

Schizophrenia, Attractors and Working Memory

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CCN Lecture 5

This lecture and the next

- Working memory models as example of dynamical models and attractor dynamics in biological neural networks;
- Application to Computational Psychiatry - Schizophrenia.
- Reading: Chapter 3 of Computational Psychiatry Primer.

1) What is schizophrenia?

2) What is working memory?

3) Attractor Paradigm for working memory/ delayed activity

- Hopfield Net

- Back to the Ring Model

4) Application to schizophrenia

1) What is Schizophrenia ?

- ❖ Major neuropsychiatric disorder
- ❖ 1% lifetime risk
- ❖ can start as sudden psychotic episode usually in early adulthood or gradual withdrawal and decline.

1. Positive symptoms: (appearance of abnormal experiences - psychosis)

- Threatening voices
- Thought Echoing
- Visual Hallucination
- Delusions



1) What is Schizophrenia ?

2. Disorganisation:

- Pressure of thought

“ in my mind there ran like and endless clockwork a compulsive torturing uninterrupted chain of ideas”

- Incongruent behaviour

“ I really would like to slap your face, people like you are usually called SOB’s”

- disturbed sense of self (“mirror sign”)



1) What is Schizophrenia ?

3. Negative Symptoms: (deficits in normal experience)

- Poverty of thought / perseverance
- Loss of Initiative
- anhedonia: loss of pleasure
- social withdrawal

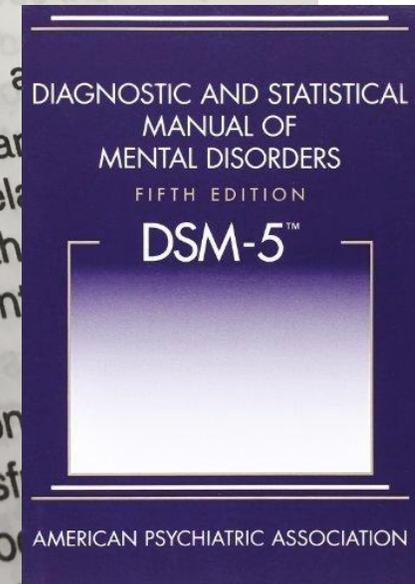
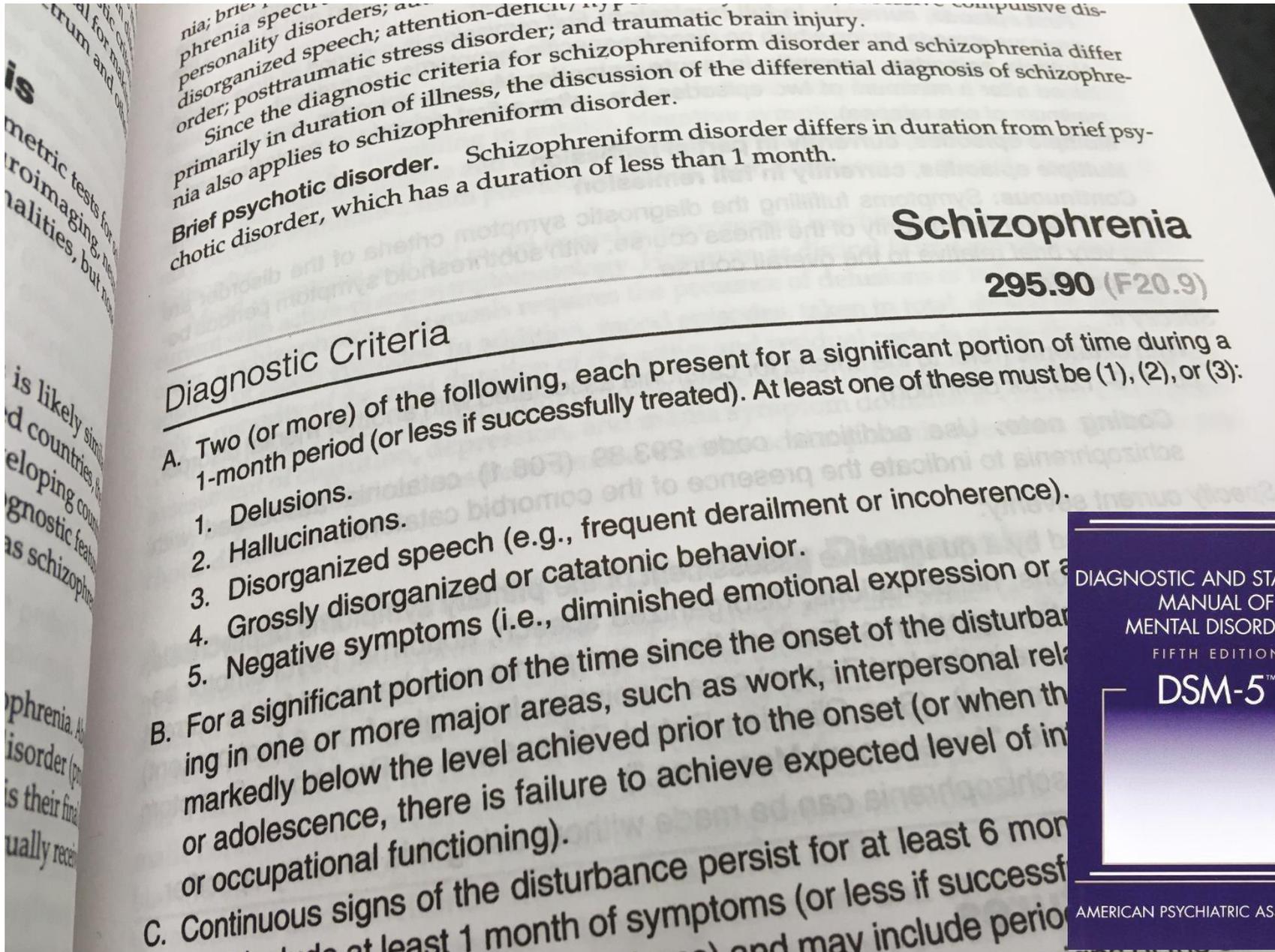
- **Cognitive deficits**

Difficulty in maintaining context, in learning associations.

working memory deficits



1) What is Schizophrenia ?



1) What is Schizophrenia ?

- **Cause is unknown** : Combination of genetic and environmental factors
- **No cure**. Medication alleviate the (+) symptoms somewhat.
- **Disconnection hypothesis.**
- **Dopamine hypothesis:** misfiring of dopaminergic neurons.
 - **Antipsychotic drugs** work on the dopamine system.
 - **Amphetamines** (DA agonist) can mimic some aspect of the disorder.
- **NMDA hypothesis:** Impairments in NMDA transmission.
 - **Ketamine** (NMDA receptor antagonist) mimic aspects of the disorder.
- Positive symptoms are most fascinating but negative/ cognitive symptoms create most suffering and disability.
- 10% suicide rate.

- Building upon ongoing advances in the **behavioural and neurobiological sciences**
- "RDoC is an attempt to create **a new kind of taxonomy** for mental disorders by bringing the power of modern research approaches in genetics, neuroscience, and behavioral science to the problem of mental illness."
[Elevag et al 2016]
- From a categorical to a **dimensional** approach.

NIMH Research Domain Criteria (RDoC)									
Functional Domains									
Negative Valence Systems (e.g., fear, anxiety, loss)		Positive Valence Systems (e.g., reward, learning, habit)		Cognitive Systems (e.g., attention, perception, memory)		Systems for Social Processes (e.g., attachment, communication, perception of self & others)		Arousal and Regulatory Systems (e.g., arousal, circadian rhythms)	
Units of Analysis									
Genes	Molecules	Cells	Circuits	Physiology	Behavior	Self-Reports	Paradigms	Genes	Molecules

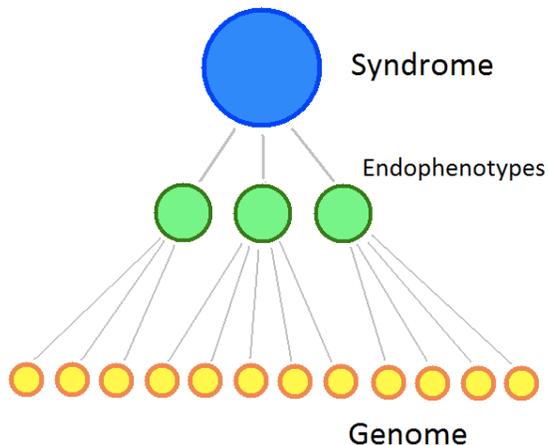
Linking Neurobiology and Psychiatry

- Working memory as a possible **endophenotype**?



endophenotype

n. a type of biological marker that is simpler to detect than genetic sequences and that may be useful in researching vulnerability to a wide range of psychological and neurological disorders. Endophenotypes may be a useful link between genetic sequences and their external emotional, cognitive, or behavioral manifestations. [coined in the 1970s by U.S. behavioral geneticist and clinical psychologist Irving I. Gottesman (1930-) and British psychiatrist James Shields (d. 1978)]



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Working memory impairment as an endophenotypic marker of a schizophrenia diathesis

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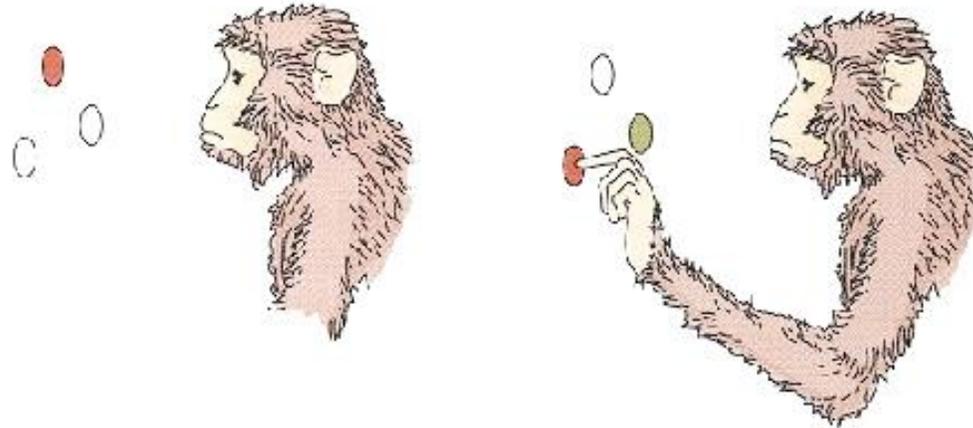
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ABSTRACT

This review focuses on the viability of working memory impairment as an endophenotypic marker of a schizophrenia diathesis. It begins with an introduction of the construct of working memory. It follows the operational criteria for defining an endophenotype. Research findings regarding the working memory impairment of schizophrenia and schizophrenia-spectrum patients, first-degree relatives of schizophrenia patients, and schizophrenia-spectrum patients, first-degree relatives of schizophrenia patients, are reviewed in terms of the criteria for being considered an endophenotypic marker. Research findings regarding the working memory deficit (namely, encoding, maintenance, and retrieval) in terms of which aspects are likely to be the best candidates for endophenotypes. Literature regarding working memory performance in bipolar disorder and major depression is reviewed in terms of relative specificity to schizophrenia. Despite some unresolved issues, it appears that working memory impairment is a very promising candidate for an endophenotypic marker of a schizophrenia diathesis.

2) What is working memory ? (a.k.a. short-term memory)

- The ability to hold information over a time scale of seconds to minutes
- a critical component of all cognitive functions (language, thoughts, planning etc..)



Delayed match-to sample task:
remember 'red'



- central to many learning disabilities and impaired in many mental disorders

<http://news.bbc.co.uk/1/hi/health/4101431.stm>

<http://www.youtube.com/watch?v=M1Tui0Gbvq4&feature=related>

2) What is working memory ?

Test yours!

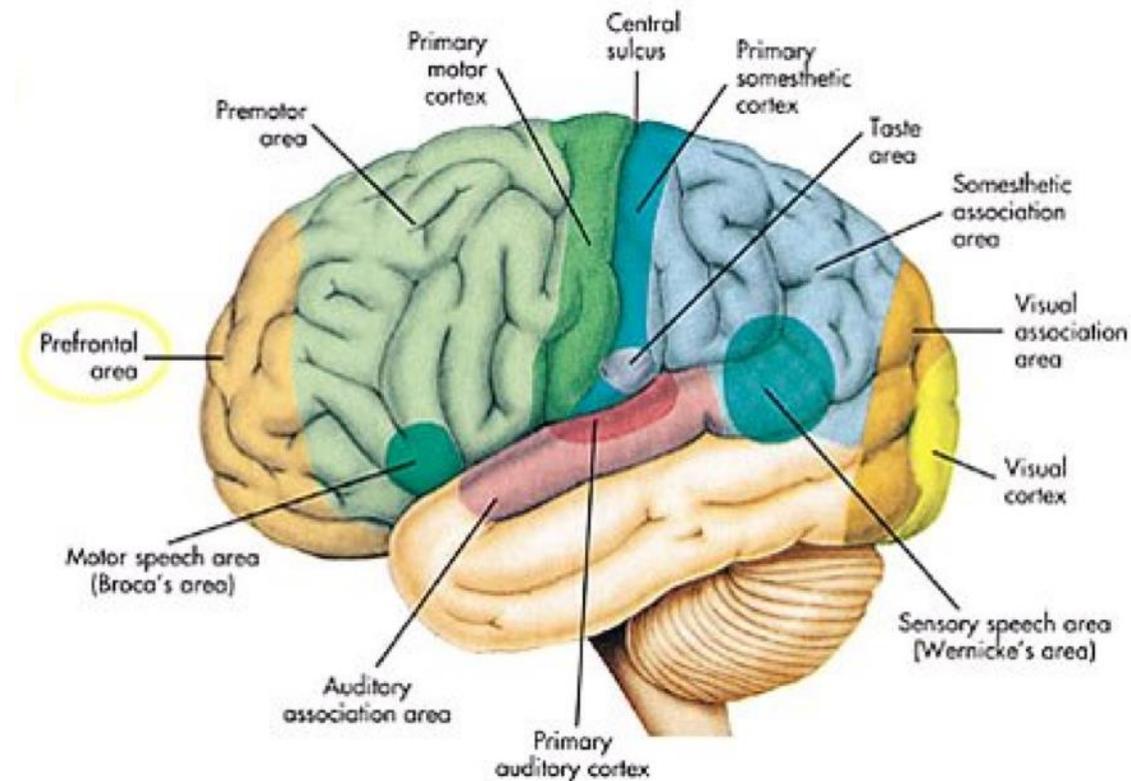
https://www.ted.com/talks/peter_doolittle_how_your_working_memory_makes_sense_of_the_world

2:21-3:49



Sustained activity in PFC (1)

- **Lesion** and inactivation studies demonstrate crucial role of **Prefrontal Cortex** (PFC) in working memory, in particular dorsolateral PFC (PFDl).



Sustained activity in PFC (2)

Oculo-motor delayed response task:
remember location of cue.

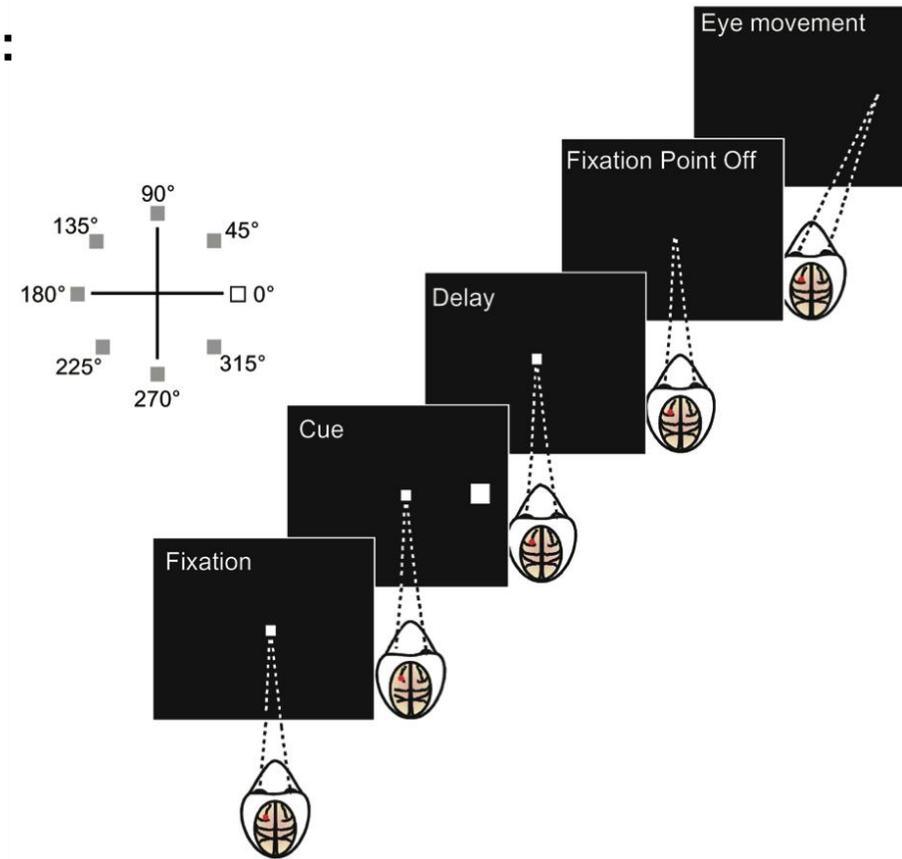
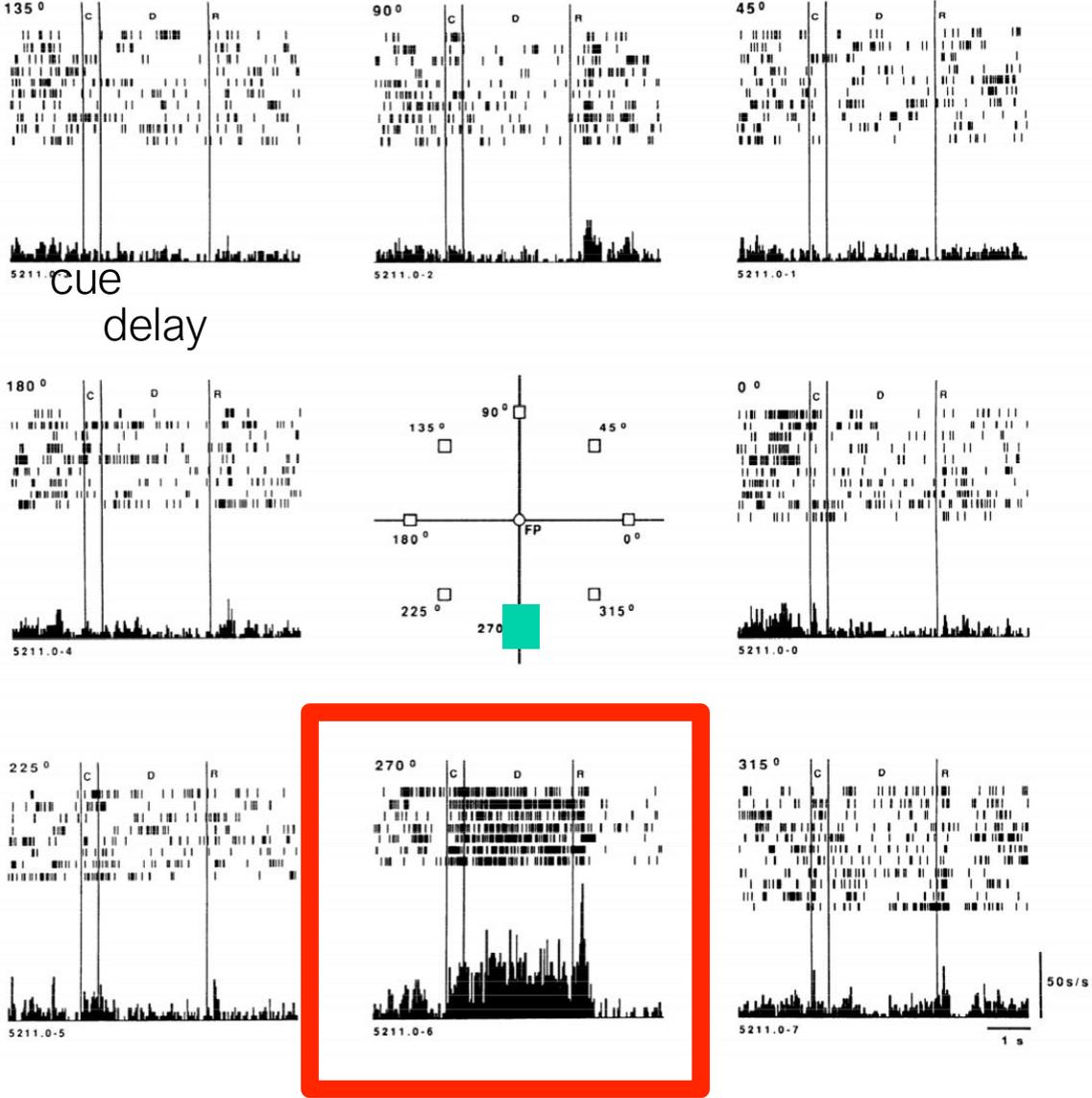


Fig. 1. Successive frames illustrate the sequence of events in the oculomotor delayed-response task. Trials begin with the appearance of a fixation point at the center of the screen, which the monkey is required to foveate throughout the trial. A spatial cue is subsequently presented, typically at one of eight locations (*left*). After a delay period of a few seconds, the fixation point is turned off and the monkey is required to indicate the location of the cue by moving his eyes accordingly on the screen.

Sustained activity in PFC (2)

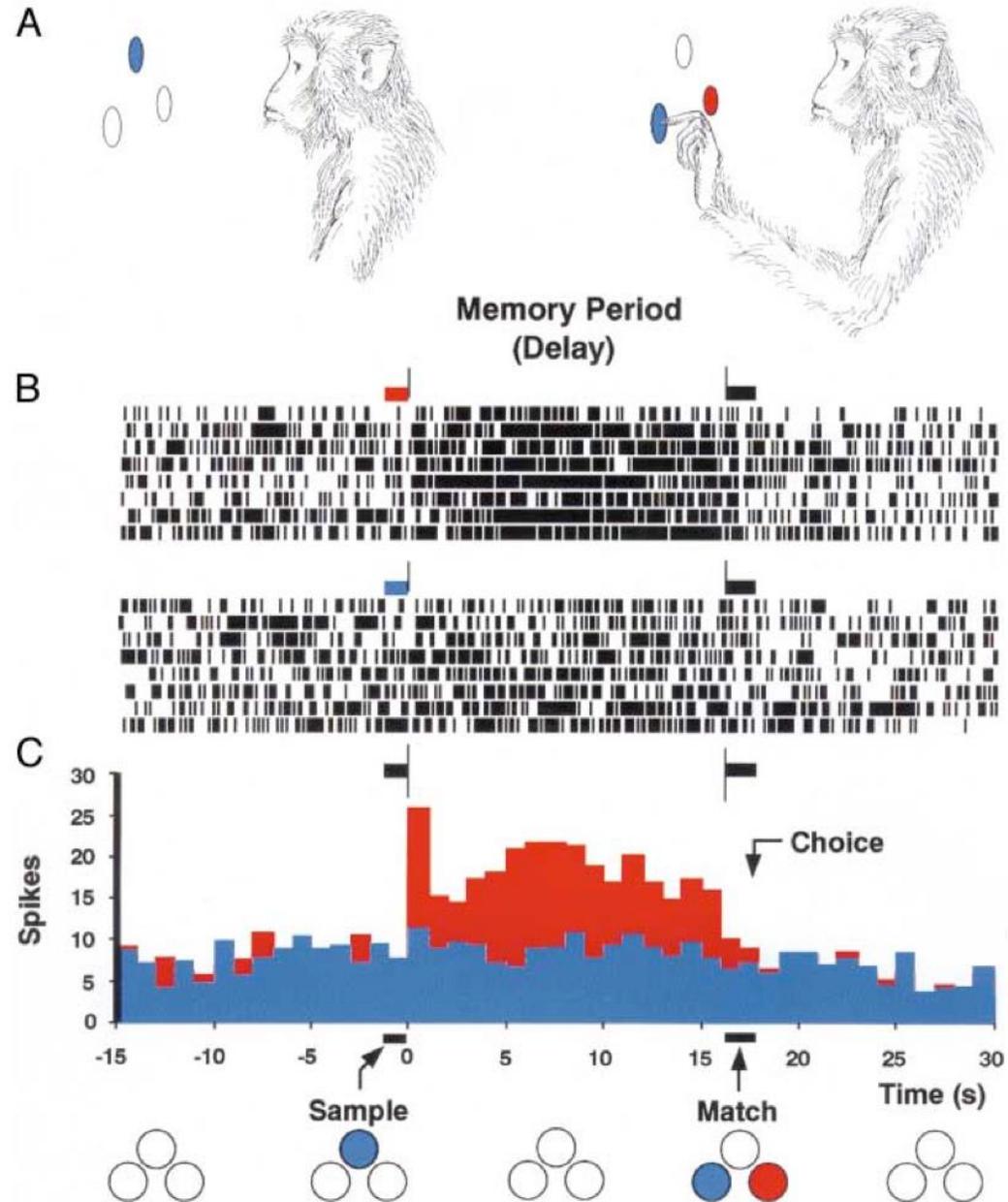


Patricia Goldman-Rakic
1937-2003



Funahashi et al, 1989

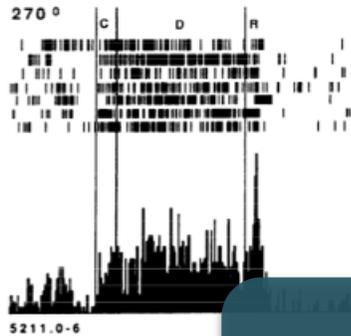
Sustained activity in inferotemporal Cx (IT)



Fuster and Jervey
1982

Working memory vs Long-term memory and Linking propositions

- Long-term memory: **molecular or structural changes - connectivity between neurons.**
- Short-term/ working memory: **Dynamical process** that has not yielded to molecular characterisation: **"Sustained" a.k.a. "Delay" or "Persistent" Activity.**



Delay activity



WM

Towards a theory of Working Memory/ Sustained Activity

- A theory of working memory should answer:
 - How is it initiated?
 - Why does it persist ?
 - What makes it specific?
 - How does it end?

 - Reason for capacity limit?
 - Relationship with attention, long-term memory?
- Mechanism : reverberations through connections (which?), or cellular?
- Lots of experimental and theoretical work to answer these questions

Delay activity could be due to reverberations through...

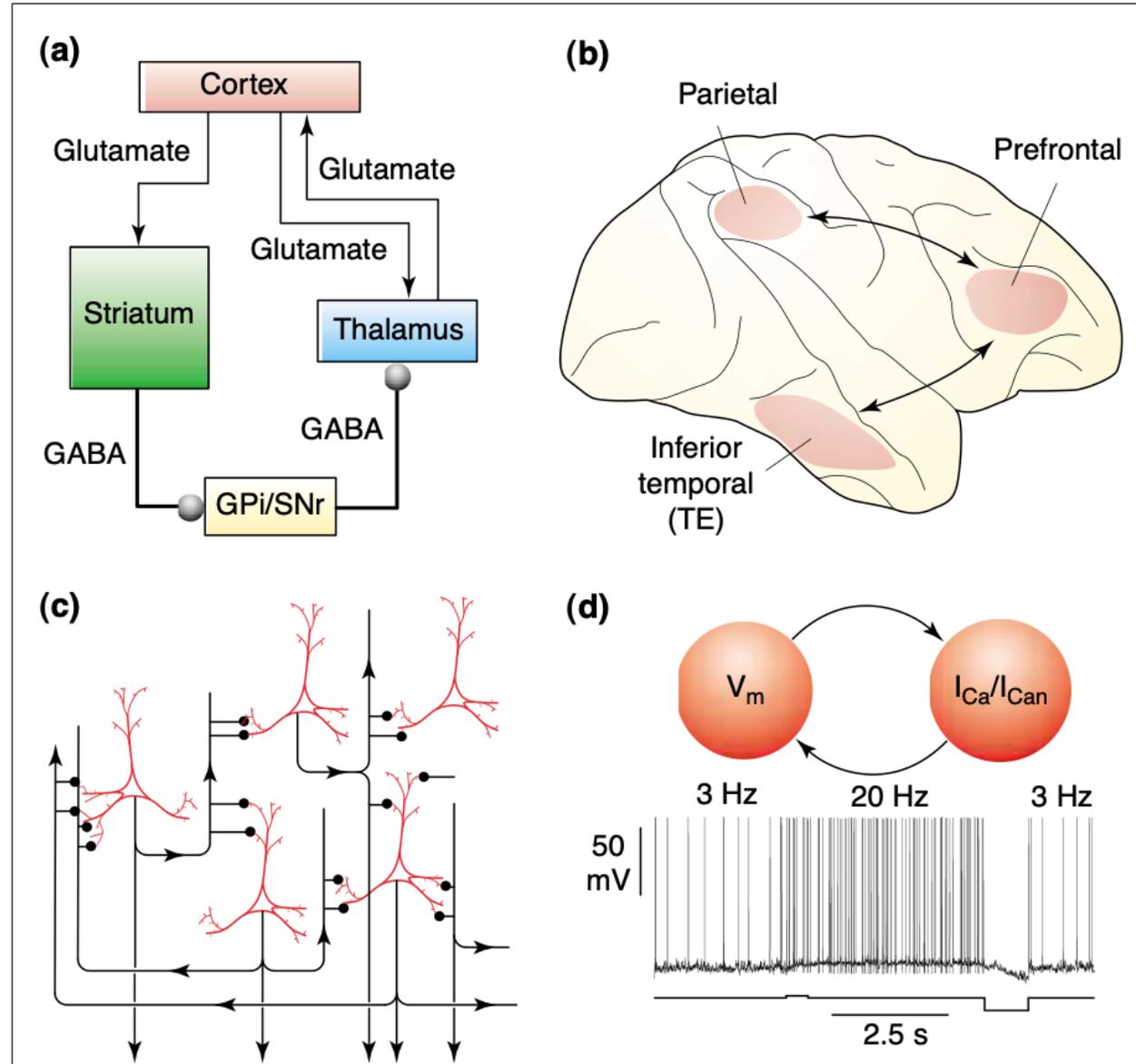


Fig. 1. Various scenarios for the anatomical substrate of excitatory reverberation in the cortex. (a) A closed thalamo-cortical loop and/or cortico-striato-thalamic-cortical circuit. In the latter case, cortical excitation of the caudate nucleus leads to an inhibition of the output from the cells of the basal ganglia and thus a disinhibition of thalamic neurons, which in turn send increased excitation back to the cortex. Abbreviations: GPi, internal segment of the globus pallidus; SNr, substantia nigra pars reticulata. (b) Reciprocal interactions between two cortical areas, prefrontal and posterior cortices (respectively prefrontal and inferotemporal cortices) for spatial (resp. object) visual memory. (c) Excitatory recurrent collaterals within a local circuit. (d) Intrinsic regenerative dynamics of single neurons. Positive feedback between membrane depolarization/spike discharges and active inward currents (voltage-gated I_{Ca} /Ca²⁺-gated I_{Can}) can produce persistent activity that outlasts a transient input current pulse (J. Tegnér and X.-J. Wang, unpublished).

3) Attractor Paradigm for Persistent Activity

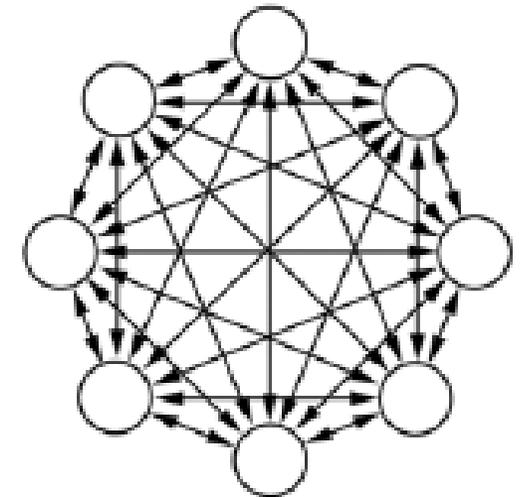
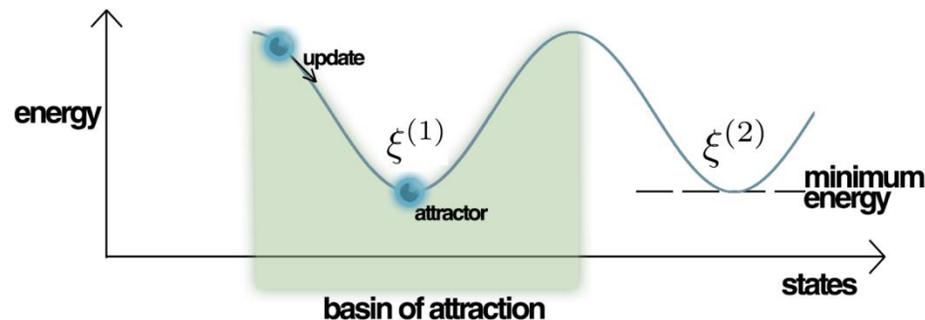
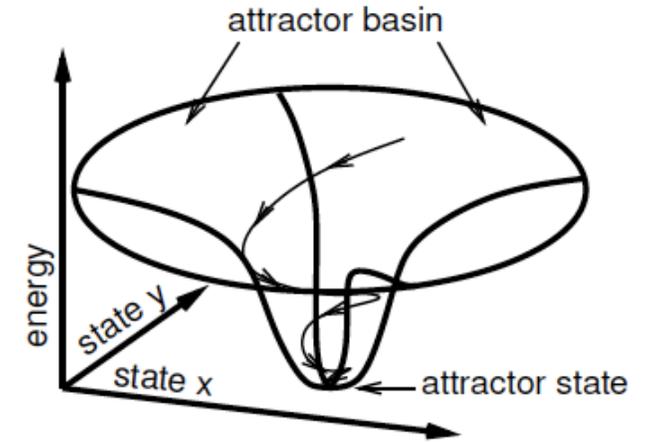
• Since 1970s it is proposed that delay activity patterns can be theoretically described by **dynamical attractors** in **recurrent neural networks**.

• **Hopfield networks** (John Hopfield (1982))

$$s_i = \begin{cases} 1 & \text{if } \sum_j w_{ij}s_j > \theta_i, \\ -1 & \text{otherwise.} \end{cases}$$

$$w_{ij} = \frac{1}{N} \sum_{k=1}^{k=N} \xi_i^k \xi_j^k$$

$$E = -\frac{1}{2} \sum_{i < j} w_{ij} s_i s_j + \sum_i \theta_i s_i$$



Hopfield Networks (1)

- A Hopfield net is a form of **recurrent artificial neural network** invented by John Hopfield (1982).
- Hopfield nets typically have **binary** (1/-1 or 1/0) **threshold units**:

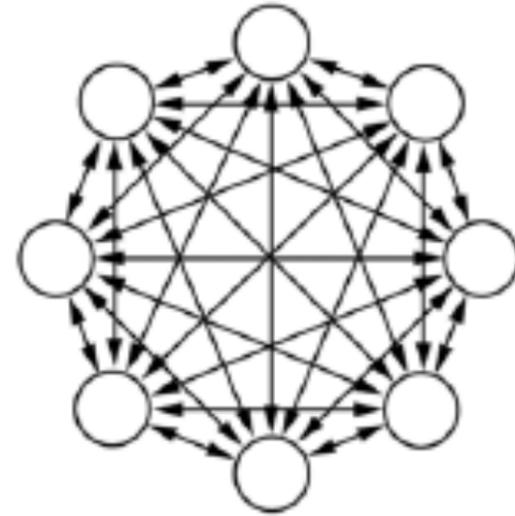
$$s_i = \begin{cases} 1 & \text{if } \sum_j w_{ij}s_j > \theta_i, \\ -1 & \text{otherwise.} \end{cases}$$

where s_j state of unit j , and θ_i is the threshold

The weights have to follow: $w_{ii}=0$, $w_{ij}=w_{ji}$

- Hopfield nets have a scalar value associated with each state of the network referred to as the "**energy**", E , of the network, where:

$$E = -\frac{1}{2} \sum_{i < j} w_{ij} s_i s_j + \sum_i \theta_i s_i$$



Hopfield Networks (2)

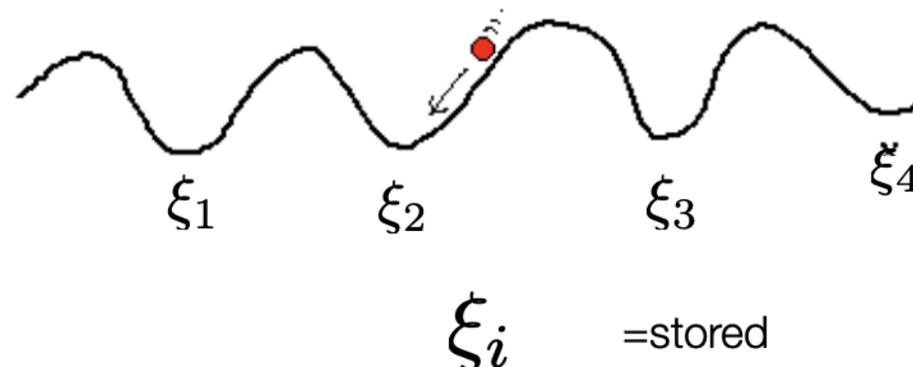
- **Initialization** is done by setting the values of the units to the start pattern (e.g. noisy or incomplete image).
- **Running**: at each step, pick a node at random and update (asynchronous update).

$$s_i = \begin{cases} 1 & \text{if } \sum_j w_{ij}s_j > \theta_i, \\ -1 & \text{otherwise.} \end{cases}$$

- Energy is guaranteed to go down & network to settle in local energy minima.
- **Learning**: the weights are set so as to 'shape' those local minima.

$$w_{ij} = \frac{1}{N} \sum_{k=1}^{k=N} \xi_i^k \xi_j^k$$

- Network will converge to stored state even if it is given only part of the state:



Associative memories

- The Hopfield network is an **associative/content addressable memory**. It can be used to recover from a distorted input the trained state that is most similar to that input.
- e.g. if we train a Hopfield net with 5 units so that the state (1, 0, 1, 0, 1) is an energy minimum. and we give the network the state (1, 0, 0, 0, 1) it will converge to (1, 0, 1, 0, 1).



Reminiscent of human memory?

The Nobel Prize in Physics 2024

The Royal Swedish Academy of Sciences has decided to award the Nobel Prize in Physics 2024 to

John J. Hopfield

Princeton University, NJ, USA

Geoffrey Hinton

University of Toronto, Canada

“for foundational discoveries and inventions that enable machine learning with artificial neural networks”

They trained artificial neural networks using physics

This year’s two Nobel Laureates in Physics have used tools from physics to develop methods that are the foundation of today’s powerful machine learning. John Hopfield created an associative memory that can store and reconstruct images and other types of patterns in data. Geoffrey Hinton invented a method that can autonomously find properties in data, and so perform tasks such as identifying specific elements in pictures.

When we talk about artificial intelligence, we often mean machine learning using artificial neural networks. This technology was originally inspired by the structure of the brain. In an artificial neural network, the brain’s neurons are represented by nodes that have different values. These nodes influence each other through connections that can be likened to synapses and which can be made stronger or weaker. The network is *trained*, for example by developing stronger connections between nodes with simultaneously high values. This year’s laureates have conducted important work with artificial neural networks from the 1980s onward.

John Hopfield invented a network that uses a method for saving and recreating patterns. We can imagine the nodes as pixels. The *Hopfield network* utilises physics that describes a material’s characteristics due to its atomic spin – a property that makes each atom a tiny magnet. The network as a whole is described in a manner equivalent to the energy in the spin system found in physics, and is trained by finding values for the connections between the nodes so that the saved

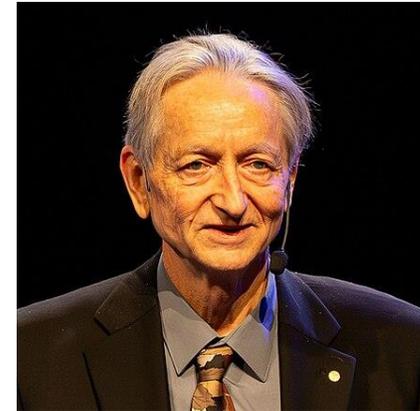
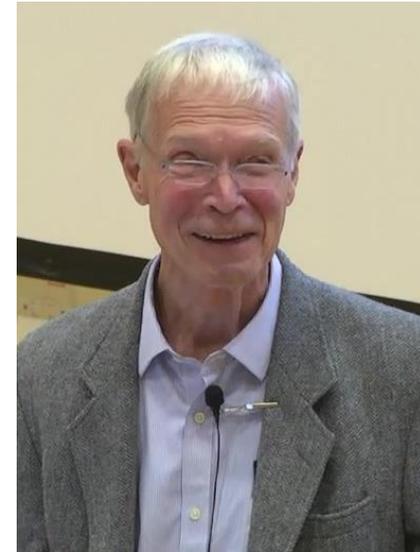
images have low energy. When the Hopfield network is fed a distorted or incomplete image, it methodically works through the nodes and updates their values so the network’s energy falls. The network thus works stepwise to find the saved image that is most like the imperfect one it was fed with.

Geoffrey Hinton used the Hopfield network as the foundation for a new network that uses a different method: the *Boltzmann machine*. This can learn to recognise characteristic elements in a given type of data. Hinton used tools from statistical physics, the science of systems built from many similar components. The machine is trained by feeding it examples that are very likely to arise when the machine is run. The Boltzmann machine can be used to classify images or create new examples of the type of pattern on which it was trained. Hinton has built upon this work, helping initiate the current explosive development of machine learning.

“The laureates’ work has already been of the greatest benefit. In physics we use artificial neural networks in a vast range of areas, such as developing new materials with specific properties,” says Ellen Moons, Chair of the Nobel Committee for Physics.

John J. Hopfield, born 1933 in Chicago, IL, USA. PhD 1958 from Cornell University, Ithaca, NY, USA. Professor at Princeton University, NJ, USA.

Geoffrey Hinton, born 1947 in London, UK. PhD 1978 from The University of Edinburgh, UK. Professor at University of Toronto, Canada.



Hoffmans' model of Schizophrenia (1989): the pruning hypothesis

❖ Synapses progressively reduce throughout childhood and adolescence.

❖ Schizophrenia: pruning process gone too far.

VOL. 15, NO. 3, 1989

Cortical Pruning and the Development of Schizophrenia: A Computer Model

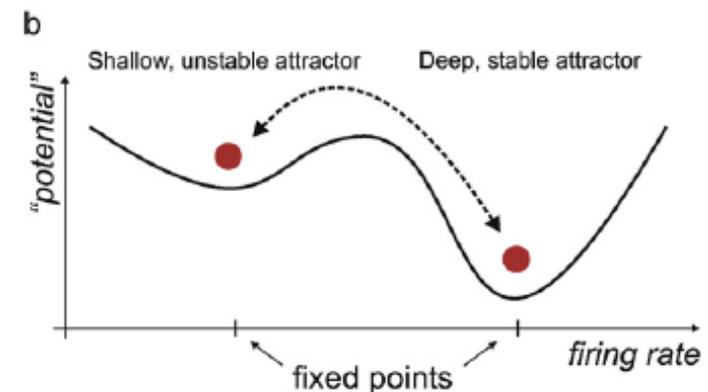
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by Ralph E. Hoffman and Steven K. Dobscha

Abstract

Schizophrenic patients tend to demonstrate reduced cerebral metabolism in frontal areas. Studies of human brain development reveal that synapses in the cerebral cortex are progressively reduced throughout childhood and adolescence, with parallel reductions in cerebral metabolism. This relationship is not surprising, since synaptic density is a primary factor determining regional metabolic requirements. The elimination of

ter being physical phenomena. Computer simulations of neural network interactions are now offering clues about how biologically characterized events can accomplish cognitive tasks. Variants of these models may reveal how disordered neural processes can yield particular kinds of disordered cognition. This report critically examines a number of biological studies of schizophrenic patients in the context of recently emerging knowledge of human brain development. A particular neuroanatomical disturbance



Hoffmans' model of Schizophrenia (1989): the pruning hypothesis

- ❖ Synapses progressively reduce throughout childhood and adolescence.
- ❖ Schizophrenia: pruning process gone too far?
- ❖ Study of pruning in Hopfield nets leads to :
merging of attractors (thought disorder?),
spurious attractors (hallucinations?)..

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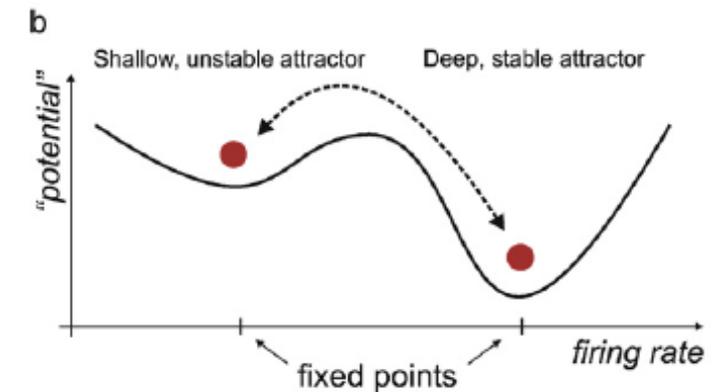
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Intermediate Summary

- **Schizophrenia** is a very serious illness characterised by “positive” (hallucinations, delusions) and negative symptoms.
- One neurobiological correlate of the illness is impairment in **working memory**.
- Short-term/ working memory: **Dynamic process** - “Sustained” a.k.a. “Delay” or “Persistent” Activity.
- **Attractor Networks** as (main) model of working memory / sustained activity
- **Hopfield Network** as example of a point attractor model (Lab 2).
- Next time: How do we go from here to providing biologically plausible spiking models, comparable to recordings in Prefrontal cortex?