

# Marr's Theory of the Hippocampus

Computational Models of Neural Systems  
Lecture 3.3

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# David Marr: 1945-1980



David Marr  
1970 – Cambridge, England

# Marr and Computational Neuroscience

- In 1969-1970, Marr wrote three major papers on theories of the cortex:
  - *A Theory of Cerebellar Cortex*
  - *A Theory for Cerebral Neocortex*
  - *Simple Memory: A Theory for Archicortex*
- *A fourth paper, on the input/output relations between cortex and hippocampus, was promised but never completed.*
- Subsequently he went on to work in computational vision.
- His vision work includes a theory of lightness computation in retina, and the Marr-Poggio stereo algorithm.

# Introduction to Marr's Archicortex Theory

- The hippocampus is in the “relatively simple and primitive” part of the cerebrum: the archicortex.
  - The *piriform* (olfactory) cortex is also part of archicortex.
- Why is archicortex considered simpler than neocortex?
  - Evolutionarily, it's an earlier part of the brain.
  - Fewer cell layers (3 vs. 6)
  - Other reasons? [connectivity?]
- Marr claims that neocortex can learn to classify inputs (category formation), whereas archicortex can only do associative recall.
  - Was this conclusion justified by the anatomy?

# What Does Marr's Hippocampus Do?

- Stores patterns immediately and efficiently, without further analysis.
- Later the neocortex can pick out the important features and memorize those.
- It may take a while for cortex to decide which features are important.
  - Transfer is not immediate.
- Hippocampus is thus a kind of medium-term memory used to train the neocortex.

# An Animal's Limited History

- If 10 fibers out of 1000 can be active at once, that gives  $C(1000,10)$  possible combinations =  $2.6 \times 10^{23}$ .
- Assume a new pattern every 1 ms.
  - Enough combinations to go for  $10^{12}$  years.
- So: assume patterns will not repeat during the lifetime of the animal.
- Very few of the many possible events (patterns) will actually be encountered.
- So events will be well-separated in pattern space, not close together.

# Numerical Constraints

Marr defined a set of numerical constraints to determine the shape of simple memory theory:

1. Capacity requirements
2. Number of inputs
3. Number of outputs
4. Number of synapse states = 2 (binary synapses)
5. Number of synapses made on a cell
6. Pattern of connectivity
7. Level of activity (sparseness)
8. Size of retrieval cue

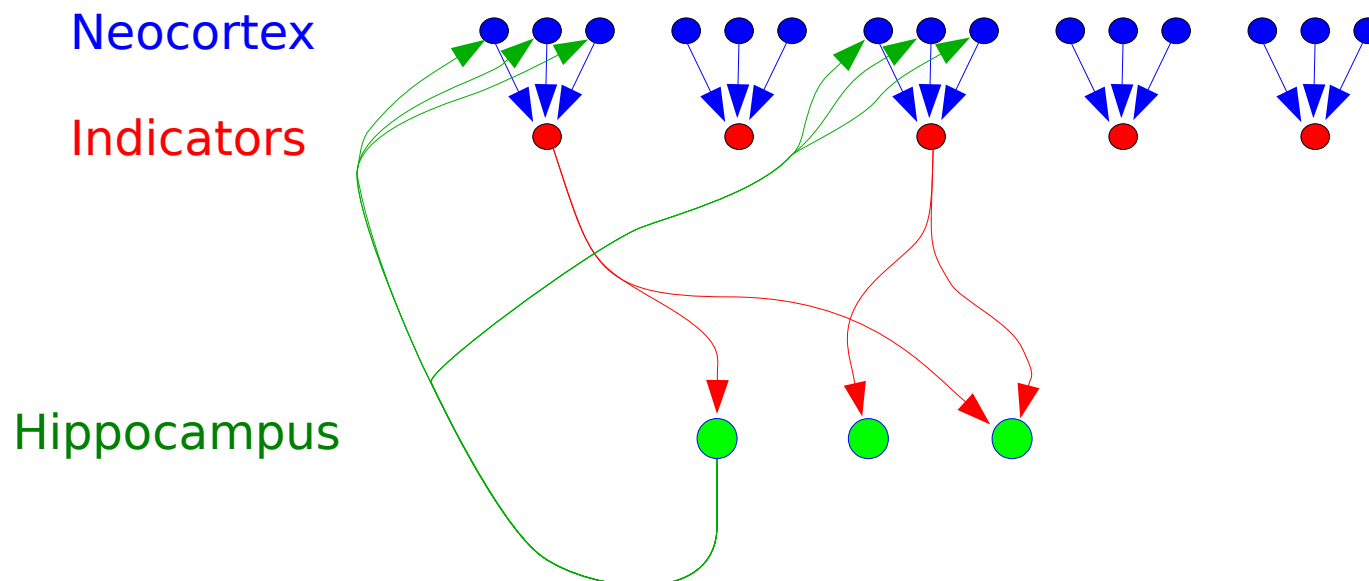
# N1. Capacity Requirements

- A simple memory only needs to store one day's worth of experiences.
- They will be transferred to neocortex at night, during sleep.
- There are 86,400 seconds in a day.
- A reasonable upper bound on memories stored is:

**100,000 events per day**

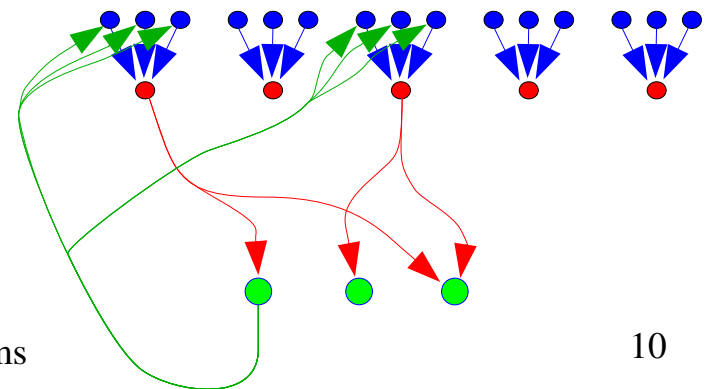
## N2. Number of Inputs

- Too many cortical pyramids ( $10^8$ ): can't all have direct contact with the hippocampus.
- Solution: introduce indicator cells as markers of activity in each local cortical region, about  $0.03 \text{ mm}^2$ .
- Indicator cells funnel activity into the hippocampal system.



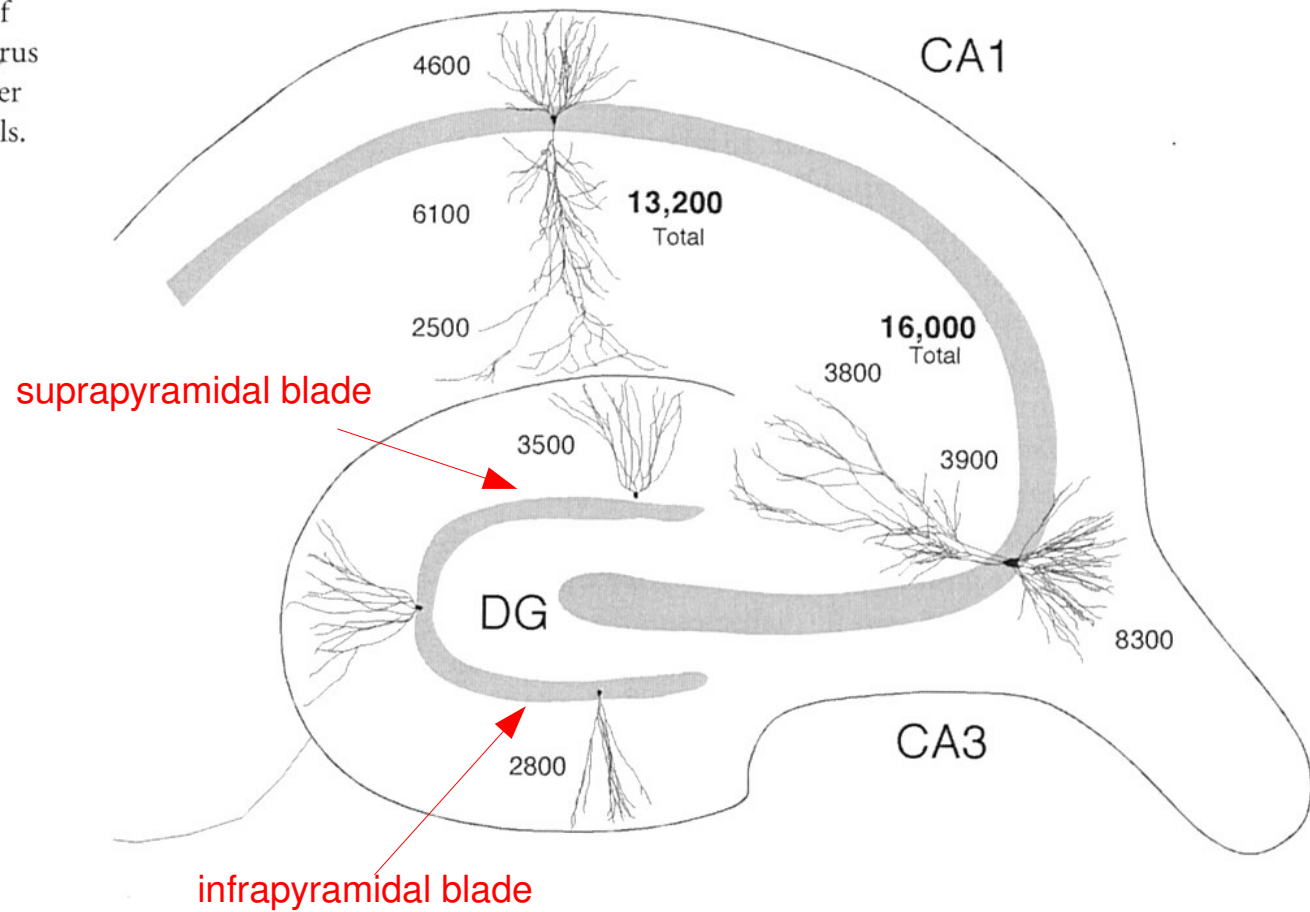
# Indicator Cells

- Indicator cells funnel information into hippocampus.
- Don't we lose information?
  - Yes, but the loss is recoverable if the input patterns aren't too similar (low overlap).
- The return connections from hippocampus to cortex must be direct to all the cortical pyramids, not to the indicator cells.
- But that's okay because there are far fewer hippocampal axons than cortical axons (so there's room for all the wiring), and each axon can make many synapses.



# Dendritic Arborization of Principal Cells

**Figure 3-15.** Dendritic arborization of the principal cells in the rat dentate gyrus (granule cells) and hippocampus proper (pyramidal cells). See the text for details.



# How Many Input Fibers?

- Roughly 30 indicator cells per mm<sup>2</sup> of cortex.
- Roughly 1300 cm<sup>2</sup> in one hemisphere of human cortex, of which about 400 cm<sup>2</sup> needs direct access to simple memory. Thus,

**About 10<sup>6</sup> afferent fibers enter simple memory.**

- This seems a reasonable number.

## N3. Number of Outputs

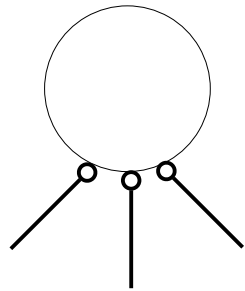
- Assume neocortical pyramidal cells have fewer than  $10^5$  afferent synapses.
- Assume only about  $10^4$  synaptic sites available on the pyramidal cell for receiving output from simple memory.
- Hence, if every hippocampal cell must contact every cortical cell, there can be at most  $10^4$  hippocampal cells in the memory. Too few!
  - If 100,000 memories stored, each memory could only have 10 cells active (based on the constraint that each cell participates in at most 100 memories.) Too few cells for accurate recall.
- Later this constraint was changed to permit  $10^5$  cells in the simple memory.

# N4. Binary Synapses

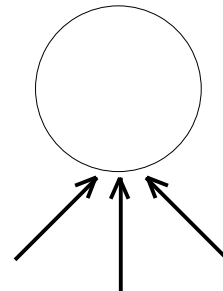
- Marr assumed a synapse is either on or off (1 or 0).
- Real-valued synapses aren't required for his associative memory model to work.
  - But they could increase the memory capacity.
- Assuming binary synapses simplifies the capacity analysis to follow.

# Types of Synapses

- Hebb synapses are binary: *on or off*.
- Brindley synapses have a fixed component in addition to the modifiable component.



Hebb synapses



Brindley synapses

- Synapses are switched to the *on* state by simultaneous activity in the pre- and post-synaptic cells.
- This is known as the Hebb learning rule.

# N5. Number of Synapses

- The number of synapses onto a cell is assumed to be high, but bounded.
- Anatomy suggests no more than 60,000.
- In most calculations he uses a value of  $10^5$ .

## N6. Pattern of Connectivity

- Some layers are subdivided into blocks, mirroring the structure of projections in cortex, and from cortex to hippocampus.
- Projections between such layers are only between corresponding blocks.
- Within blocks, the projection is random.

## N7. Level of Activity

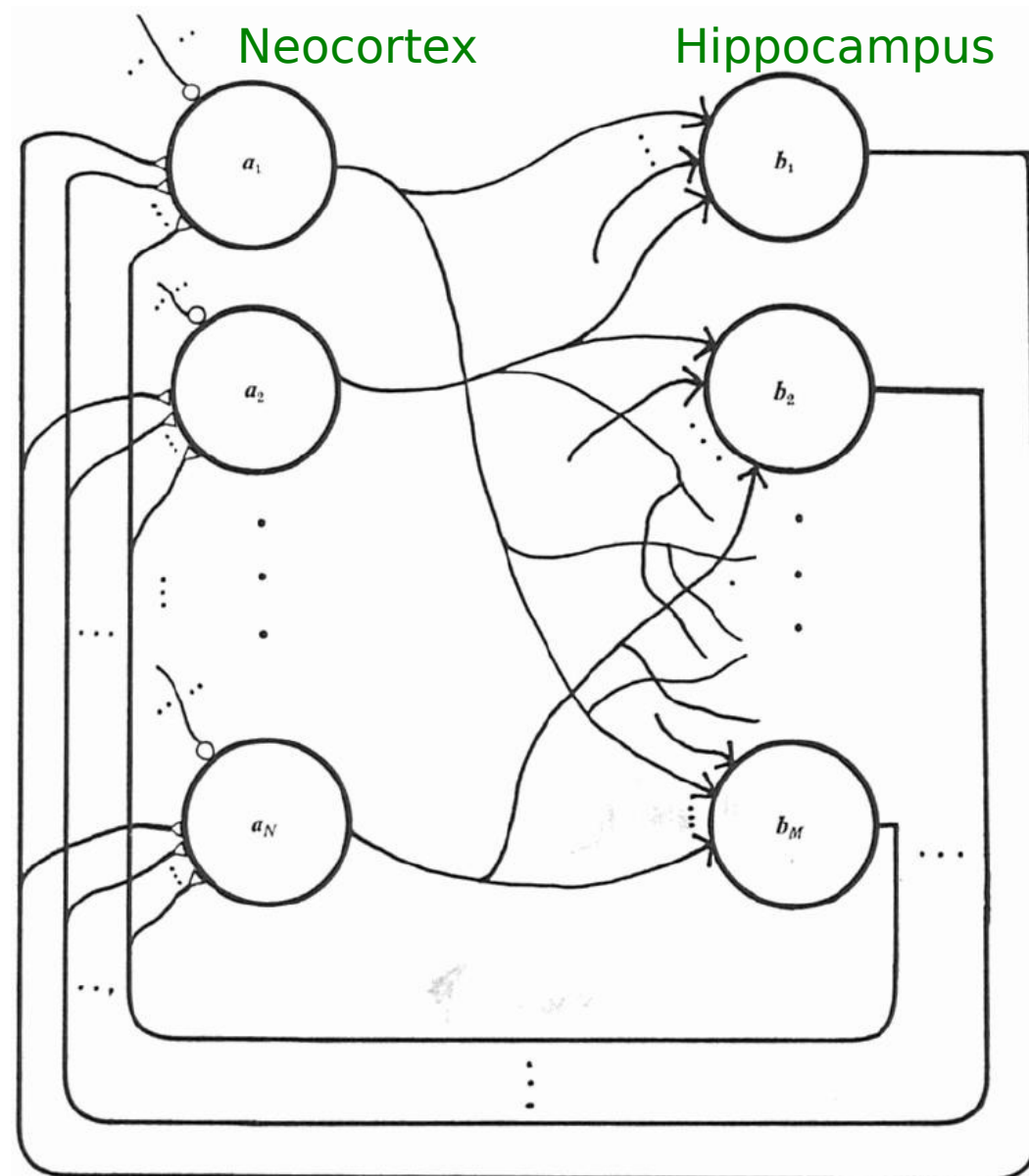
- Activity level (percentage of active units) should be low so that patterns will be sparse and many events can be stored.
- Inhibition is used to keep the number of active cells constant.
- Activity level must not be too low, because inhibition depends on an accurate sampling of the activity level.
- Assume at least 1 cell in 1000 is active.
- That is,  $\alpha > 0.001$ .

## N8. Size of Retrieval Cue

- Fraction of a previously stored event required to successfully retrieve the full event.
- Marr sets this to  $1/10$ .
- This constitutes the minimum acceptable cue size.
- If the minimum cue size is increased, more memories could be stored with the same level of accuracy.

# Marr's Two-Layer Model

- Event  $E$  is on cells  $a_1 \dots a_N$  (the cortical cells)
- Codon formation on  $b_1 \dots b_M$  (evidence cells in HC)
- Inputs to the  $b_j$  use Brindley synapses
- Codon formation is a type of competitive learning (anticipates Grossberg, Kohonen)
- Recurrent connections to the  $a_i$  use Hebb synapses



# Simple Representations

- Only a small number of afferent synapses are available at neocortical pyramids for the simple memory function; the rest are needed for cortical computation.
- In order to recall an event  $E$  from a subevent  $X$ :
  - Most of the work will have to be done within the simple memory itself.
  - Little work can be done by the feedback connections to cortex.
- No fancy transformation from **b** back to **a**.
- Thus, for subevent  $X$  to recall an event  $E$ , they should both activate the same set of **b** cells.

# Recalling An Event

- How to tell if a partial input pattern is a cue for recalling a learned event, or a new event to be stored?
- Assume that events  $E$  to be stored are always much larger (more active units) than cues  $X$  used for recall.
- Smaller pattern means not enough dendritic activation to trigger synaptic modification, so only recall takes place.

# Codon Formation

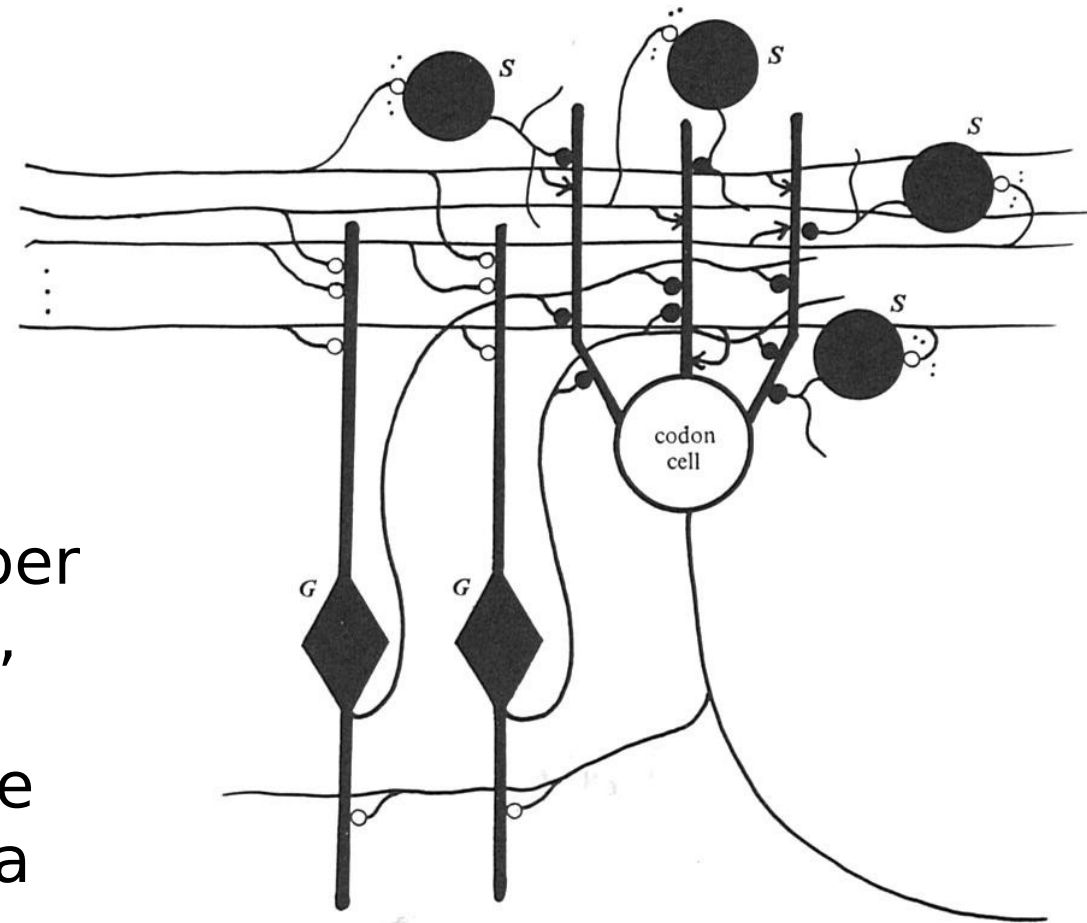
- Memory performance can be improved by orthogonalizing the set of key vectors.
  - The **b** cells do this. How?
- Project the vector space into a higher dimensional space.
- Each output dimension is a conjunction of a random  $k$ -tuple of input dimensions (so non-linear).
- In cerebellum this was assumed to use fixed wiring. In cortex it's done by a learning algorithm.
- Observation from McNaughton concerning rats:
  - Entorhinal cortex contains about  $10^5$  projection cells.
  - Dentate gyrus contains  $10^6$  granule cells.
  - Hence, EC projects to a higher dimensional space in DG.

# Codon Formation

- For each input event  $E$ , different  $\mathbf{b}$  cells will receive different amounts of activation.
- Activation level depends on which  $\mathbf{a}$  cells connect to that  $\mathbf{b}$  cell.
- We want the pattern size  $L$  to be roughly the same for all events.
- Solution: choose only the  $L$  most highly activated  $\mathbf{b}$  cells as the simple representations for  $E$ .
- How to do this?
  - Adjust the thresholds of the  $\mathbf{b}$  cells so that only  $L$  remain active.

# Inhibition to Control Pattern Size

- S and G cells are inhibitory interneurons.
- S cells sample the input lines and supply feed-forward inhibition to the codon cells.
- G cells' modifiable synapses track the number of patterns learned so far, and raise the inhibition accordingly. They sample the codon cell's output via an axon collateral.



# Threshold Setting

- Two factors cause the activation levels of **b** cells to vary:
  - 1) Amount of activity in the **a** cells (not all patterns are of the same size, due to partial cues)
  - 2) Number of potentiated synapses from **a** cells onto the **b** cell. This value gradually increases as more patterns are stored.
    - More cells can become active as more weights are set.
- Solution:
  - 1) S-cells driven by codon cell afferents compute an inhibition term based on the total activity in the  $a_i$  fibers. Assumes no synapses have been modified.
  - 2) G-cells driven by codon cell axon collaterals use negative feedback to compensate for effects of weight increases.
- Together, S and G cells provide subtractive inhibition to maintain a pattern size of L over the **b** units.

# Recall From a Subevent

- If subevent  $X$  is fully contained in  $E$ , the best retrieval strategy is to lower the codon threshold until roughly  $L$  of the **b** cells are active.
- But if  $X$  only partially overlaps with  $E$ , some spurious input units will have synapses onto codon units. A better strategy is for codon cells to take into account the fraction  $f$  of their  $A$  active synapses that have been modified by learning (meaning they are part of some previously-stored pattern).
- Unmodified synapses that are active during recall can only be a source of noise.
- Thus, a **b** cell should only fire if a sufficient proportion  $f$  of its active synapses have been modified, meaning they are part of at least one stored pattern — perhaps the correct one,  $E$ .

# Recall From a Subevent

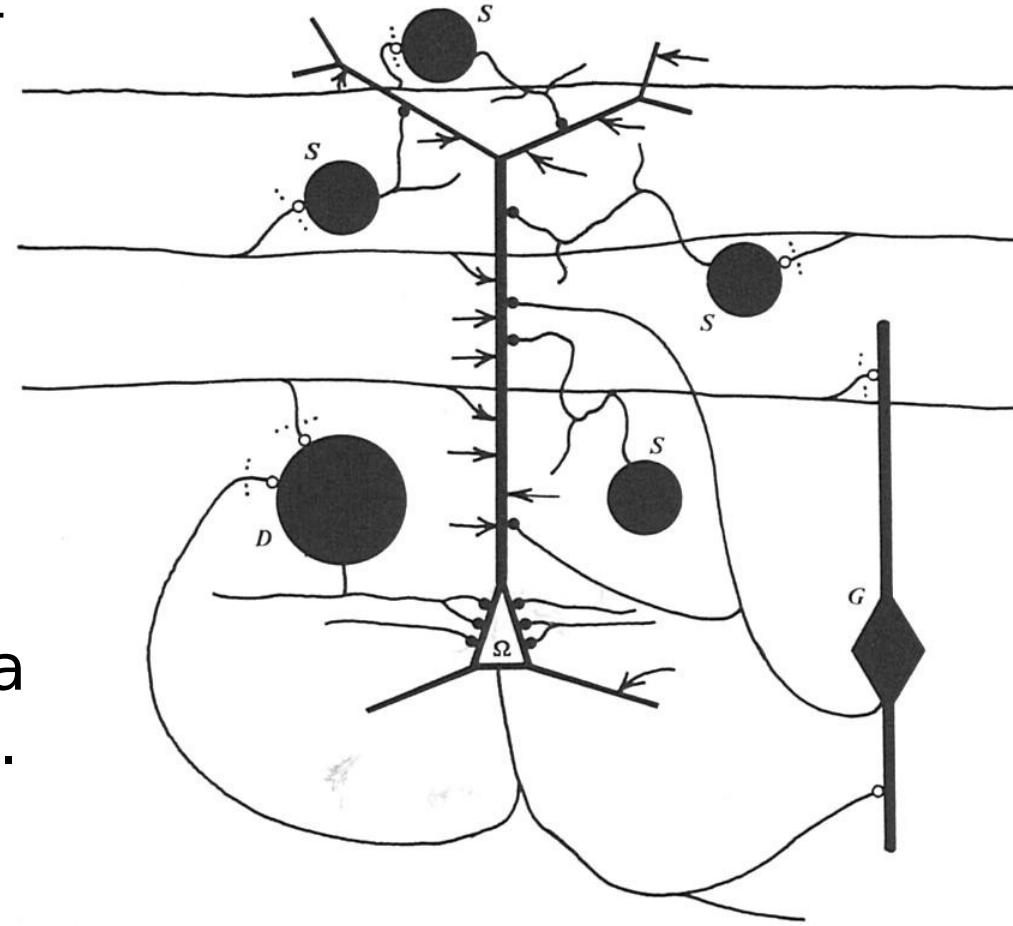
- A cell should only fire if it's being driven by enough modified synapses.
- $A$  = number of active synapses.
- $f$  = fraction of synapses that have been modified.
- The cell's division threshold is equal to  $fA$ .
- Let  $S$  be the summed activation of the cell:

$$S = \sum_i a_i w_i$$

- The cell should fire if  $S > fA$ , or  $S / (fA) > 1$ .

# D-Cells

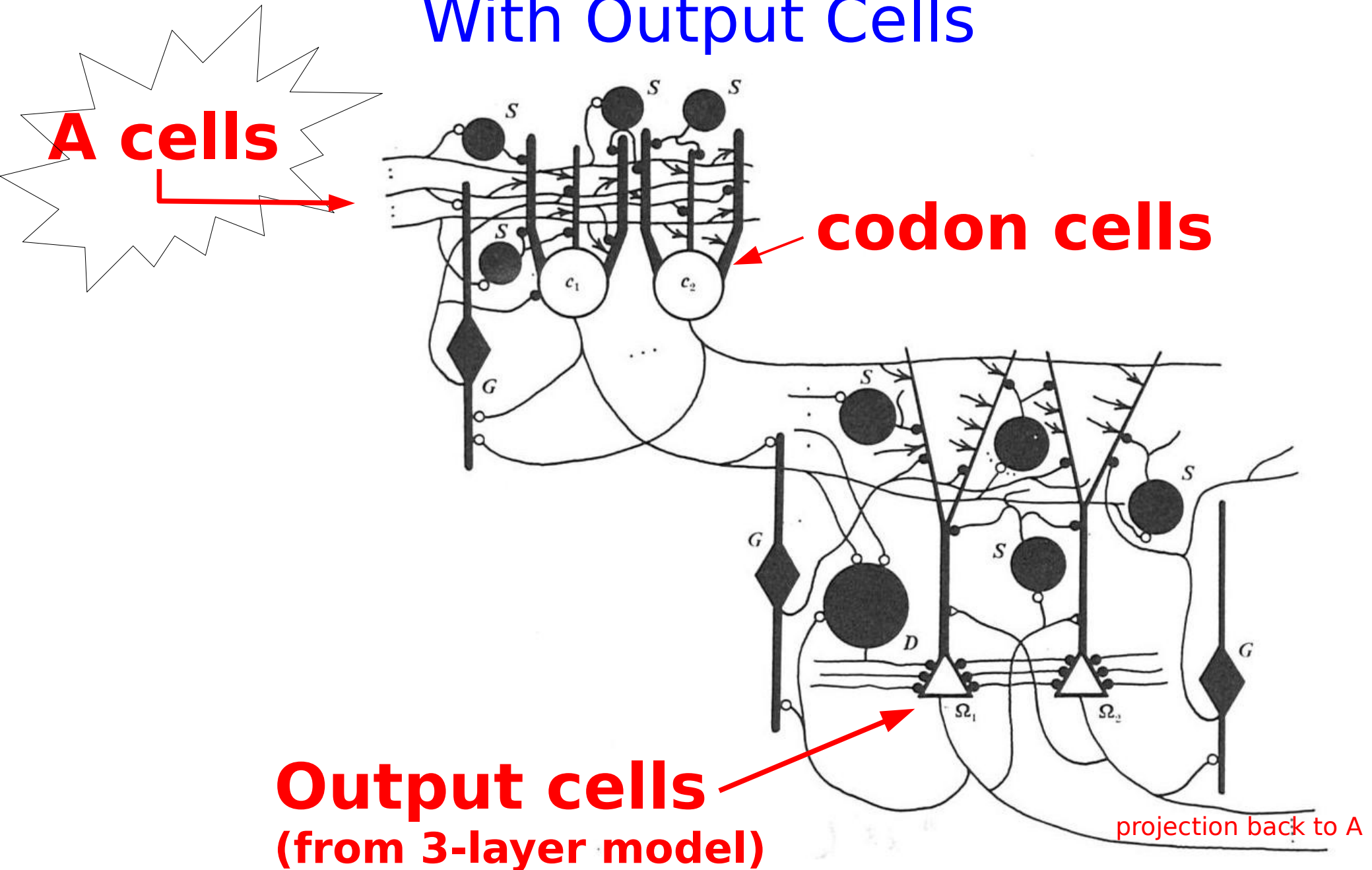
- D cells compute  $fA$  and pass it as an inhibitory input to the pyramidal cells.
- D cells apply their inhibition directly to the cell body, like basket cells in cerebellum.
- This type of inhibition causes a division instead of subtraction.
- McNaughton: division can be achieved by shunting inhibition, e.g., the chloride-dependent  $GABA_A$  channel.



# Dual Thresholds

- Cells have two separate thresholds:
  - The absolute threshold  $T$ , controlled by inhibition from S and G cells, should be close to the pattern size  $L$ , but must be reduced when given a partial cue.
  - The division threshold  $fA$ , controlled by inhibition from D cells.
- Marr's calculations show that both types of thresholding are necessary for best performance of the memory.
- How to set these thresholds? No procedure is given.
  - Willshaw & Buckingham try several methods, e.g., *staircase strategy*: start with small  $f$  and large  $T$ . Gradually reduce  $T$  until enough cells are active, then raise  $f$  slightly and repeat.

# 3 Layer Model: A Simple Memory With Output Cells



# Inadequacy of the Simple Model

- Assume that  $N = 10^6$   $a_i$  afferents.
- Assume each neocortical pyramid can accept  $10^4$  synapses from the  $b_j$  cells.
- Assume upper bound of 200 learned events per cell, due to limitation on number of afferent synapses. (Marr derived this from looking at Purkinje cells in cerebellum.)
  - Use 100 events/cell as a conservative value.
- If capacity  $n = 10^5$  events, and each **b** cell participates in 100 of them, then activity  $\alpha = 10^{-3}$ . With  $10^4$  **b** cells, only 10 can be active per event.
  - Too few for reliable representation. Threshold setting would be too difficult with such a small sample size.

# What's Wrong With This Argument?

- The simple model is inadequate because the activity level is too low: only 10 active units per stored event.
- But this is because Marr assumes only  $10^4$  evidence (codon) cells. Why?
  - Limited room for afferent synapses back to the cortical cells.
- This is based on the notion that every evidence (codon) cell must connect back to *every* cortical cell.
- Later in the paper he relaxes this restriction and switches to  $10^5$  evidence cells.

# Combinatorics 1: Permutations

- How many ways to order 3 items: A, B, C?

\_\_\_\_\_

- Three choices for the first slot.
- Two choices left for the second.
- One choice left for the third.

  B        A        C  

- Total choices =  $3 \times 2 \times 1 = 3! = 6$ .

# Combinatorics 2: Choices

- How many ways to choose 2 items from a set of 5?

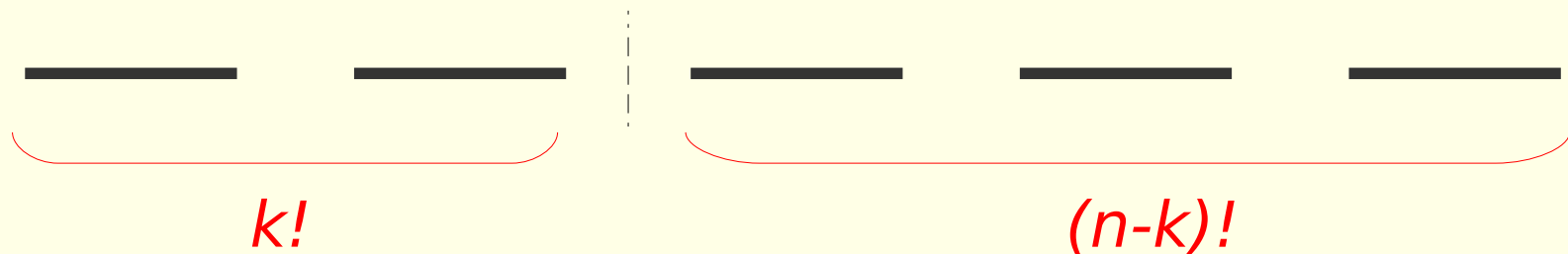
In formal notation, what is the value of  $\binom{5}{2} = C(5,2)$  ?

- Five choices for first item. Four choices for the second.
- Permutations of the chosen item are equivalent: combination B,E is the same as combination E,B
- So total ways to choose two items is  $(5 \times 4)/(2!) = 10$ .
- Since  $5! = 5 \times 4 \times 3 \times 2 \times 1$ , we can get  $5 \times 4$  from  $5!/3!$

$$\binom{5}{2} = \frac{5!}{3! \cdot 2!} = \frac{5!}{3! \cdot 2!}$$

# Choices (continued)

- How many ways to choose  $k=2$  items from  $n=5$  ?
- Allocate 5 slots giving  $n! = 120$  permutations:



- All permutations of the  $k$  chosen items are equivalent, so divide by  $k! = 2$ .
- All permutations of the  $(n-k)$  unchosen items are equivalent, so divide by  $(n-k)! = 6$ .

$$\binom{n}{k} = \frac{n!}{k! \cdot (n-k)!}$$

# Review of Probability

- Suppose a coin has a probability  $z$  of coming up heads.
- The probability of tails is  $(1-z)$ .
- What are the chances of seeing  $h$  heads in a row?

$$z^h$$

- What are the chances of seeing exactly  $h$  heads in a row, followed by exactly  $t$  tails?

$$z^h \cdot (1-z)^t$$

- What about seeing exactly  $h$  heads total in  $N$  tosses?

$$\binom{N}{h} \cdot z^h \cdot (1-z)^{(N-h)}$$

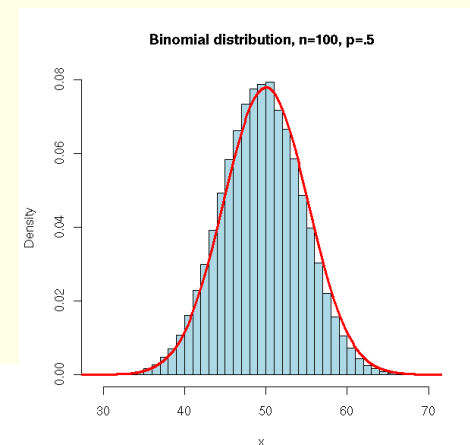
# Binomial Distribution

- How many heads should we expect in  $N=100$  tosses of a biased ( $z=0.2$ ) coin?
  - Expected value is  $E\langle h \rangle = Nz = 20$ .
- What is the probability of a particular sequence of tosses containing exactly  $h$  heads?

$$P\left[\langle t_1, t_2, \dots, t_N \rangle\right] = z^h \cdot (1-z)^{N-h}$$

- The probability of getting exactly  $h$  heads in any order follows a binomial distribution:

$$\text{Binomial}(N; z)[h] = \binom{N}{h} \cdot z^h \cdot (1-z)^{N-h}$$



# Marr's Notation

$P_i$	Population of cells.
$N_i$	Number of cells in population $P_i$
$L_i$	Number of active cells for a pattern in $P_i$
$\alpha_i$	Fraction of active cells: $L_i/N_i$
$R_i$	Threshold of cells in $P_i$
$S_i$	Number of afferent synapses of a cell in $P_i$
$Z_i$	Contact probability: likelihood of synapse from cell in $P_{i-1}$ to $P_i$
$\Pi_i$	Probability that a particular synapse in $P_i$ has been modified
$E\langle x \rangle$	Expected (mean) value of $x$
$n$	Number of stored memories

# Response to an Input Event

- Assume afferents to  $P_i$  distribute uniformly with probability  $Z_i$ .
- $L_{i-1}$  = number of active afferents.
- What is the expected pattern size in this population?

$$E\langle L_i \rangle = N_i \sum_{r=R_i}^{L_{i-1}} \binom{L_{i-1}}{r} \cdot (Z_i)^r \cdot (1-Z_i)^{L_{i-1}-r}$$

- What do the terms in this formula mean?

# Response to an Input Event

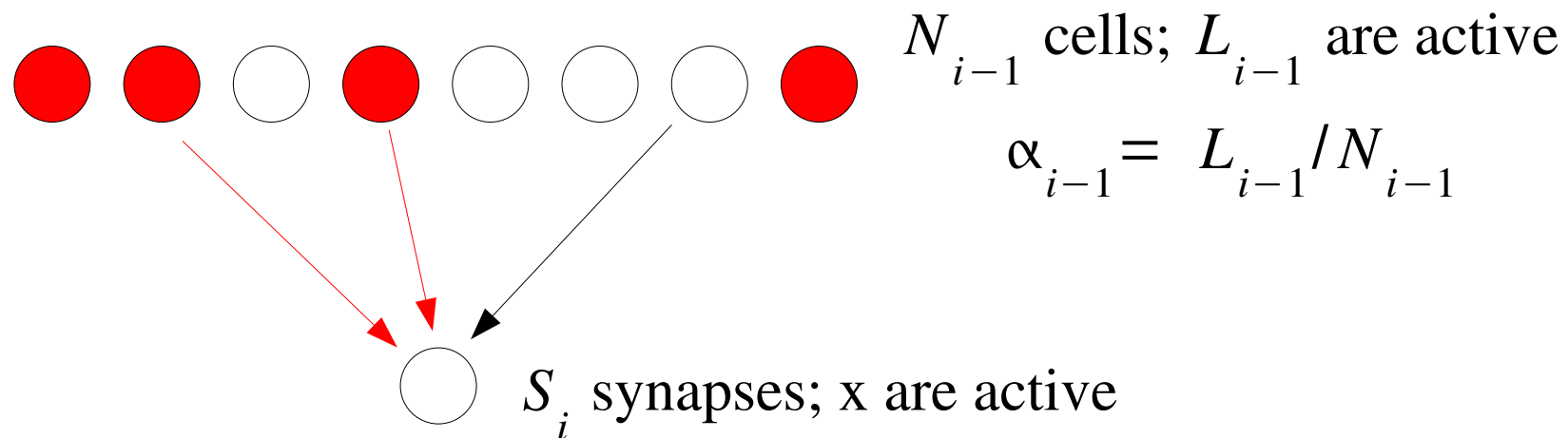
$$E \langle L_i \rangle = N_i \sum_{r=R_i}^{L_{i-1}} \binom{L_{i-1}}{r} \cdot (Z_i)^r \cdot (1-Z_i)^{L_{i-1}-r}$$

probability a unit has EXACTLY  $r$  active input fibers

probability a unit has AT LEAST  $R_i$  active input fibers (so is active)

- One term of the summation is the probability that a cell will receive an input of size exactly  $r$ , given  $L_{i-1}$  active fibers in the preceding layer.
- $r$  is number of active fibers;  $R_i$  is the threshold.
- Must have  $r \geq R_i$  in order for the layer  $i$  cell to fire. Also,  $r \leq L_{i-1}$ , the pattern size for layer  $i-1$ .
- Large  $R_i$  keeps us on the tail of the binomial distribution.
- The value of  $\alpha_i = L_i / N_i$  will be small.

# Counting Active Synapses



Number of active synapses  $x$  is binomially distributed.

$$P(x) = \binom{S_i}{x} \cdot (\alpha_{i-1})^x \cdot (1 - \alpha_{i-1})^{S_i - x}$$

$$E\langle x \rangle = \alpha_{i-1} S_i$$

# Constraint on Modifiable Synapses

Activity  $\alpha_{i-1} = L_{i-1}/N_{i-1}$ .

Proportion of synapses active at each active cell of  $P_i$  is at least equal to the mean  $\alpha_{i-1}$  because the active cells are on the tail of the distribution.

The amount by which it exceeds this decreases as  $S_i \alpha_{i-1}$  grows.

Probability that a (pre,post)-synaptic pair of cells is simultaneously active is  $\alpha_{i-1} \alpha_i$ .

After  $n$  events, probability that a particular synapse of  $P_i$  is facilitated is:

$$\Pi_i = 1 - (1 - \alpha_{i-1} \alpha_i)^n$$

If  $\alpha_{i-1}$  is small, then  $\alpha_{i-1} \alpha_i$  is smaller, so this gives roughly

$$\Pi_i \approx 1 - \exp(-n \alpha_{i-1} \alpha_i)$$

because for small  $\epsilon$ ,  $(1 - \epsilon)^n \approx \exp(-n \epsilon)$

# Constraint on Modifiable Synapses

- For modifiable synapses to be useful, not all should be modified after  $n$  events are stored.
  - Otherwise we could just make all of them fixed.
- Suppose we want at most  $1 - (1/e)$  of them to be modified, which is about 63%.

$$\begin{aligned}\Pi_i &\leq 1 - (1/e) \\ &= 1 - \exp(-1) \\ &\approx 1 - \exp(-n \alpha_{i-1} \alpha_i)\end{aligned}$$

- Thus we have computational constraint C1:

$$n \alpha_{i-1} \alpha_i \leq 1$$

# Condition for Full Representation

- Activity in  $P_i$  must provide an adequate representation of the input event.
- Weak criterion of adequacy: change in input fibers (active cells in  $P_{i-1}$ ) should produce a change in the cells that are firing in  $P_i$ .
- Cells in  $P_i$  just above threshold  $\rightarrow$  losing one input will shut off the cell.

# Condition for Full Representation

Probability  $P$  that an arbitrary input fiber doesn't contact any active cell of  $P_i$  (so  $P_i$  doesn't care if it's shut off) is:

$$(1-\epsilon)^n \approx \exp(-n\epsilon)$$
$$P = (1 - Z_i)^{L_i}$$
$$P \approx \exp\left(-\alpha_i N_i \cdot S_i / N_{i-1}\right)$$

$$L_i = \alpha_i N_i$$
$$Z_i = S_i / N_{i-1}$$

Let's require  $P < e^{-20}$  (about  $2 \times 10^{-9}$ ). Then with a little bit of algebra we have computational constraint C2:

$$S_i \alpha_i N_i \geq 20 N_{i-1}$$

# Summary of Constraints

- To store lots of memories, patterns must be sparse.

$$\text{Constraint C1: } n \alpha_i \alpha_{i-1} < 1$$

- For the encoding to always distinguish between input patterns, outputs must change in response to any input change.
  - There must be enough units and synapses to assure this.

$$\text{Constraint C2: } S_i \alpha_i N_i \geq 20 N_{i-1}$$

- Assumes output cells are just above threshold so losing 1 input fiber will turn them off. They must be on the tail of the binomial distribution for this to hold.

# What's Next?

- Move to a larger, three-layer, block-structured model.
- Add recurrent connections.
- Derive conditions under which recurrent connections improve recall results.
- Map this model onto the circuitry of the hippocampus.

# The Three-Layer Model

Noisy cue X

Pattern C induced by collaterals

Representation of event  $E_0$

50,000 units

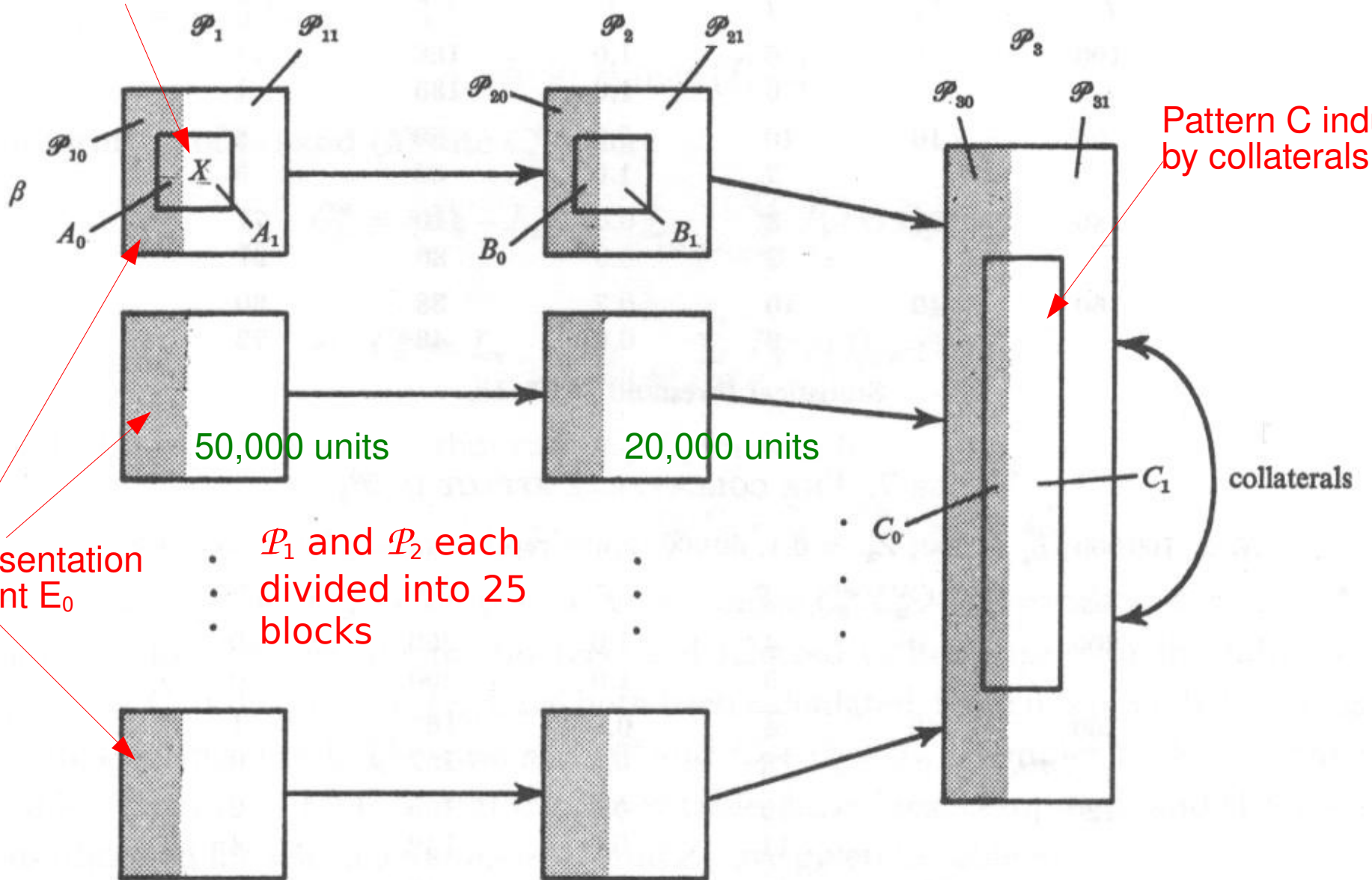
20,000 units

$\mathcal{P}_1$  and  $\mathcal{P}_2$  each divided into 25 blocks

$\mathcal{P}_1: 1.25 \times 10^6$   
Input Cells

$\mathcal{P}_2: 500,000$   
Evidence Cells (Codons)

$\mathcal{P}_3: 100,000$   
Output Cells



# Willshaw and Buckingham's Model

- Willshaw and Buckingham (1990) implemented a simplified 1/100 scale model of Marr's architecture
- Didn't bother partitioning  $\mathcal{P}_1$  and  $\mathcal{P}_2$  into blocks.
- $\mathcal{P}_1 = 8000$  cells,  $\mathcal{P}_2 = 4000$  cells, and  $\mathcal{P}_3 = 1024$  cells.
- For two-layer version, omit  $\mathcal{P}_2$ .
- Performance was similar for both architectures.
- Memory capacity was roughly 1000 events.
  - Partial cue of 8% gave perfect retrieval 66% of the time.
  - In two-layer net, 16% cue gave perfect retrieval 99% of the time.
  - In three-layer version, 25% cue gave 100% perfect retrieval.

# Willshaw and Buckingham's Three-Layer Model Parameters

$$\begin{array}{lll} a_1=0.03 & a_2=0.03 & a_3=0.03 \\ N_1=8000 & N_2=4000 & N_3=1024 \\ & S_2=1333 & S_3=2666 \end{array}$$

calculated:

$$\begin{array}{lll} L_1=240 & L_2=120 & L_3=30 \\ & Z_2=0.17 & Z_3=0.67 \\ & \Pi_2=0.41 & \Pi_3=0.41 \end{array}$$

# Assessment of Marr's Theory

- Strong points:
  - Sparse, topographic connectivity: more biologically realistic.
  - Multiple inhibitory mechanisms: subtraction and division.
  - Predicts when recurrent collaterals will help retrieval.
  - Anticipated many important findings: LTP, division operations, information transfer during sleep.
- Weak points:
  - Ignores the trisynaptic circuit ( $EC \rightarrow DG \rightarrow CA3 \rightarrow CA1$ ). It seems like  $\mathcal{P}_1$  is neocortex,  $\mathcal{P}_2$  is EC or DG, and  $\mathcal{P}_3$  is CA3.
    - $\mathcal{P}_2$  has more cells than  $\mathcal{P}_3$ , like DG has more than CA3
    - But mossy fiber projection to CA3 is too sparse for  $\mathcal{P}_2 \rightarrow \mathcal{P}_3$
  - Claim that three layers of cells are necessary was unjustified.
  - Unanswered question: how are memories transferred from hippocampus to the neocortex?