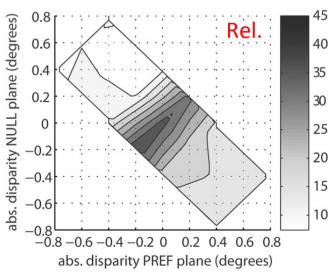




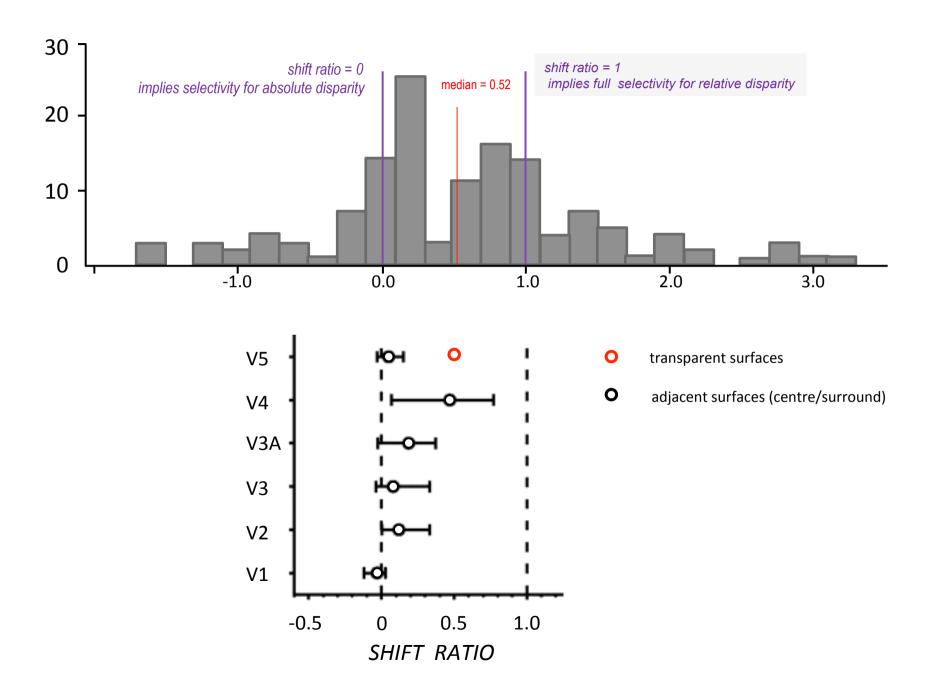
Prediction for absolute disparity tuning Prediction for relative disparity tuning

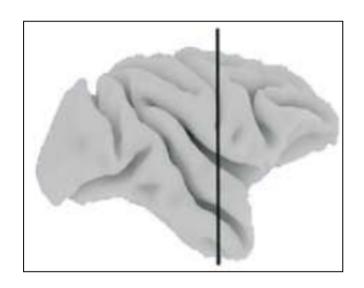


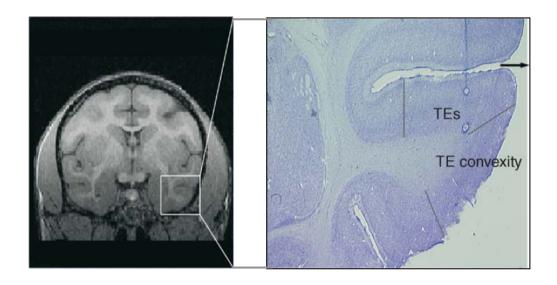


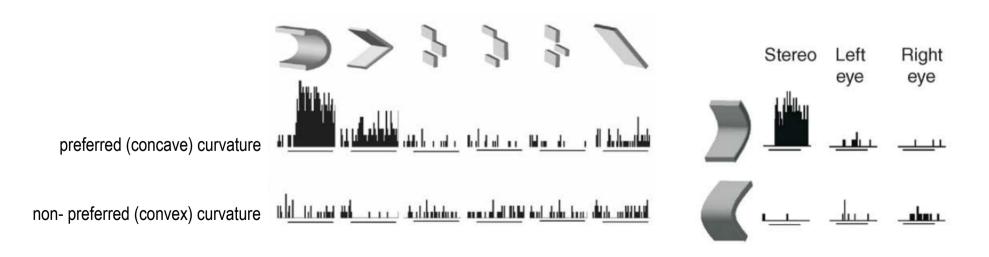


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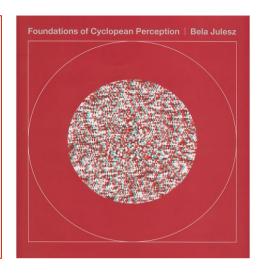
- especially for relative disparity between nearby, adjacent surfaces: e.g. for detecting slant, curvature.

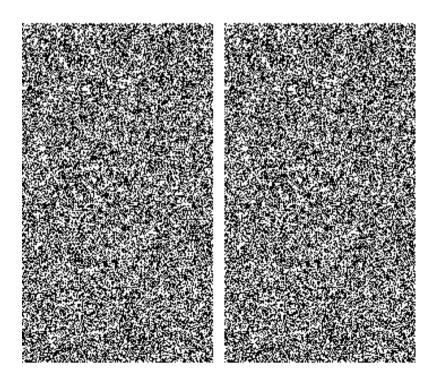
In 'DORSAL SYSTEM' may help top discriminate depth planes e.g. in detecting movement behind foliage (?)

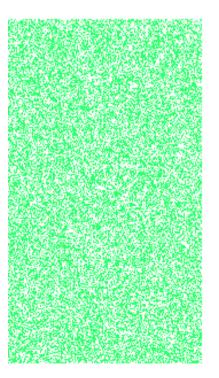
# **Cyclopean vision**

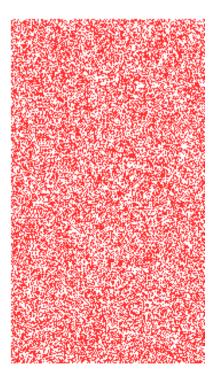
Random dot stereograms demonstrate that stereoscopic mechanisms can process 'raw' visual texture. A preliminary stage of contour extraction or object recognition is not required; rather, stereoscopic mechanisms can produce their own contours and provide an independent mode of form recognition.

To do so the brain must solve the (so-called) 'correspondence problem' – find the correct dot in the right eye to match each dot in the left eye (or vice versa).







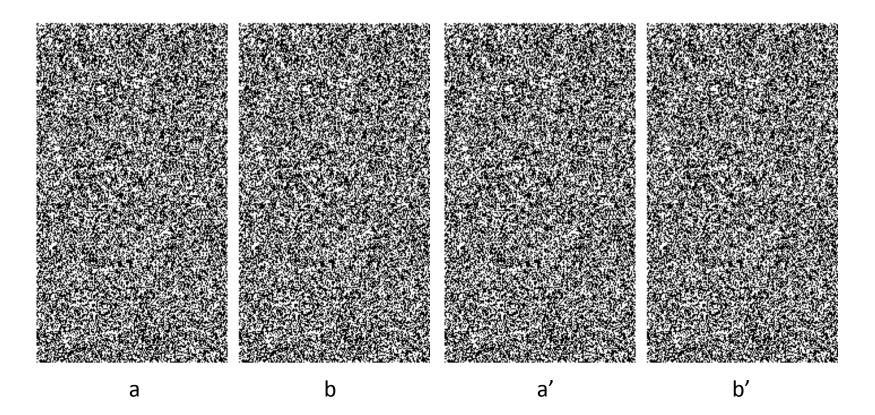


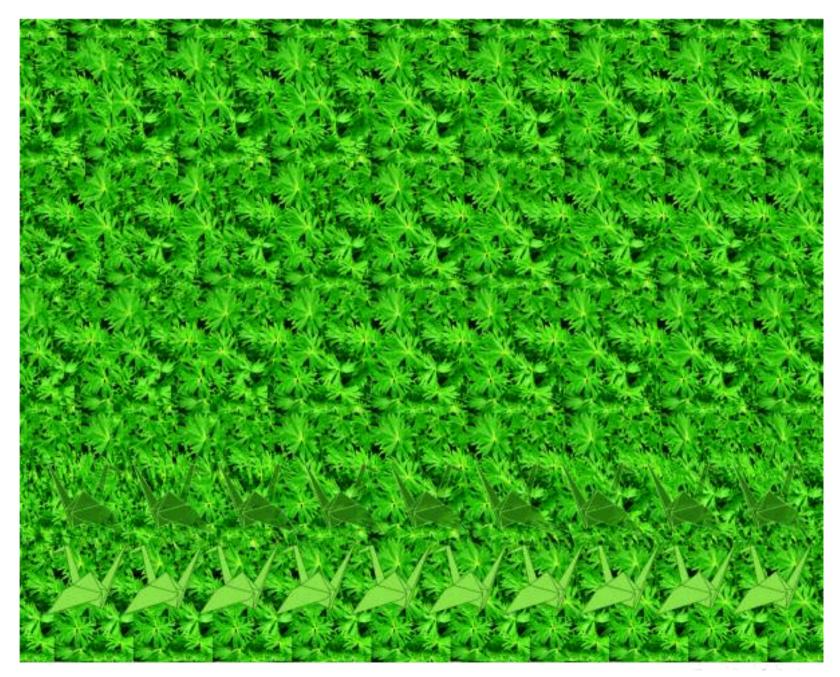
## Cyclopean vision – 'free fusion'

Random dot stereograms demonstrate that stereoscopic mechanisms can process 'raw' visual texture. A preliminary stage of contour extraction or object recognition is not required; rather, stereoscopic mechanisms can produce their own contours and provide an independent mode of form recognition.

To do so the brain must solve the (so-called) 'correspondence problem' – find the correct dot in the right eye to match each dot in the left eye (or vice versa).

Below, **a** & **b**, and **a'** & **b'**, are two identical stereograms, that display a number (from 0-9) in front of the background, when 'solved' by converging. At the same time, the same number will appear as a cut-out (i.e. behind the background) in the stereo pair formed by **b** and **a'**. Foreground and background depth is reversed if the stereograms are solved by diverging the eyes, i.e. to focus on a plane behind the screen. Typically, this is more difficult.





'Magic eye' pictures are based on exactly the same principle (but normally require divergent free fusion to give the intended depth percept).