

# Lecture 15: Generalization, geometry, and causality

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# Roadmap

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- ▶ Representations that support invariant prediction have a distinctive parallel geometry that is attested in neural recordings, supporting the idea that the brain organizes its representational architecture to support causal generalization.
- ▶ Several mechanisms for learning invariant predictors, with connections to dreaming and oscillatory plasticity rules.

# Recognizing cows with different backgrounds

Convolutional neural networks trained for object recognition struggle when cows appear in unusual backgrounds like beaches. Because cows appear mainly in pastures, the networks learn to rely on information contained in the background—a spurious correlation that should be ignored. How does the brain know what to ignore?



(A) **Cow: 0.99**, Pasture: 0.99, Grass: 0.99, No Person: 0.98, Mammal: 0.98



(B) No Person: 0.99, Water: 0.98, Beach: 0.97, Outdoors: 0.97, Seashore: 0.97



(C) No Person: 0.97, **Mammal: 0.96**, Water: 0.94, Beach: 0.94, Two: 0.94

[Beery et al 2018]

# Causality and generalization

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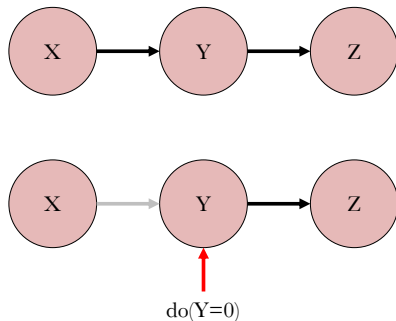
- ▶ The essence of the problem is causality: we understand intuitively that pastures don't cause cows to appear in images. Only cows cause the appearance of cows!
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- ▶ Because cows often graze in pastures, the presence of a pasture makes it more likely that a cow will be there. In other words, we could think of the pasture as a cause of the cow's presence, which in turn causes its appearance in a photo of the pasture.
- ▶ If the farmer arrives and leads the cow back to its stable, the pasture will still appear in the photo but the cow will not. The farmer has “intervened” on the causal structure, severing the correlation between the pasture and the cow appearance.

# Causal model

(Top) The model represented as a directed graph, where nodes represent variables and arrows represent causal dependencies. (Bottom) Intervening on variable  $Y$ , represented by  $\text{do}(Y = 0)$ , sets  $Y$  to 0 and removes the arrow from  $X$  to  $Y$ . In terms of the example,  $X$  is the pasture,  $Y$  is the cow presence, and  $Z$  is the cow's appearance in the photo. The farmer's intervention is represented by the "do" operator.



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- ▶ In contrast,  $Y$  is a cause of  $Z$  because its effect on  $Z$  is invariant across interventions on  $X$  (a cow will appear in a photo if it's present, even if the photo is taken on a beach).

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- ▶ A system that learns invariant causal mechanisms will generalize correctly to new contexts, whereas a system that learns spurious correlations will not.
- ▶ We will see how causal invariance is achieved by neural representations with a particular geometry in the high-dimensional space of population activity.
- ▶ This geometry is abstract in the sense that it maintains its structure across contexts, reflecting the underlying causal invariants that are unaffected by interventions on the other variables that collectively comprise the context.

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- ▶ They all have in common the assertion that causal relationships are “law-like” in the sense that they generalize across many contexts.
- ▶ Conversely, contexts are interventions that leave the causal relationships intact.
- ▶ Because each context is associated with a different distribution over observations, knowledge of causal relationships enables a form of “out of distribution” generalization.

## Study question

Why is generalization fundamentally a causal problem rather than a statistical one?

# The trouble with empirical risk minimization

- ▶ Recall the empirical risk minimization setup. We are given a dataset of  $M$  input-label pairs,  $\{x_m, s_m\}_{m=1}^M$ , where  $x_m$  is an input (e.g., an image) and  $s_m$  is its label (e.g., an object category or a continuous feature).

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- ▶ Goal is to find a conditional distribution  $q(s|x)$ , a *predictor*, that minimizes the empirical risk:

$$\hat{L}(q) = \frac{1}{M} \sum_m L(q, s_m, x_m)$$

where  $L(q, s, x)$  is a loss function.

## The trouble with empirical risk minimization

- ▶ Regression example using continuous labels ( $s \in \mathbb{R}$ ) and two continuous inputs ( $x = [x_a, x_b]$ ). Data-generating process ( $s$  is an effect of  $x_a$  and a cause of  $x_b$ ):

$$x_a \sim \mathcal{N}(0, \sigma_a^2), \quad x_b \sim \mathcal{N}(s, \sigma_b^2), \quad s \sim \mathcal{N}(x_a, \sigma_s^2)$$

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- ▶ Ideally, we would like our classifier to correctly identify the causal structure, relying only on  $x_a$  to predict  $s$ .
- ▶ Unfortunately, this will not generally be the case for empirical risk minimization. The Bayes-optimal predictor, obtained by minimizing by the cross-entropy loss  $L(q, s, x) = -\log q(s|x)$ , is the posterior,  $q(s|x) = p(s|x)$ , which in this case is a Gaussian with mean:

$$\hat{s} = \frac{\sigma_b^2}{\sigma_s^2 + \sigma_b^2} x_a + \frac{\sigma_s^2}{\sigma_s^2 + \sigma_b^2} x_b$$

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- ▶ In the limit  $\sigma_b^2 \rightarrow 0$ ,  $x_b$  becomes a deterministic effect of  $s$ , and its coefficient goes to 1, whereas the coefficient for  $x_a$  (the correct causal predictor) goes to 0.

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- ▶ Empirical risk minimization learns a spurious correlation rather than an invariant cause.
- ▶ In essence, the problem is that there is no way to learn invariant causes without some source of variance.

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- ▶ Naively, you might think that you could just pool all the contexts together and apply the standard empirical risk minimization approach. However, this can still lead to fitting spurious correlations if the variance of the non-causal variables is small.
- ▶ Need a predictor that performs well simultaneously in all the contexts, which eliminates non-causal predictors by stress-testing them in contexts where they fail.

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- ▶ To guarantee that causal variables can be identified, the predictor needs access to a sufficiently rich feature representation, such that at least some of the features correspond to causal variables.
- ▶ Feature representation computed by a non-linear encoder  $\phi(x)$ . The label distribution is assumed to be a log-linear decoder:

$$q(s|\phi(x)) \propto \exp[\beta_s + w_s \cdot \phi(x)],$$

where  $w_s$  is a weight vector and we have included a bias term  $\beta_s$ .

# Invariant risk minimization

- ▶ Binary classification, with  $s \in \{1, 2\}$ , with log odds:

$$\log \frac{q(s = 1 | \phi(x))}{q(s = 2 | \phi(x))} = \beta + w \cdot \phi(x)$$

where  $w = w_1 - w_2$  and  $\beta = \beta_1 - \beta_2$ .

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- ▶ When the class-conditional distribution over features is isotropic Gaussian,  $\phi(x) | s \sim \mathcal{N}(\bar{\phi}_s, \sigma^2 \mathbf{I})$ , the Bayes-optimal weight vector is:

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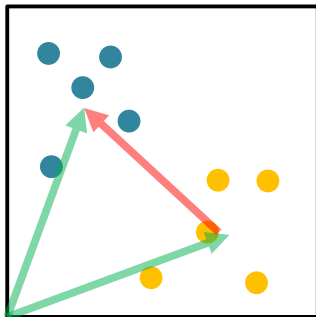
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- ▶ The optimal weight vector is proportional to the *coding direction*—the direction in feature space that maximally separates the two classes, which in this case corresponds to the difference between the class-conditional means.

# Binary classification

The two green arrows represent the vectors pointing at the class-conditional means. The red arrow shows the coding direction obtained by vector subtraction.



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- ▶ This will change the optimal bias, but will *not* change the optimal weight vector.
- ▶ Thus,  $w^*$  defines an invariant predictor: it can be applied across all contexts.

## Context-dependent flexibility

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- ▶ These settings require context-dependent flexibility, where the weight vector is allowed to vary across contexts.
- ▶ But we still want to seek causal invariants that support abstraction across contexts.

## Context-dependent flexibility

- ▶ We relax the invariant risk minimization problem:

$$w_e^* = \operatorname{argmin}_w \hat{L}_e(w) + \lambda \|\bar{w} - w\|^2, \quad \bar{w} = \frac{1}{N} \sum_e w_e$$

where  $\hat{L}_e(w)$  is the empirical risk given weight vector  $w$  in context  $e$ , and  $\bar{w}$  is the average weight vector across all  $N$  contexts.

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- ▶ The second term regularizes each context-dependent weight towards the average weight; the parameter  $\lambda$  controls the strength of this regularization. When  $\lambda$  is large,  $w_e^*$  will tend to be invariant across contexts.

## Context-dependent flexibility

- ▶ Suppose we have two contexts,  $e \in \{1, 2\}$ . If we sum the regularization terms across contexts and apply the Law of Cosines, we get:

$$\|\bar{w} - w_1\|^2 + \|\bar{w} - w_2\|^2 = \|w_1\|^2 + \|w_2\|^2 - 2\|w_1\|\|w_2\|\cos(\theta)$$

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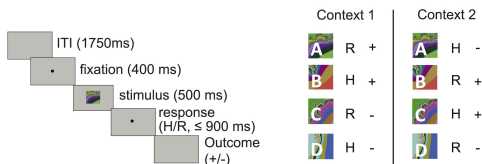
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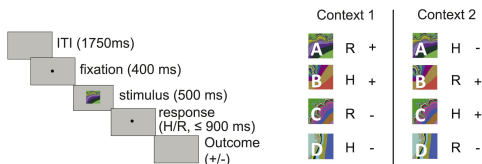
- ▶ The regularizer penalizes both large weights (the first two terms) *and* large angles between the weight vectors.
- ▶ Recall that the optimal unregularized weight vector is proportional to the coding direction (the difference between the class-conditional means). This suggests that if we also optimize an encoder  $\phi(x, e)$  defined jointly over sensory inputs and context, we should find representations where the coding directions are approximately parallel across contexts (i.e.,  $\theta \approx 0$ ).

# Representational geometry in the brain



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- ▶ Context switched every 50-70 trials.

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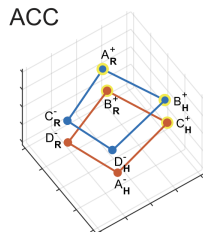
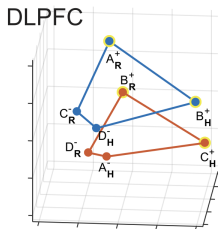
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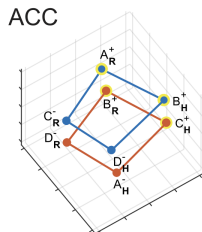
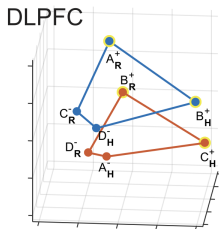
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- ▶ For a linear decoder, this only requires that there is some weight vector  $w$  that separates the correct and incorrect actions for each context.
- ▶ However, this admits spurious correlations that can lead to poor generalization.

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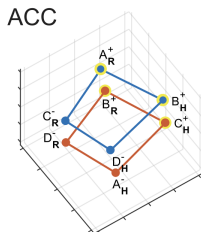
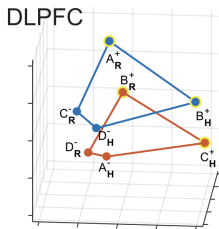
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- ▶ This was indeed the case in the prefrontal cortex and hippocampus: neural representations for the two contexts look like approximately translated copies of one another.
- ▶ Suggests that representations in these areas are optimized for extracting invariant predictors.

## Representational geometry in the brain

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- ▶ Fully factorized case: shattering dimensionality is the rank of the representation matrix  $\Phi$ ; if all the columns (features) are linearly independent, the rank is the number of features. Usually much smaller than the number of possible dichotomies ( $2^M$ ).

# Representational geometry in the brain

- ▶ To achieve greater flexibility, add an “interaction” term  $\psi(x, e)$ :

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- ▶ Thus, sacrificing a small amount of parallelism can enable a huge gain in flexibility.
- ▶ Consistent with observations from Bernardi et al [2020]: PFC and hippocampus exhibited high shattering dimensionality, despite also having high parallelism.

# Offline mechanisms for causal learning

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# Offline mechanisms for causal learning

- ▶ Our treatment of invariant prediction has relied on an extrinsic source of variance: an agent has to actually experience different contexts in order to discover invariant predictors.
- ▶ Fortunately, the brain is not truly a prisoner of experience—it can synthesize counterfactual data, providing itself with an alternative source of variance.

# Learning from randomized data: a function of dream sleep?

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- ▶ Originally developed in robotics, where systems are first trained on simulated data before being deployed in the real world.

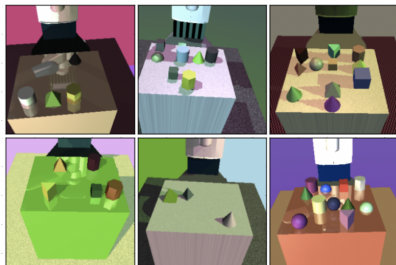
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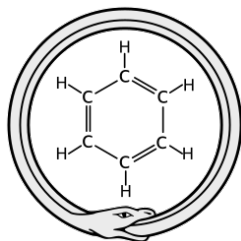
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- ▶ Domain randomization addresses this problem by randomizing aspects of the simulator (e.g., viewpoint, color, texture) that are independent of the underlying physical laws. This provides the source of variance needed to learn invariant predictors.

# Images generated from a randomized simulator



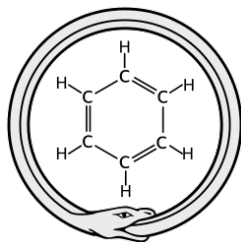
[Tobin et al 2017]

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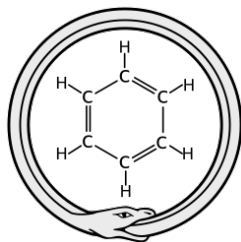
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- ▶ Dreams often occur in response to repetitive task training. People trained on a virtual maze navigation task reported task-related mental imagery during sleep, and the occurrence of this imagery predicted later performance on a test with random starting positions [Wamsley et al 2010].
- ▶ Suggests that dreaming does not merely improve memory—it also improves generalization

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- ▶ If sleeping generates variation unavailable during waking, it could ameliorate overfitting by breaking spurious correlations.

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- ▶ These findings are in broad agreement with the hypothesis that offline activation generates a form of domain randomization.

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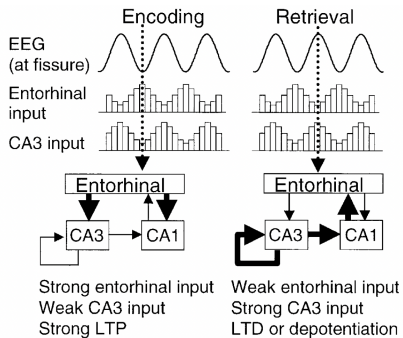
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- ▶ This phase also coincides with the strongest recurrent activity in subfield CA3 and greater long-term depression at hippocampal synapses.
- ▶ Opposite profile at the trough of the theta oscillation: lower firing rates, stronger feedforward activity from entorhinal cortex, and greater long-term potentiation.

# Separation of encoding and retrieval phases by theta oscillations in the hippocampus

EEG: electroencephalography; LTP: long-term potentiation; LTD: long-term depression. CA3 and CA1 are subfields of the hippocampus. The fissure is an anatomical landmark at the input to CA1.



[Hasselmo et al 2012]

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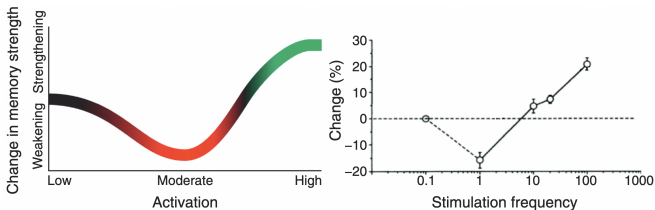
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- ▶ When inhibition is reduced near the peak, spurious correlations are revealed and then weakened by anti-Hebbian plasticity.

# The nonmonotonic plasticity hypothesis

(Left) Hypothetical relationship between activation level and plasticity. Strongly activated memories are strengthened; moderately activated memories are weakened. (Right) Long-term depression for intermediate-frequency stimulation and long-term potentiation for high-frequency stimulation.



[Ritvo et al 2019; Kirkwood et al 1996]

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- ▶ Causal invariance under intervention: a causal relationship between variables is the structure that remains intact when other aspects of the world change.
- ▶ This principle ties causality tightly to generalization, since invariance is the abstraction needed to make predictions in new contexts.

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- ▶ Parallelism cannot be perfect: some non-linear interaction between these variables (mixed selectivity) is needed to endow the system with a sufficiently rich feature set for generalization across many different prediction problems.
- ▶ Domain randomization by dreaming or contrastive learning by oscillatory plasticity can generate sources of variance needed for learning invariant predictors, escaping the prison of experience.