

A latent variable model of configural conditioning

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Similarity & Discrimination in Animal Learning

- **Similarity:** How do animals respond to novel patterns of stimuli?
- **Discrimination:** How do animals learn to discriminate between overlapping patterns of stimuli?
- We recognize these issues as tradeoff between **generalization** and **data-fitting**



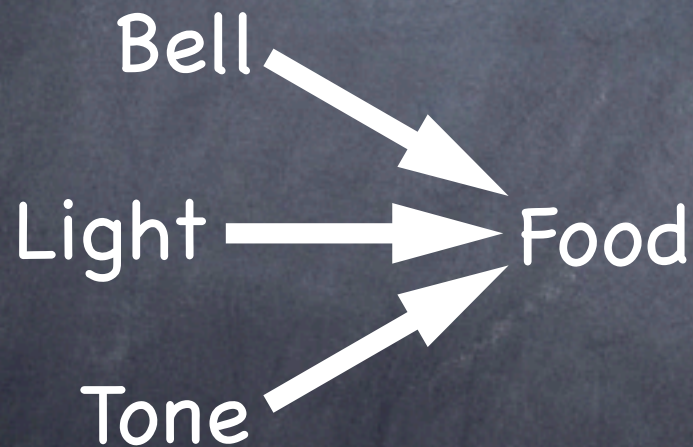
**DIGESTIVE JUICES
BEGIN TO WORK**

**SALIVA BEGINS
TO FLOW**

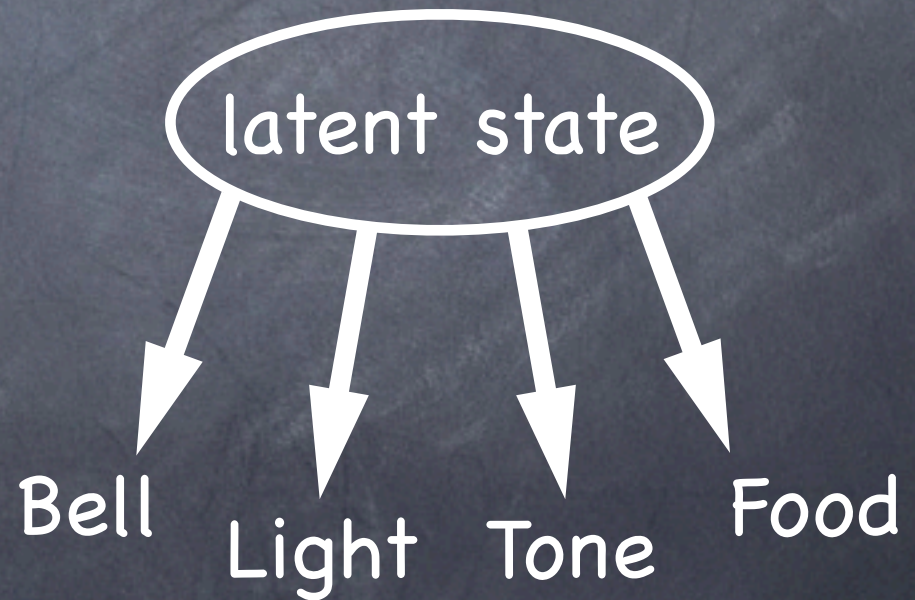
TONGUE OUT

Perspectives on modeling conditioning

Discriminative



Generative



Models of Conditioning:

Rescorla-Wagner (1972)

- Predicts reinforcement intensity as a linear function of stimuli, $X = [A \text{ (light)}, B \text{ (bell)}, \dots]$.

$$V = \sum_i w_i X_i$$

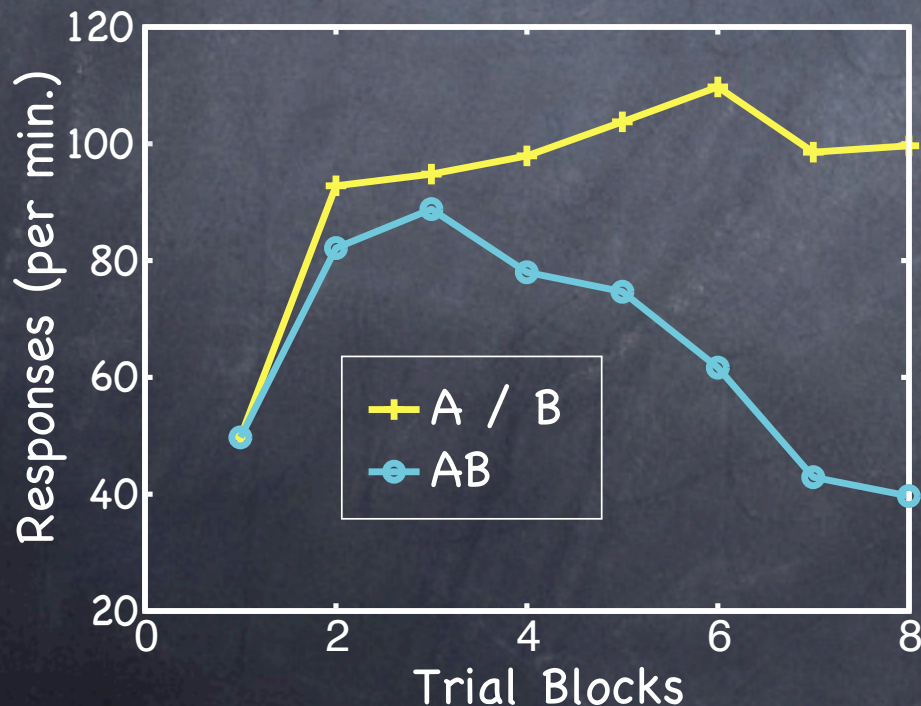
- Learning rule is gradient descent on prediction error

$$\Delta w_i = \alpha_i \beta (r - V) X_i$$

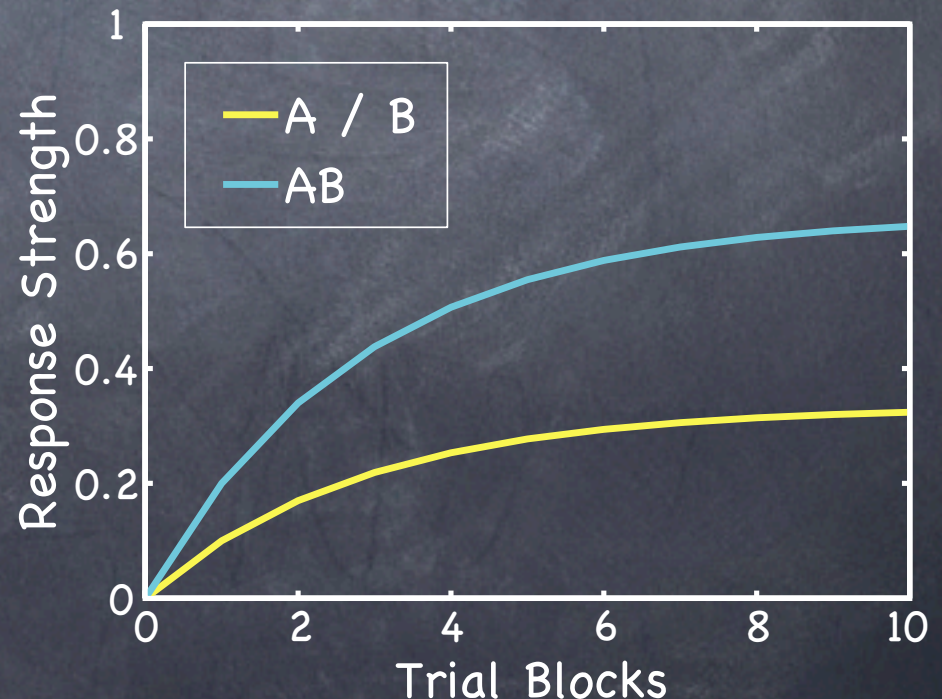
Stimulus configurations

- **Configural conditioning:** discrimination and generalization between patterns of stimuli.

Training: (**XOR**) A+ B+ AB-



Rescorla-Wagner:



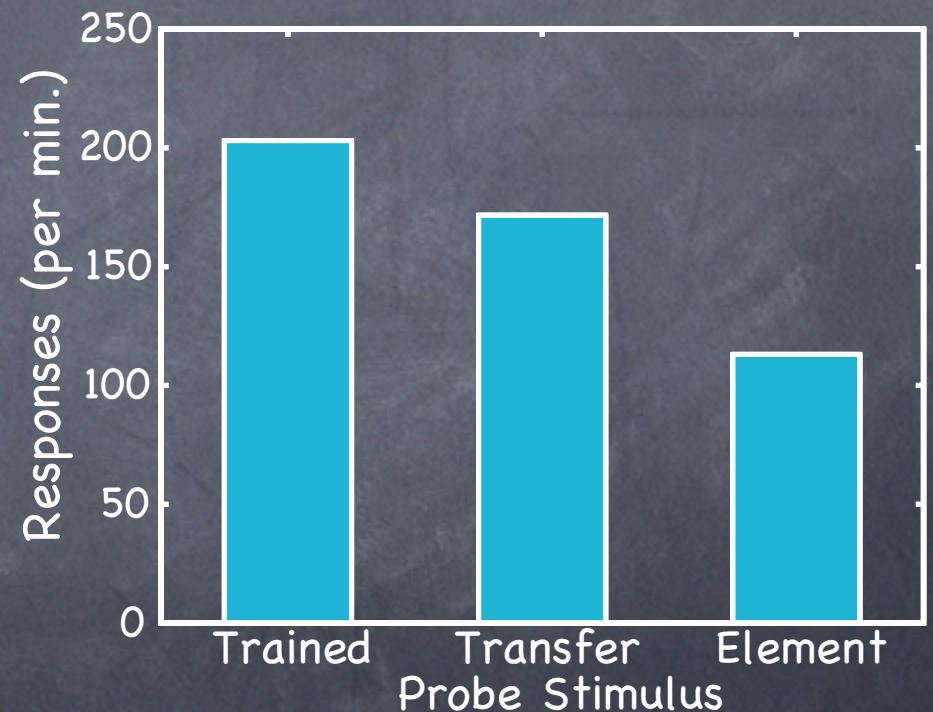
Modeling Configurations

- Two dominant perspectives:
 1. **Added elements RW**, [W&R, 1972]
 2. **Configural model** [Pearce, 1994]
- Augment stimulus representation with **configural unit**. eg. XOR: **$X=[A,B,AB]$** .
- Which units are active? Observe **AB**
 - [W&R 1972]: All units present. **$X=[A=1,B=1,AB=1]$**
 - [Pearce 1994]: Graded activation by generalization rule. **$X=[A=.5,B=.5,AB=1]$**

Expt. 1 Paired Compounds

[Rescorla, 2003]

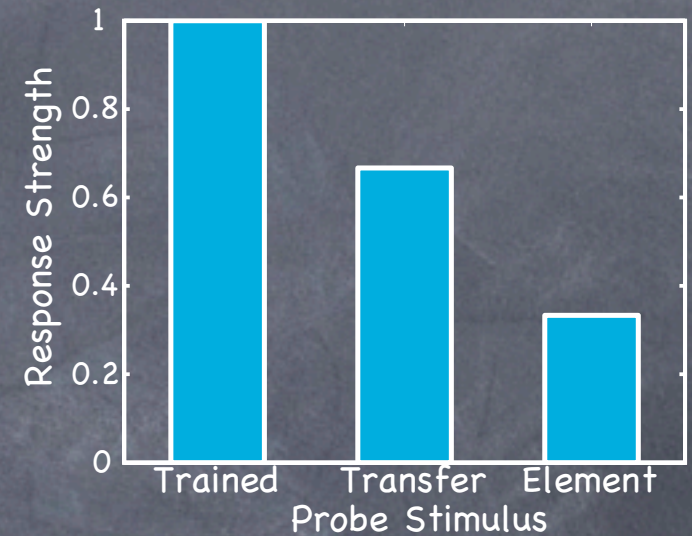
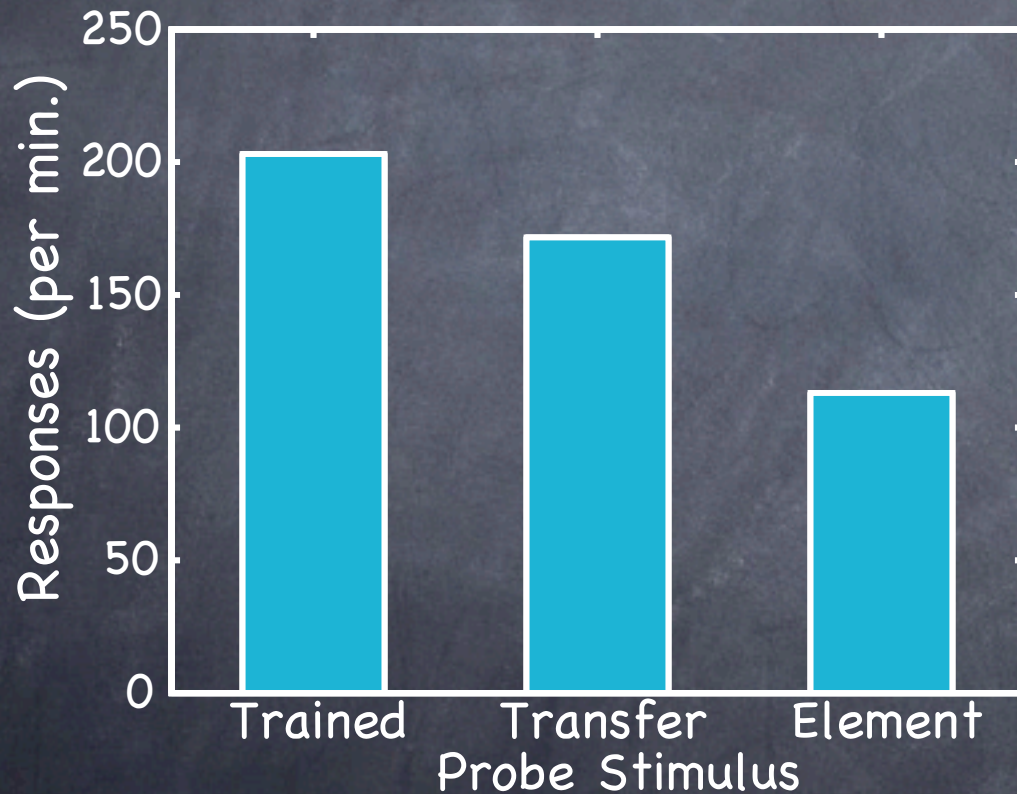
| Training Trials: | Test Trials: |
|------------------|----------------------|
| AB+ CD+ | Trained: AB, CD |
| | Transfer: AD, BC |
| | Elements: A, B, C, D |



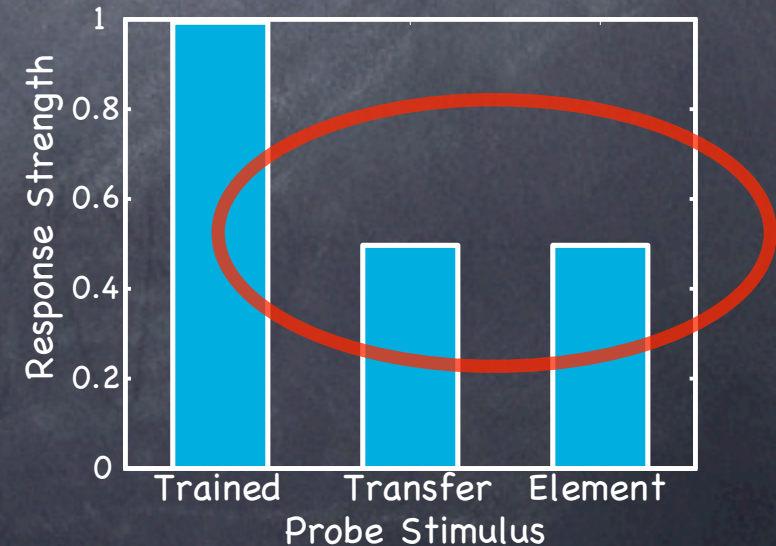
Modeling Paired Compounds

Rescorla-Wagner:

Training: AB+ CD+



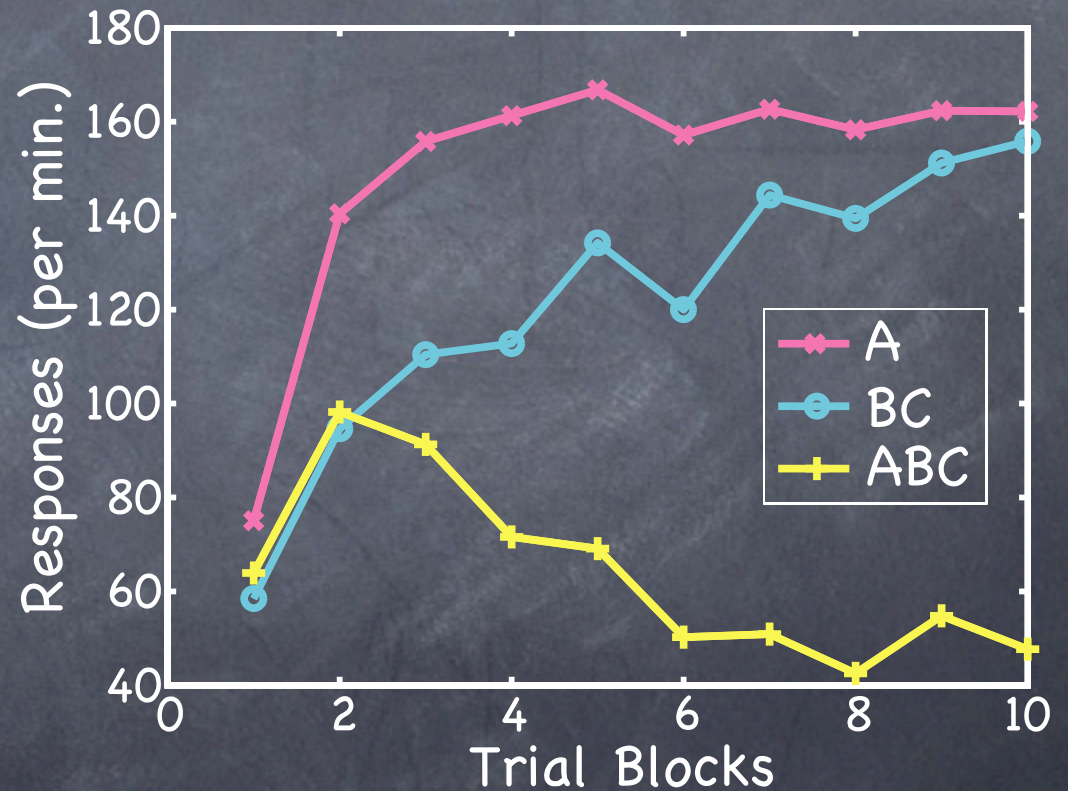
Pearce:



Expt. 2 Asymmetric XOR

[Redhead & Pearce, 1995]

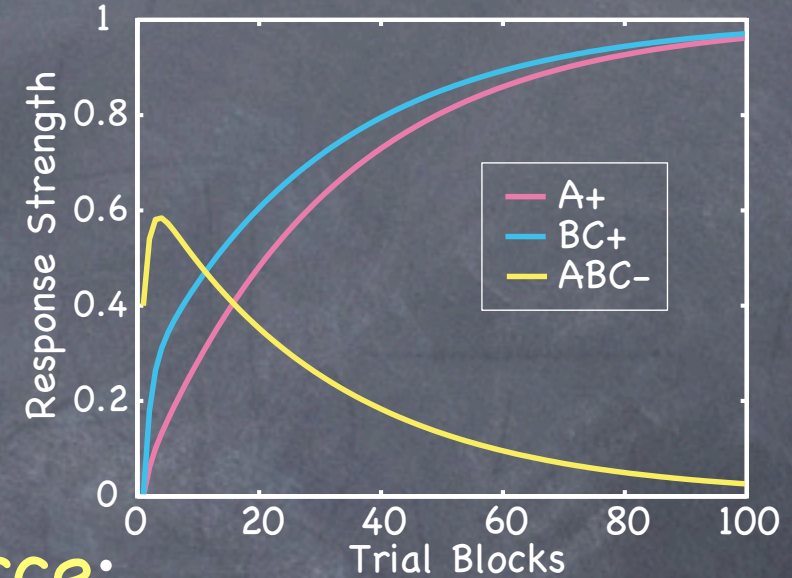
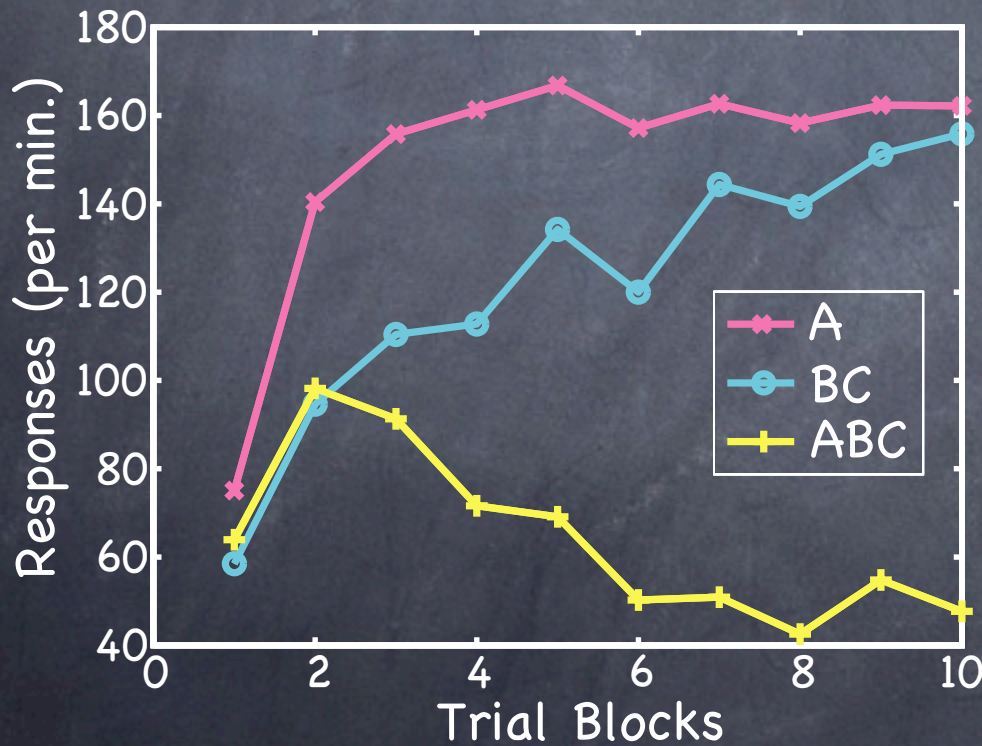
| Training Trials: | Test Trials: |
|------------------|--------------|
| A+ | A |
| BC+ | BC |
| ABC- | ABC |



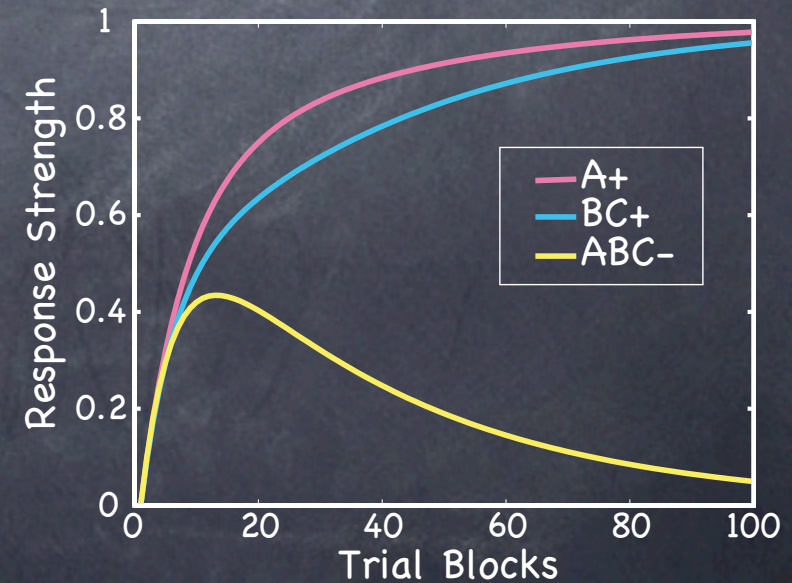
Modeling Asymmetric XOR

Rescorla-Wagner:

Training: A+ BC+ ABC-



Pearce:

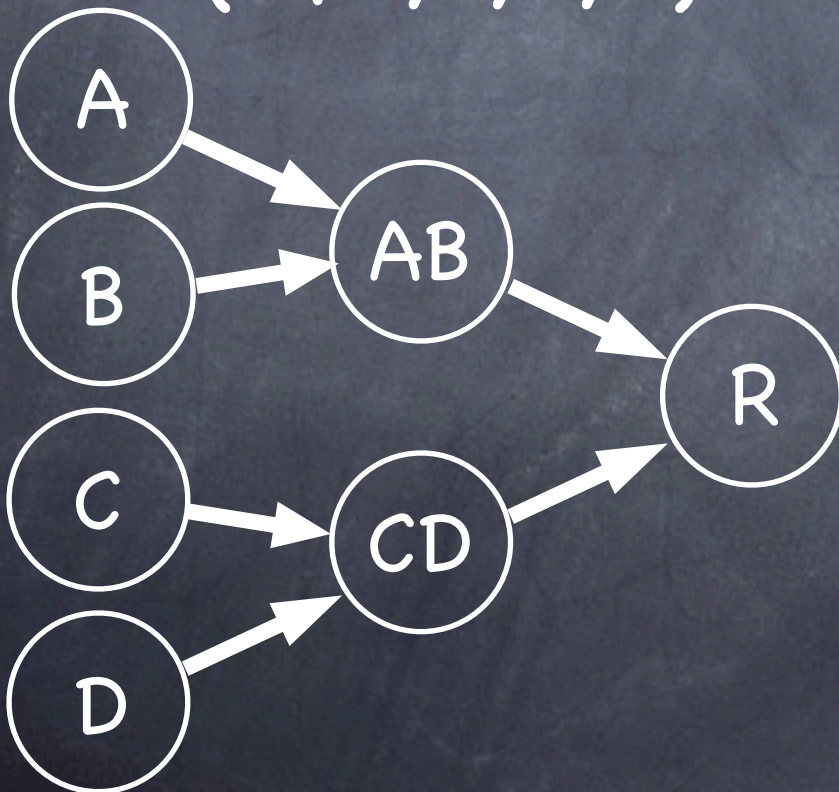


Issues in Modeling Configural Conditioning

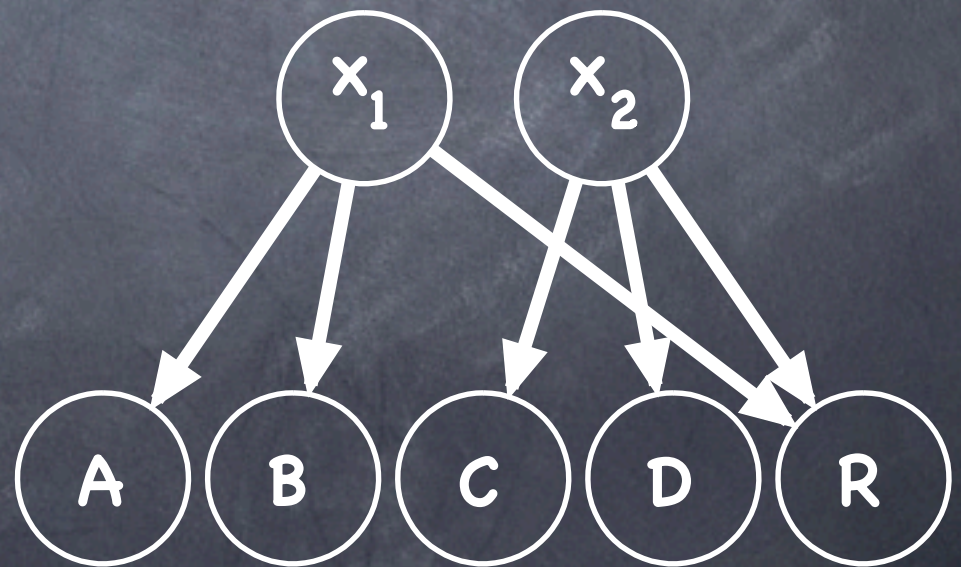
- How do we choose between the two models?
- **Similarity**: How to measure similarity between patterns of stimuli?
- **Discrimination**: How do we choose a representation that is flexible enough?
- A formal Bayesian approach can guide us

Perspectives on modeling conditioning

Discriminative
 $P(R|A,B,C,D)$



Generative
 $P(R,A,B,C,D)$

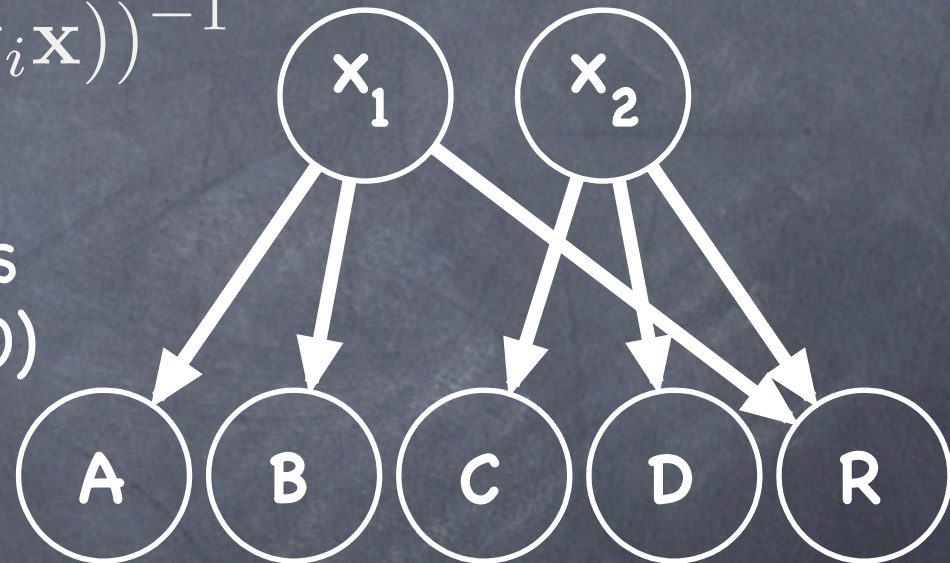


A latent variable model

- Generative model: sigmoid belief network.

$$P(S_i | \mathbf{x}) = (1 + \exp(-\mathbf{w}_i \mathbf{x}))^{-1}$$

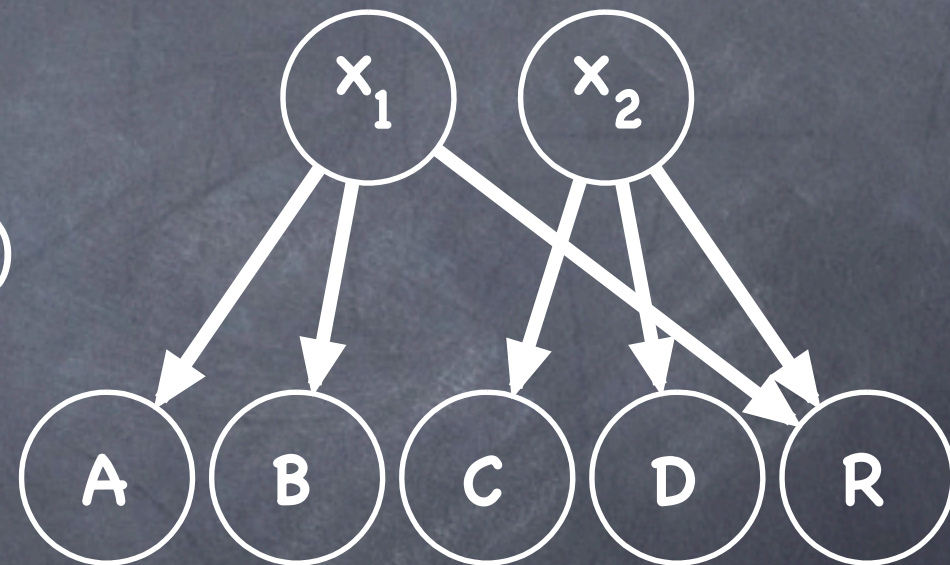
- Stimuli and Latent variables are binary (on = 1, off = 0)



- Latent variables \Rightarrow configural unit
 - correlate stimuli

Model Inference

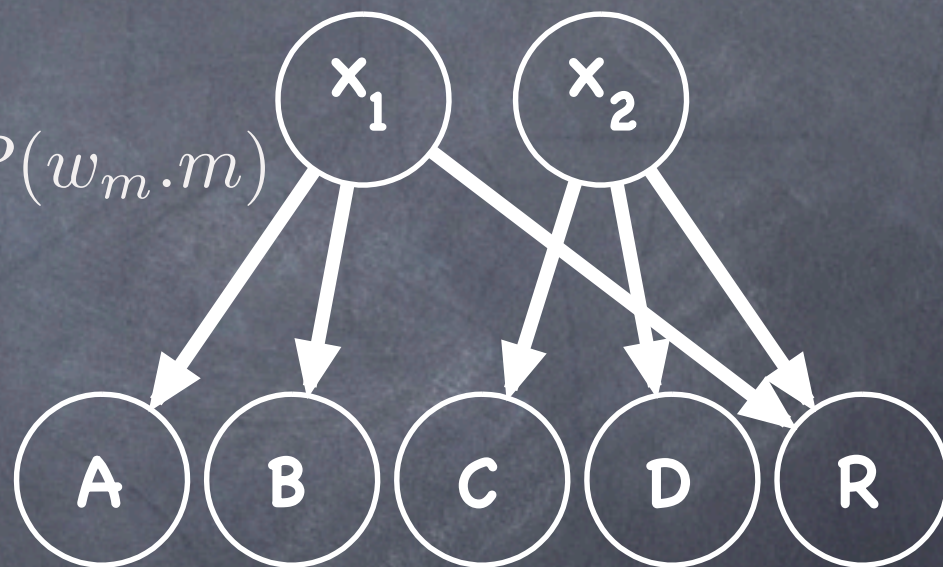
- Learning: $P(w,m \mid D)$
- Prediction: $P(R \mid \text{Stim}, D)$



Learning in the L.V. model

- Learning = Bayesian inference over **weights** & **model structure** conditional on training data

$$P(w_m, m | \mathcal{D}) \propto P(\mathcal{D} | w_m, m) P(w_m, m)$$



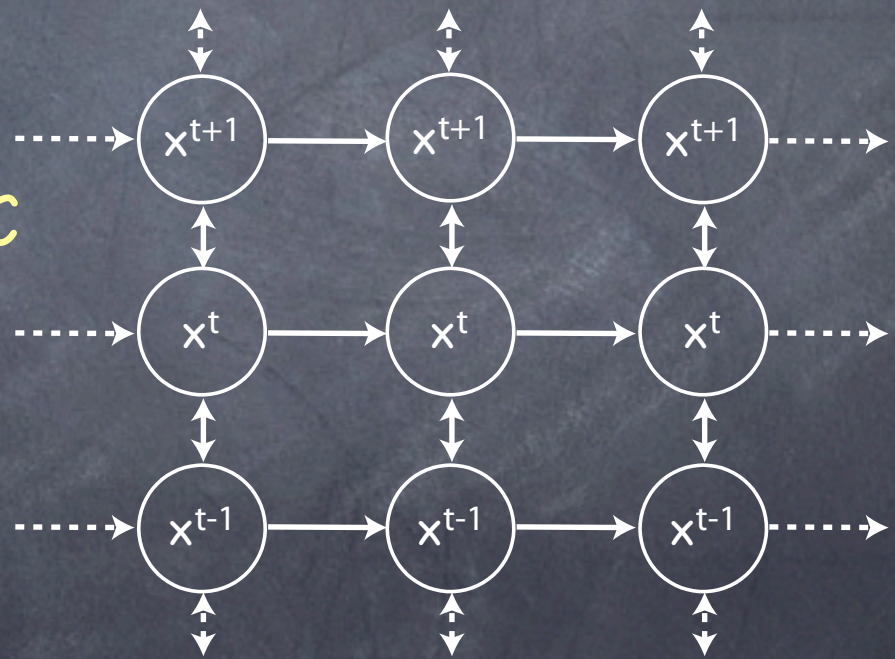
- Latent variable is unknown and unwanted so we compute the marginal likelihood:

$$P(\mathcal{D} | w_m, m) = \prod_t \sum_x \prod_i P(S_{t,i} | x, w_m, m) P(x | w_m, m)$$

Approximate inference

- Inference is analytically intractable: use reversible-jump MCMC

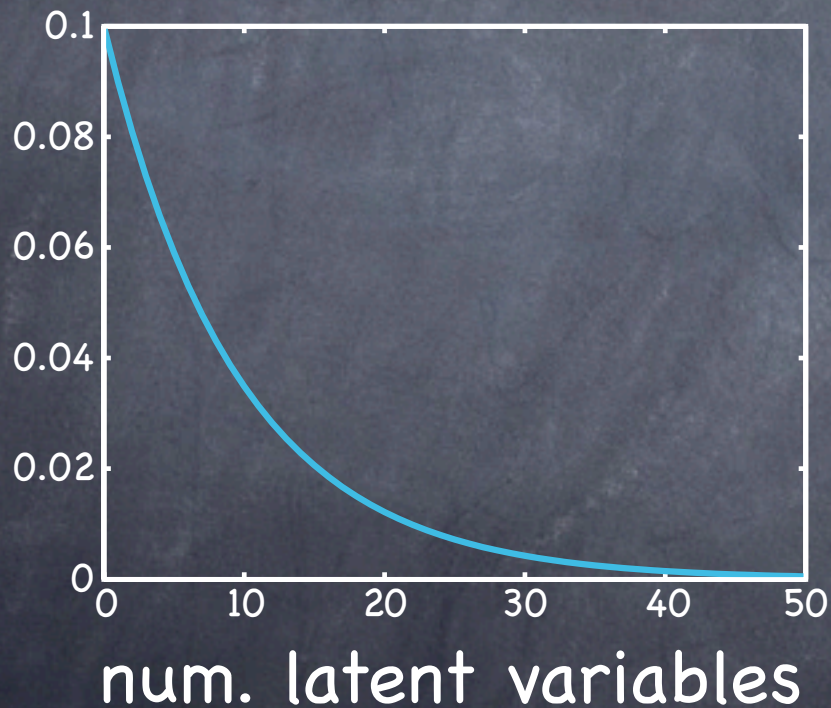
- reversible-jump mixes slowly: **Exchange MCMC** method to help



L.V. model priors

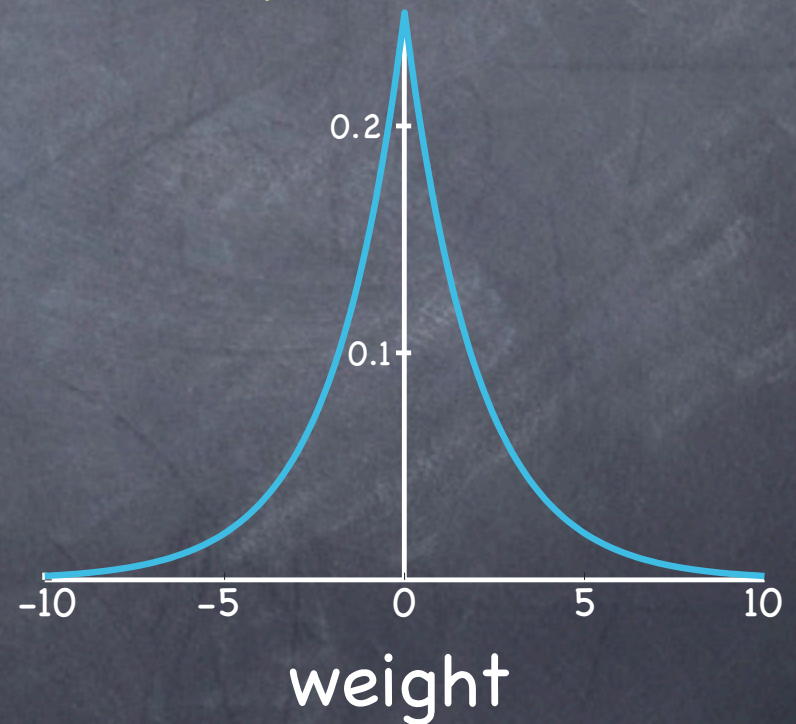
Prior over number of latent variables:

Geometric(0.1)



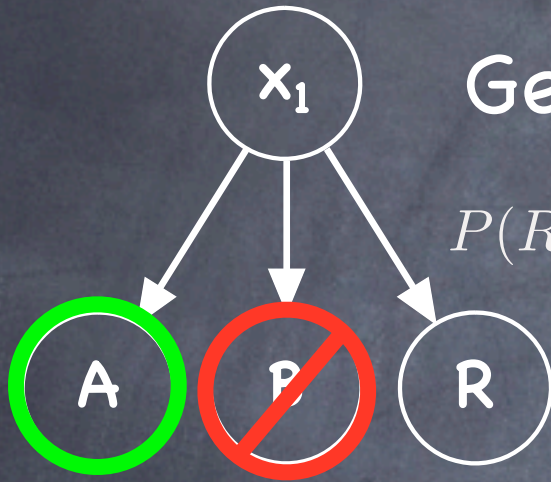
Prior over weight magnitudes:

Laplace(2.0)



Additional assumption: **Stimuli are a priori rare**

Prediction



Generalization \Rightarrow inference over latents

$$P(R | A, \bar{B}, m, \mathbf{w}_m) = \sum_{x_1} P(R | x_1, m, \mathbf{w}_m) P(\mathbf{x} | A, \bar{B}, m, \mathbf{w}_m)$$

$$P(x_1 | A, \bar{B}, m, \mathbf{w}_m) \propto P(A | x_1, m, \mathbf{w}_m) P(\bar{B} | x_1, m, \mathbf{w}_m) P(x_1 | m, \mathbf{w}_m)$$

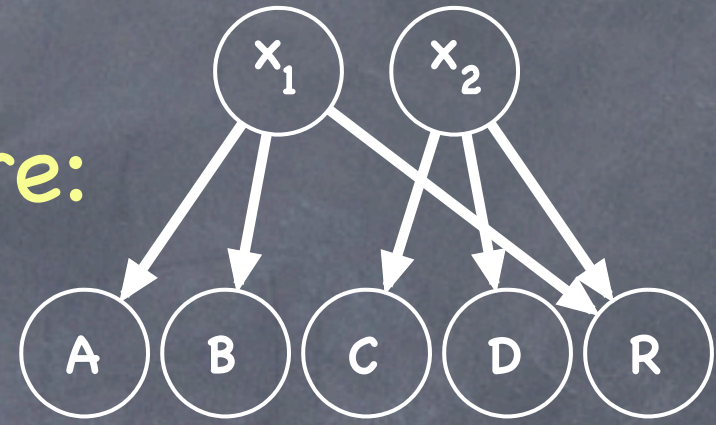
Posterior reinforcement prediction: marginalize over choice of weights and model structure.

$$P(R | Stim, m, \mathcal{D}) = \int P(R | Stim, m, \mathbf{w}_m, \mathcal{D}) P(\mathbf{w}_m | m, \mathcal{D}) d\mathbf{w}_w$$

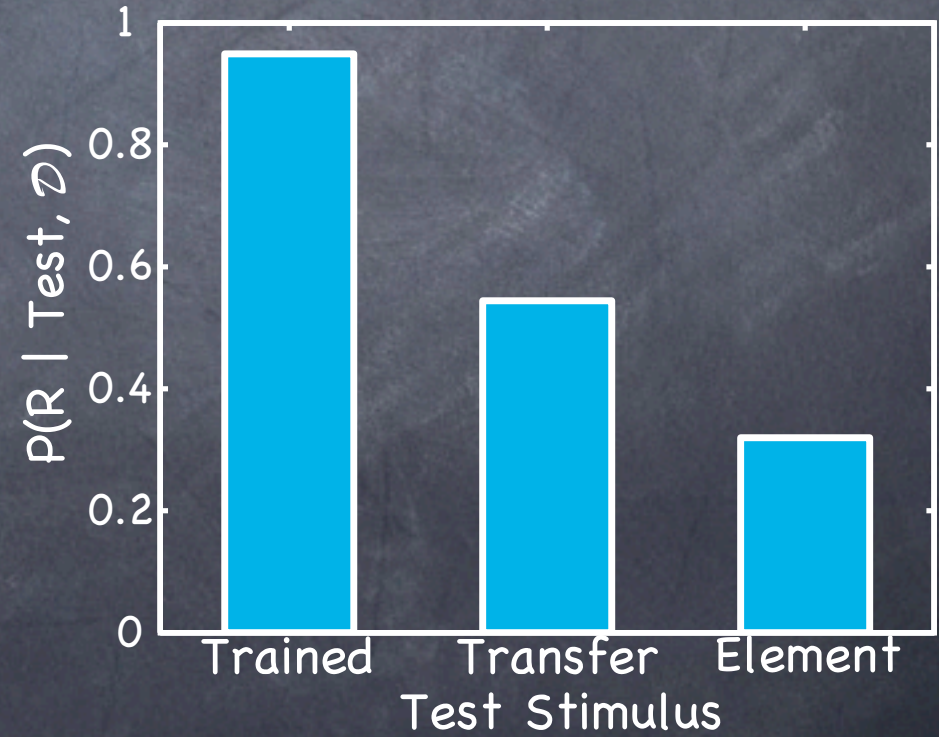
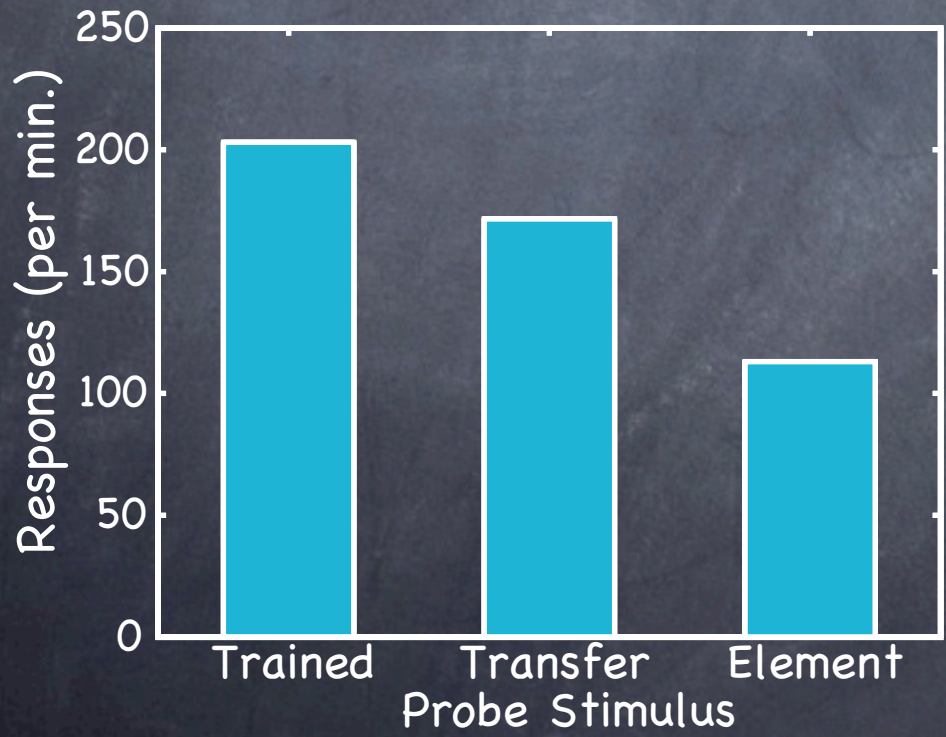
$$P(R | Stim, \mathcal{D}) = \sum_m P(R | Stim, m, \mathcal{D}) P(m | \mathcal{D})$$

L.V. Model of Paired Compounds

MAP Model Structure:

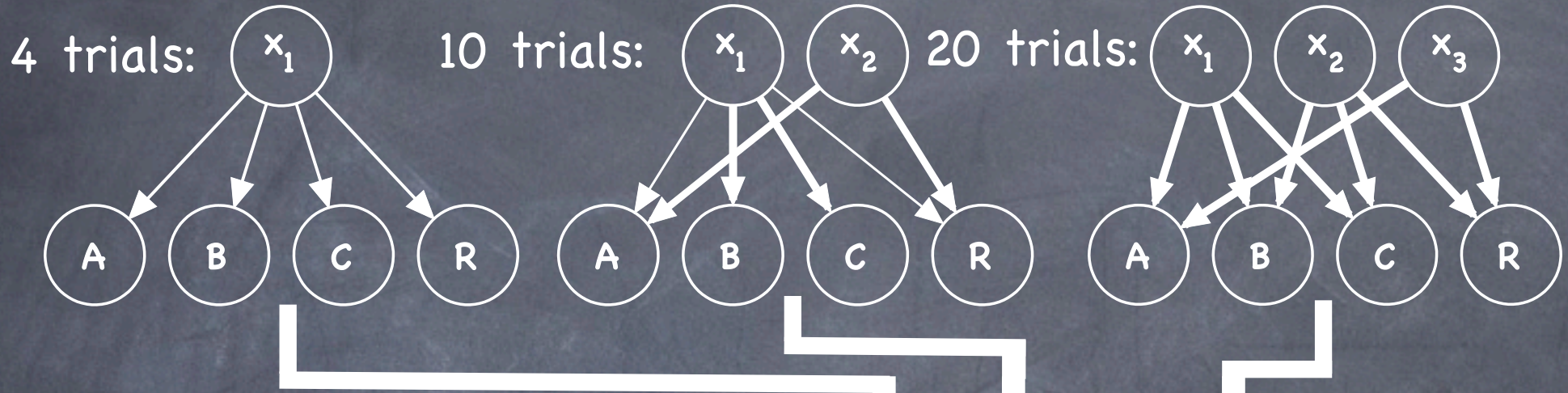


Training: AB+ CD+

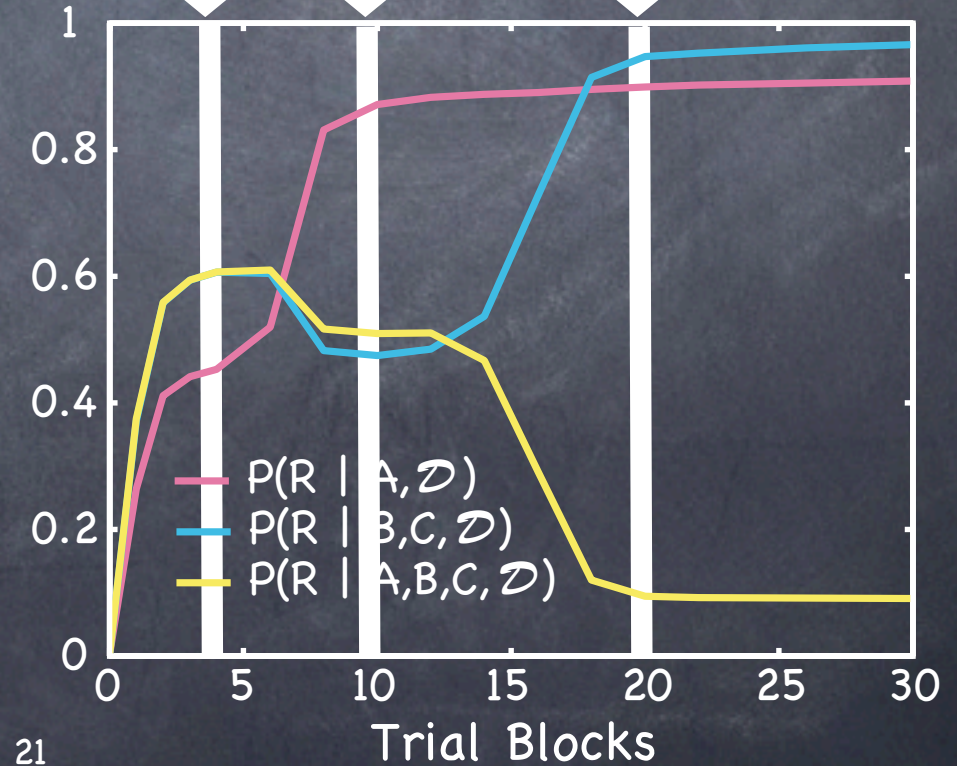
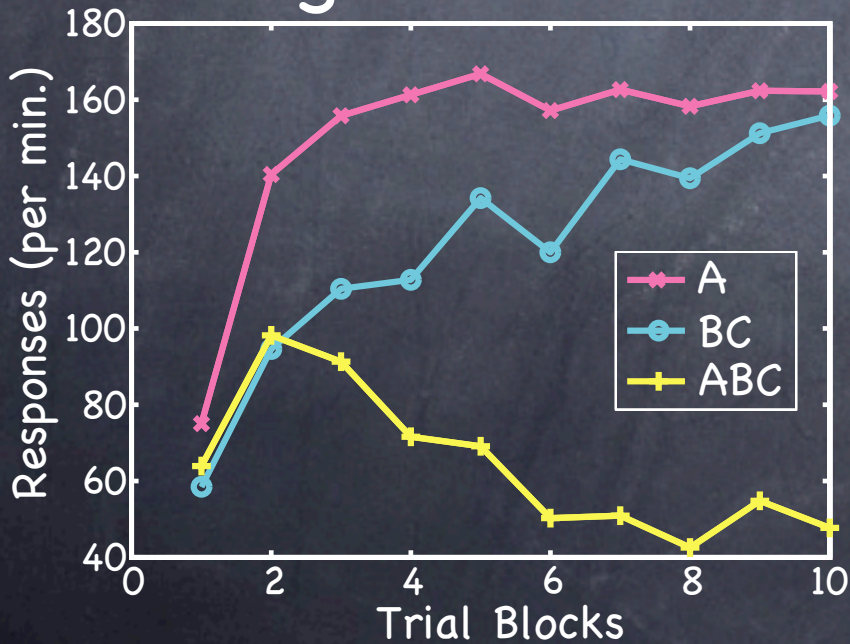


L.V. Model of Asymmetric XOR

MAP Model Structures:



Training: A+ BC+ ABC-



What's a configuration?

- Can account for experiments that are traditionally deemed “**Configural Conditioning**”
- Previous models cast configuration as result of stimuli being trained together.
- We view it as the result of model complexity pressures to group stimuli.

Expt: Second-order conditioning versus Conditioned Inhibition

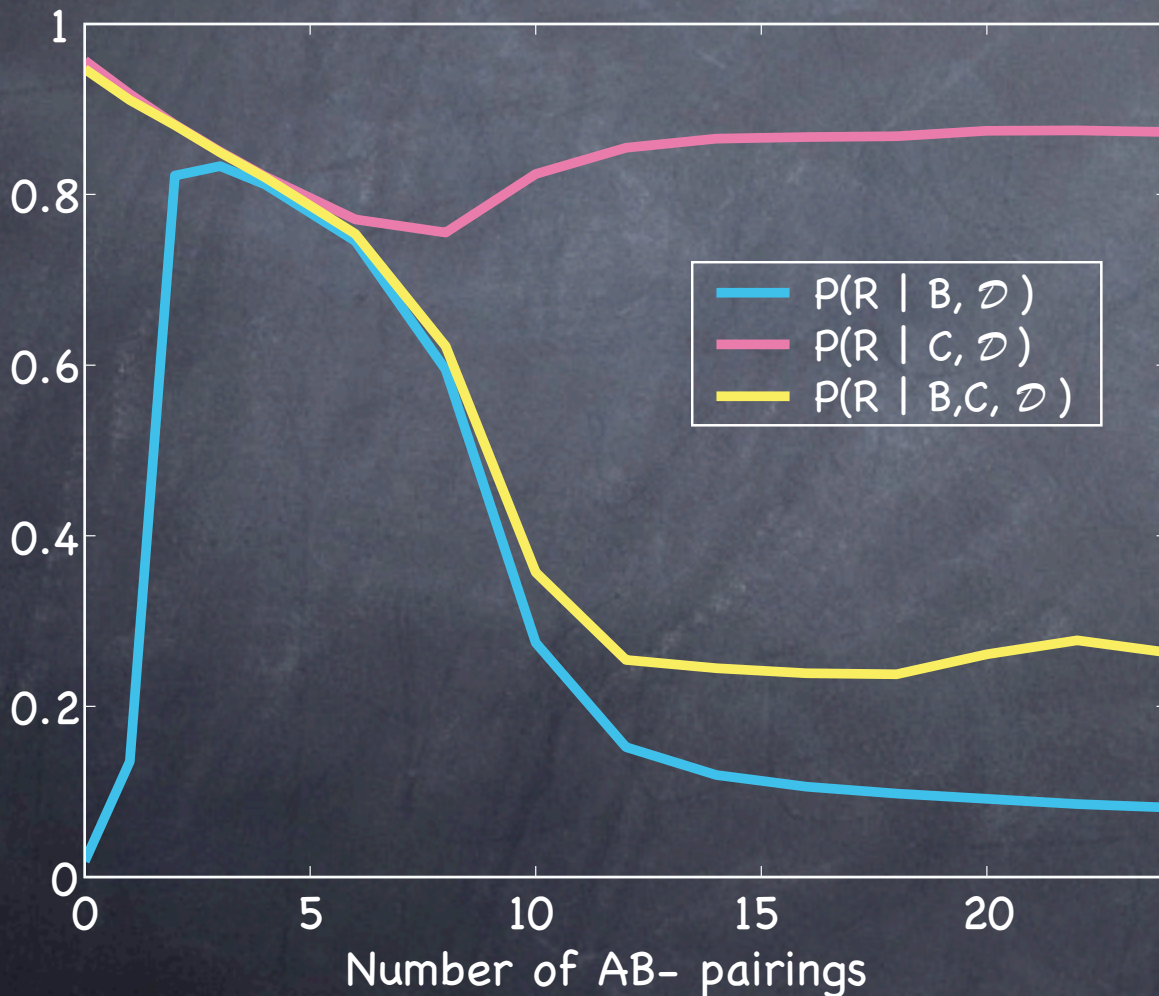
[Yin et al, 1994]

| Group | A+ | AB- | C+ | Test → Result | Test → Result |
|--------|----|-----|----|---------------|---------------|
| No B | 96 | 0 | 8 | B → _ | BC → Resp. |
| Few B | 96 | 4 | 8 | B → Resp. | BC → Resp. |
| Many B | 96 | 48 | 8 | B → _ | BC → _ |

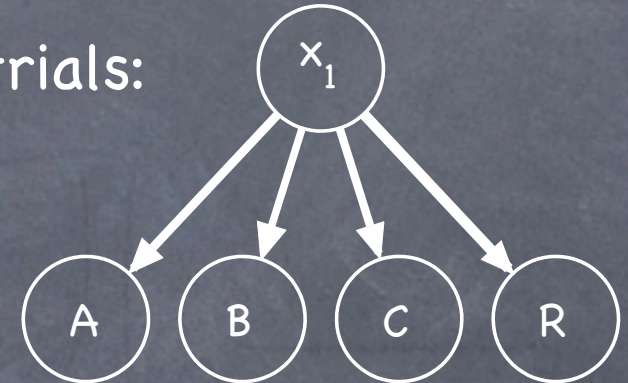
Bayesian Model of Second-Order Conditioning / Conditioned Inhibition

MAP Model Structure:

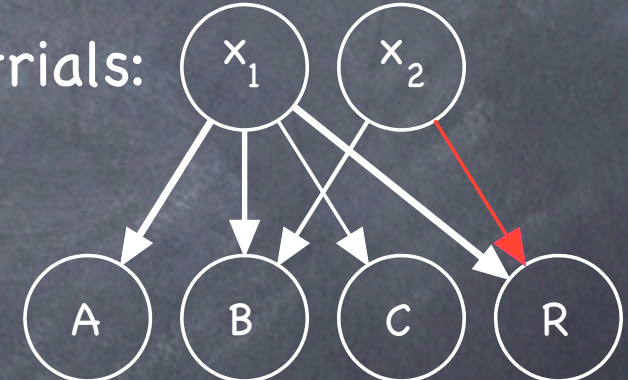
Training: A+ AB- C+



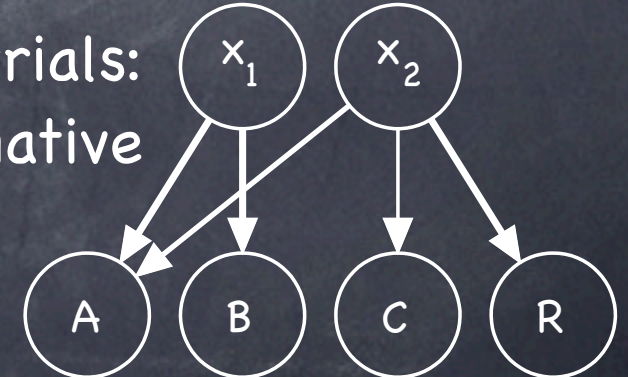
4 trials:



18 trials:



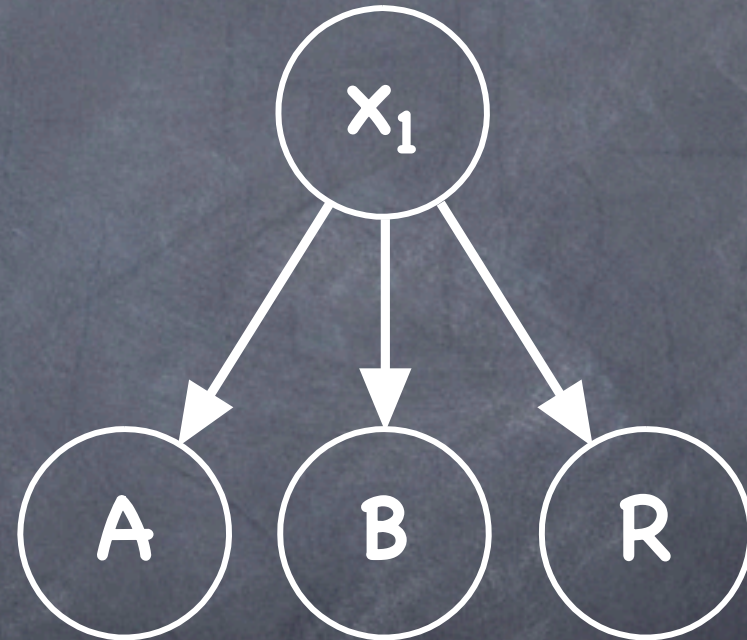
18 trials:
Alternative



Dealing with Reinforcement

Are reinforcers really just like other stimulus?

Train: $A+$ $B+$

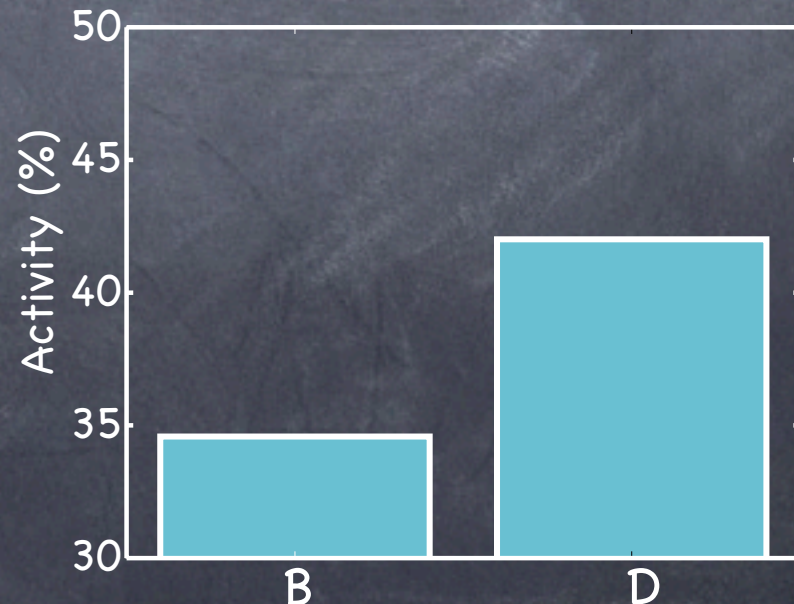
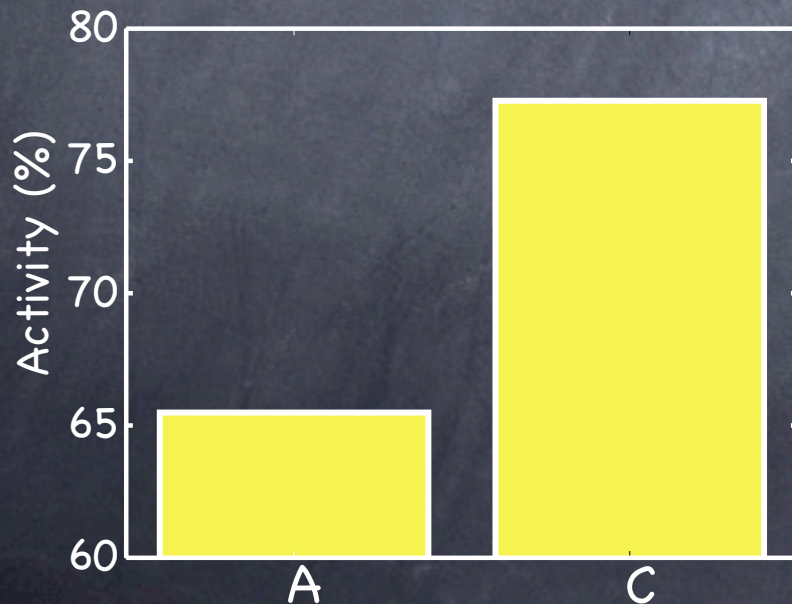


Do animals do this?

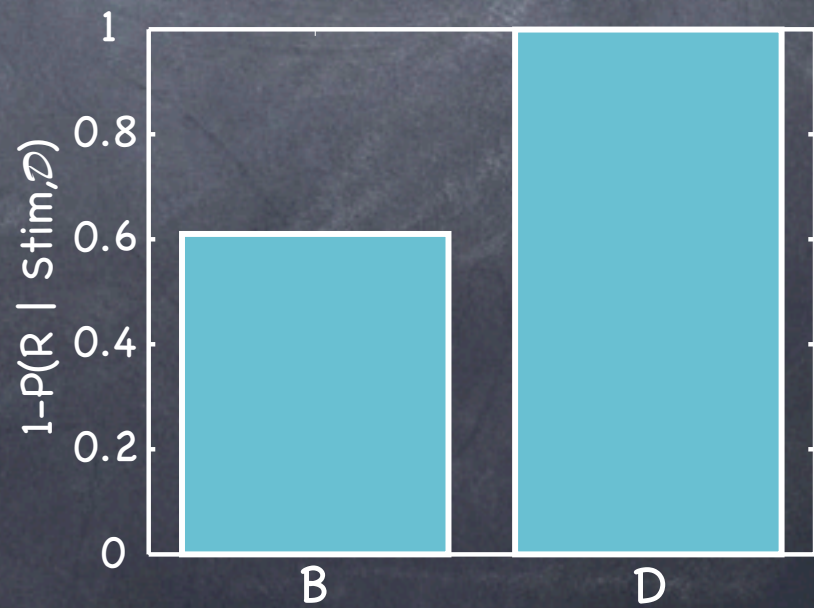
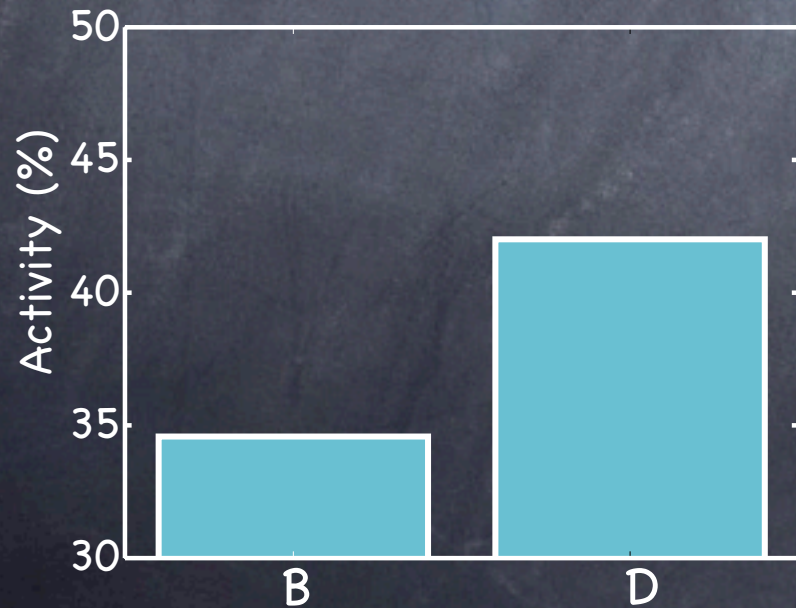
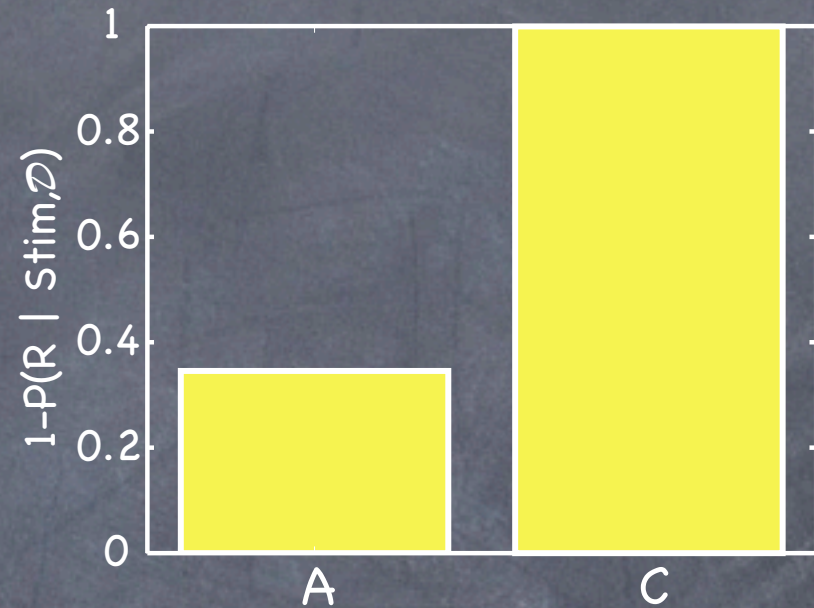
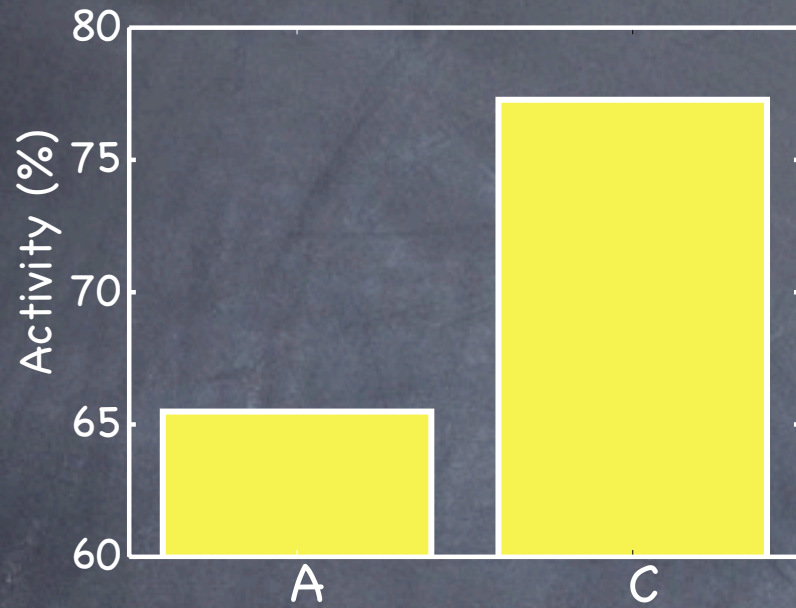
Acquired Relational Equivalence

[Honey & Watt, 1999]

| Biconditional training | | Revaluation | Test |
|------------------------|------------|-------------|--------|
| AY-food | AZ-no food | A-shock | A vs C |
| BY-food | BZ-no food | | |
| CY-no food | CZ-food | C-no shock | B vs D |
| DY-no food | DZ-food | | |

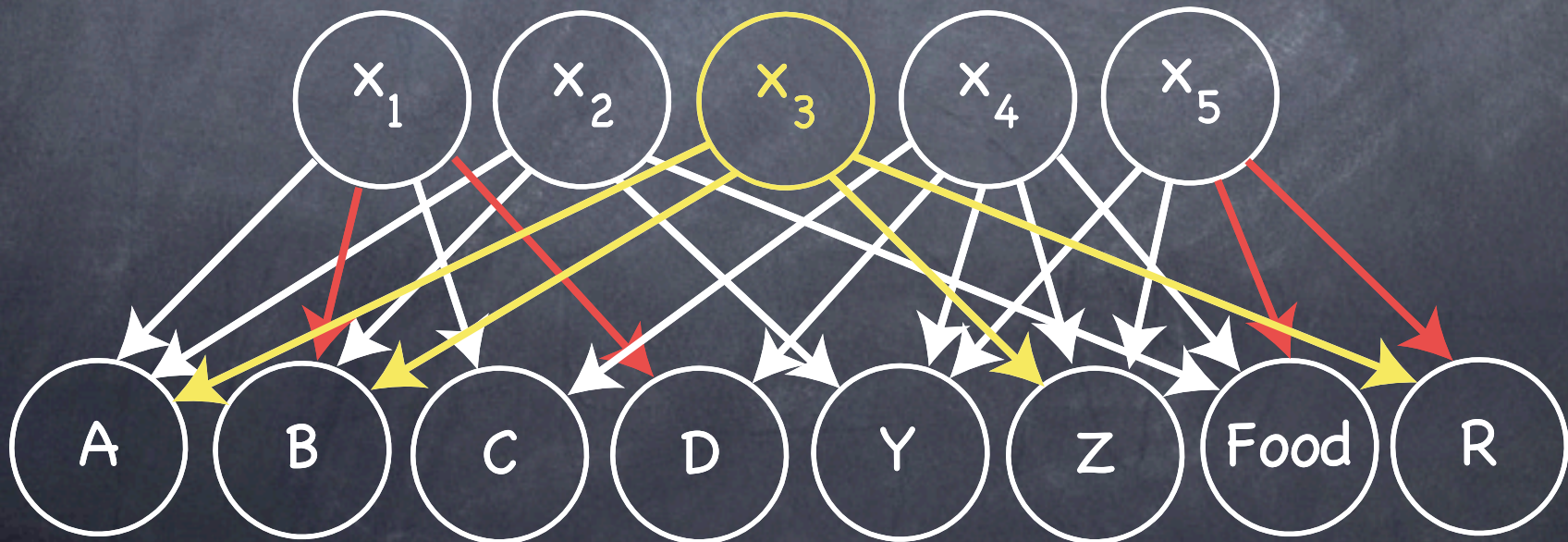


Modeling Acq. Rel. Equiv.



Modeling Acq. Rel. Equiv.

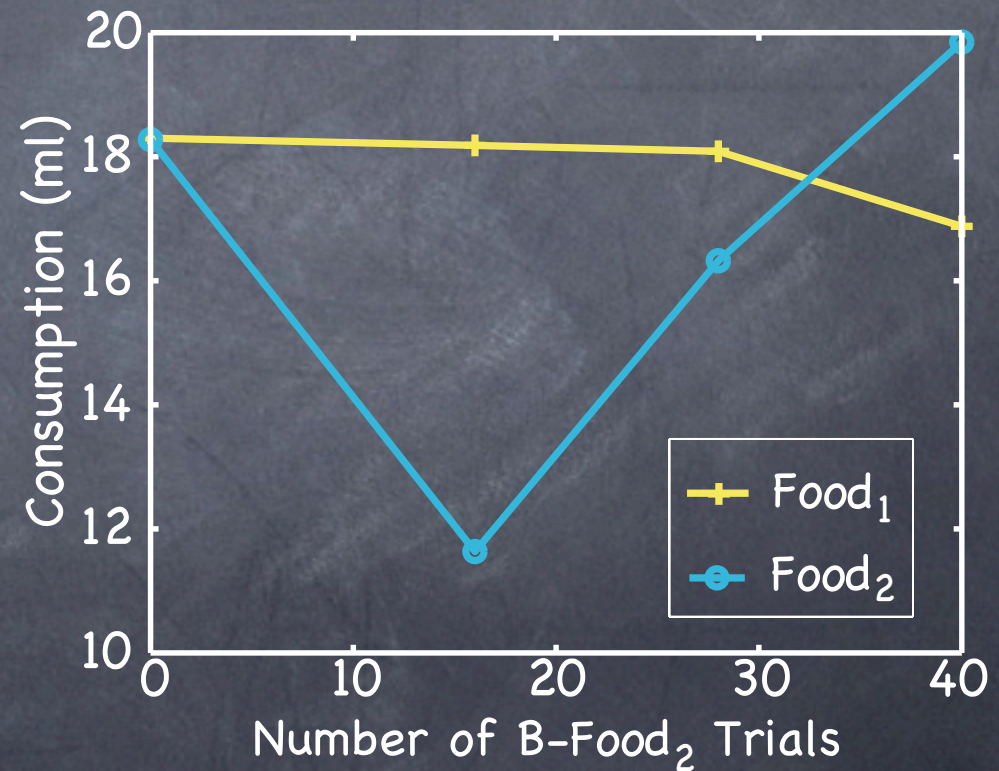
| Biconditional training | | Revaluation | Test |
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| AY-food | AZ-no food | A-shock | A vs C |
| BY-food | BZ-no food | | |
| CY-no food | CZ-food | | |
| DY-no food | DZ-food | C-no shock | B vs D |



Expt: Food Devaluation

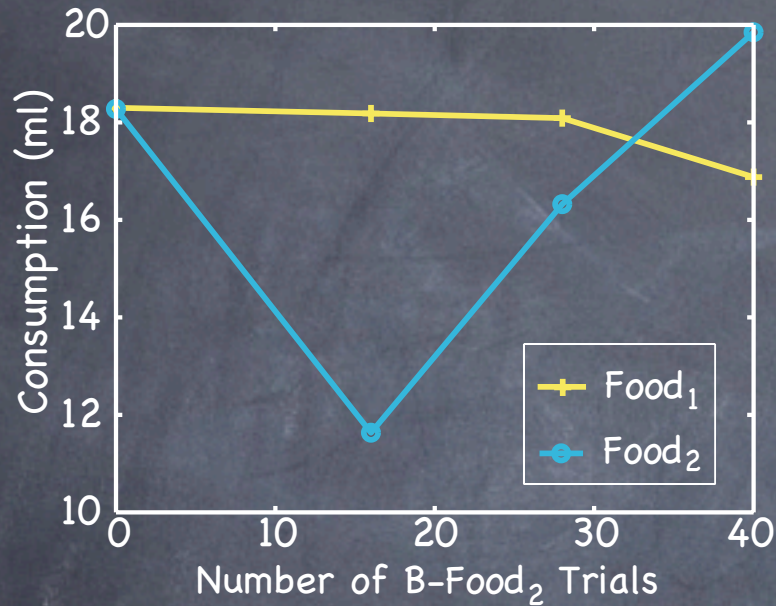
[Holland, 1998]

| Training Trials | | | Test |
|-----------------|------|---------|----------|
| Phase 1 | | Phase 2 | |
| A-F1 | B-F2 | B-R | |
| 16 | 0 | 6 | F1 F2 |
| | 16 | | |
| | 28 | | |
| | 40 | | |

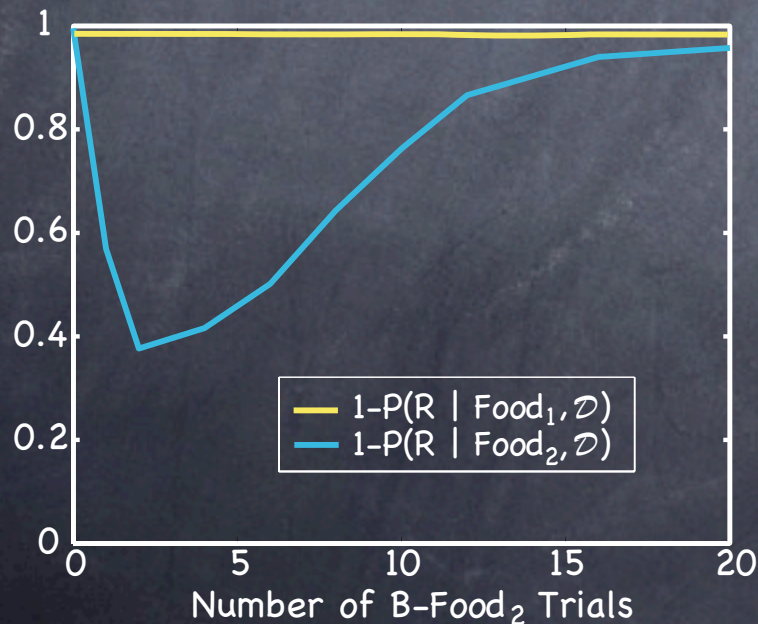
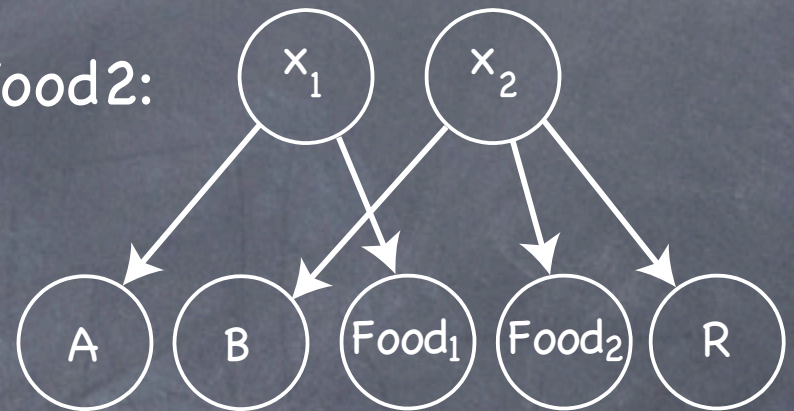


L.V. model of Devaluation

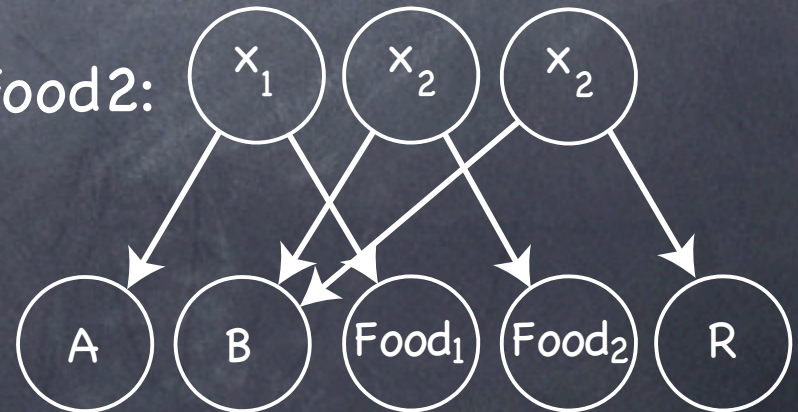
A-Food1 / B-Food2, B-R



Few B-Food2:



Many B-Food2:

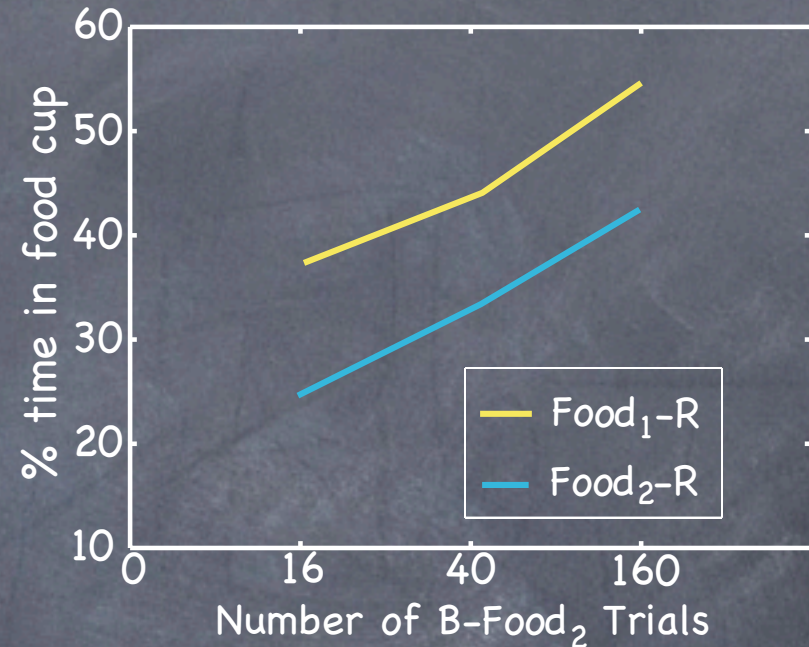


Not the whole story...

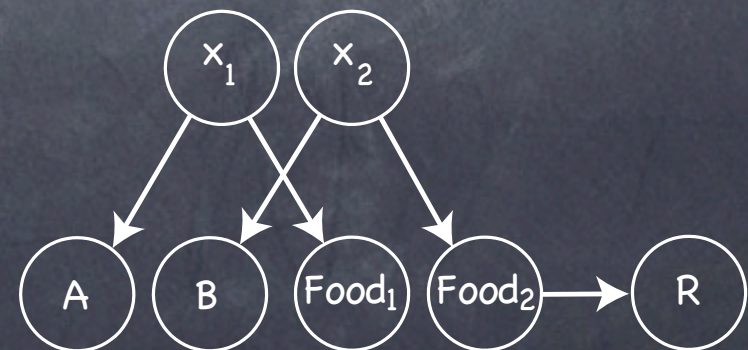
(Variant on Devaluation)

[Holland, 1998]

| Training Trials | | | Test |
|-----------------|------|----------|------|
| Phase 1 | | Phase 2 | |
| A-F1 | B-F2 | F(1,2)-R | |
| 16 | 16 | 2 | B |
| | 40 | | |
| | 160 | | |



Model Structure?



Future Directions

- Explore the priors: Experimentally manipulatable.
- Remove independent trial assumption.

Modeling Change

- Should reflect our understanding of how the world is believed to change.
- Example: Causal model **parameter drift**.
- The marginal distribution of the diffusion process should reflect your prior

Conclusions

- **Similarity** and **Discrimination** are recognized as the tradeoff between **complexity** and **data fidelity** arising in Bayesian inference.
- A latent variable is a natural (causal) setting for the study of classical conditioning.
- Account for configural conditioning data and more.