**An active inference approach to modeling structure learning: concept learning**

**as an example case**

Ryan Smith,[1] Philipp Schwartenbeck,[2] Thomas Parr,[2] Karl J. Friston,[2]

[1]Laureate Institute for Brain Research, Tulsa, OK, USA

[2]Wellcome Centre for Human Neuroimaging, Institute of Neurology, University

College London, WC1N 3BG, UK

12    Corresponding Author Information:

14    Ryan Smith

15    Laureate Institute for Brain Research

16    6655 S Yale Ave, Tulsa, OK 74136, USA

17    Email: rsmith@laureateinstitute.org

18                                   **Abstract**

Within computational neuroscience, the algorithmic and neural basis of structure

learning remains poorly understood. Concept learning is one primary example,

which requires both a type of internal model expansion process (adding novel

hidden states that explain new observations), and a model reduction process

(merging different states into one underlying cause and thus reducing model

complexity via meta-learning). Although various algorithmic models of concept

learning have been proposed within machine learning and cognitive science, many

are limited to various degrees by an inability to generalize, the need for very large

amounts of training data, and/or insufficiently established biological plausibility.

Using concept learning as an example case, we introduce a novel approach for

modeling structure learning within the active inference framework and its

accompanying neural process theory. This approach is based on the idea that a

generative model can be equipped with extra (hidden state or cause) 'slots' that can

be engaged when an agent learns about novel concepts. This can be combined with a

Bayesian model reduction process, in which any concept learning – associated with

these slots – can be reset in favor of a simpler model with higher model evidence.

We use simulations to illustrate this model's ability to add new concepts to its state

space (with relatively few observations) and increase the granularity of the

concepts it currently possesses. We also simulate the predicted neural basis of these

processes. We further show that it accomplishes a simple form of 'one-shot'

generalization to new stimuli. Although deliberately simple, these results suggest

that this general approach to modeling concept learning within active inference

research may also offer useful resources in developing neurocomputational models

of structure learning more generally.

*Keywords*: Model Expansion; Structure Learning; Concepts; Computational

Neuroscience; Active Inference

46                                              **Introduction**

47              The ability to learn the latent structure of one's environment – such as

inferring the existence of hidden causes of regularly observed patterns in co-

occurring feature observations – is central to human cognition. For example, we do

not simply observe particular sets of colors, textures, shapes, and sizes – we also

observe *identifiable object*s such as, say, a 'screwdriver'. If we were tool experts, we

might also recognize particular types of screwdrivers (e.g., flat vs. Phillip's head),

designed for a particular use. This ability to learn latent structure, such as learning

to recognize co-occurring features under conceptual categories (as opposed to just

perceiving sensory qualities; e.g., red, round, etc.), is also highly adaptive. Only if we

knew an object was a screwdriver could we efficiently infer that it affords putting

certain structures together and taking them apart; and only if we knew the specific

type of screwdriver could we efficiently infer, say, the artefacts to use it on. Many

concepts of this sort require experience-dependent acquisition (i.e., learning).

60              From a computational perspective, the ability to acquire a new concept can

be seen as a type of structure learning involving Bayesian model comparison

(Botvinick, Niv, & Barto, 2009; S. J. Gershman & Niv, 2010; MacKay & Peto, 1995;

Salakhutdinov, Tenenbaum, & Torralba, 2013; Tervo, Tenenbaum, & Gershman,

2016). Specifically, concept acquisition can be cast as an agent learning (or

inferring) that a new hypothesis (referred to here as a hidden cause or state) should

be added to the internal or generative model with which she explains her

environment, because existing causes cannot account for new observations (e.g., an

agent might start out believing that the only tools are hammers and screwdrivers,

but later learn that there are also wrenches). In other words, the structure of the

space of hidden causes itself needs to expand to accommodate new patterns of

observations. This model expansion process is complementary to a process called

Bayesian model reduction (Karl Friston & Penny, 2011); in which the agent can infer

that there is redundancy in her model, and a model with fewer states or parameters

provides a more parsimonious (i.e. simpler) explanation of observations (KJ Friston,

Lin, et al., 2017; Schmidhuber, 2006). For example, in some instances it may be

more appropriate to differentiate between fish and birds as opposed to salmon,

peacocks and pigeons. This reflects a reduction in model complexity based on a

particular feature space underlying observations and thus resonates with other

accounts of concept learning as dimensionality reduction (Behrens et al., 2018;

Stachenfeld, Botvinick, & Gershman, 2016) – a topic we discuss further below.

A growing body of work in a number of domains has approached this

problem from different angles. In developmental psychology and cognitive science,

for example, probability theoretic (Bayesian) models have been proposed to account

for word learning in children and the remarkable human ability to generalize from

very few (or even one) examples in which both broader and narrower categorical

referents could be inferred (Kemp, Perfors, & Tenenbaum, 2007; Lake,

Salakhutdinov, & Tenenbaum, 2015; Perfors, Tenenbaum, Griffiths, & Xu, 2011; Xu &

Tenenbaum, 2007a, 2007b). In statistics, a number of nonparametric Bayesian

models, such as the "Chinese Room" process and the "Indian Buffet" process, have

been used to infer the need for model expansion (S. Gershman & Blei, 2012). There

are also related approaches in machine learning, as applied to things like Gaussian

mixture models (McNicholas, 2016).

Such approaches employ various clustering algorithms, which take sets of

data points in a multidimensional space and divide them into separable clusters

(e.g., see (Anderson, 1991; Love, Medin, & Gureckis, 2004; Sanborn, Griffiths, &

Navarro, 2010)). While many of these approaches assume the number of clusters is

known in advance, various goodness-of-fit criteria may be used to determine the

optimal number. However, a number of approaches require much larger amounts of

data than humans do to learn new concepts (Geman, Bienenstock, & Doursat, 1992;

Hinton et al., 2012; LeCun, Bengio, & Hinton, 2015; Lecun, Bottou, Bengio, & Haffner,

1998; Mnih et al., 2015). Many also require corrective feedback to learn and yet fail

to acquire sufficiently rich conceptual structure to allow for generalization

(Barsalou, 1983; Biederman, 1987; Feldman, 1997; Jern & Kemp, 2013; A. B.

Markman & Makin, 1998; Osherson & Smith, 1981; Ward, 1994; Williams &

Lombrozo, 2010).

Approaches to formally modeling structure learning, including concept

learning, have not yet been examined within the emerging field of research on

Active Inference models within computational neuroscience (KJ Friston, 2010; KJ

Friston et al., 2016; KJ Friston, Lin, et al., 2017; KJ Friston, Parr, & de Vries, 2017).

This represents one potentially fruitful research avenue that has not yet been

examined and, as discussed below, may offer unique advantages in research focused

on understanding the neural basis of learning latent structure. In this paper, we

explore the potential of this approach. In brief, we conclude that structure learning

is an emergent property of active inference (and learning) under generative models

with 'spare capacity'; where spare or uncommitted capacity is used to expand the

repertoire of representations (Baker & Tenenbaum, 2014), while Bayesian model

reduction (KJ Friston, Lin, et al., 2017; Hobson & Friston, 2012) promotes

generalization by minimizing model complexity – and releasing representations to

replenish 'spare capacity'.

From a machine learning perspective, Bayesian model reduction affords the

opportunity to consider generative models with a large number of hidden states or

latent factors and then optimize the number (or indeed partitions) of latent factors

with respect to a variational bound on model evidence. This could be regarded as a

bounded form of nonparametric Bayes, in which a potentially infinite number of

latent factors are considered; with appropriate (e.g., Indian buffet process) priors

over the number of hidden states generating data features[1]. The Bayesian model

reduction approach here is bounded in the sense that an upper bound on the

number of hidden states is specified prior to structure learning. Furthermore, in

virtue of the (biologically plausible) variational schemes used for model reduction,

there is no need to explicitly compute model evidence; thereby emulating the

computational efficiency of nonparametric Bayes (S. Gershman & Blei, 2012), while

accommodating any prior over models.

In what follows, we first provide a brief overview of active inference. We then

introduce a model of concept learning (using basic and subordinate level animal

---

[1] Generally motivated by starting with a finite parametric model and taking the limit as the number of latent states with more parameters tends to infinity.

categories), as a representative example of structure learning. We specifically model

cognitive (semantic) processes that add new concepts to a state space and that

optimize the granularity of an existing state space. We then establish the validity of

this model using numerical analyses of concept learning, when repeatedly

presenting a synthetic agent with different animals characterized by different

combinations of observable features. We demonstrate how particular approaches

combining Bayesian model reduction and expansion can reproduce successful

concept learning without the need for corrective feedback – and allow for

generalization. We further demonstrate the ability of this model to generate

predictions about measurable neural responses based on the neural process theory

that accompanies active inference. We conclude with a brief discussion of the

implications of this work. Our goal is to present an introductory proof of concept –

that could be used as the foundation of future research on the neurocomputational

basis of structure learning.

150                    **An Active Inference model of concept learning**

**A primer on Active Inference**

Active Inference suggests that the brain is an inference machine that

approximates optimal probabilistic (Bayesian) belief updating across perceptual,

cognitive, and motor domains. Active Inference more specifically postulates that the

brain embodies an internal model of the world that is "generative" in the sense that

it can simulate the sensory data that it should receive if its model of the world is

correct. These simulated (predicted) sensory data can be compared to actual

observations, and differences between predicted and observed sensations can be

used to update the model. Over short timescales (e.g., a single observation) this

updating corresponds to inference (perception), whereas on longer timescales it

corresponds to learning (i.e., updating expectations about what will be observed

later). Another way of putting this is that perception optimizes beliefs about the

current state of the world, while learning optimizes beliefs about the relationships

between the variables that constitute the world. These processes can be seen as

ensuring the generative model (entailed by recognition processes in the brain)

remains an accurate model of the world that it seeks to regulate (Conant & Ashbey,

1970).

Active Inference casts decision-making in similar terms. Actions can be

chosen to resolve uncertainty about variables within a generative model (i.e.,

sampling from domains in which the model does not make precise predictions),

which can prevent anticipated deviations from predicted outcomes. In addition,

some expectations are treated as a fixed phenotype of an organism. For example, if

an organism did not continue to "expect" to observe certain amounts of food, water,

and shelter, then it would quickly cease to exist (McKay & Dennett, 2009) – as it

would not pursue those behaviors that fulfill these expectations (c.f. the 'optimism

bias' (Sharot, 2011)). Thus, a creature should continually seek out observations that

support – or are internally consistent with – its own continued existence. Decision-

making can therefore be cast as a process in which the brain infers the sets of

actions (policies) that would lead to observations consistent with its own survival-

related expectations (i.e., its "prior preferences"). Mathematically, this can be

described as selecting sequences of actions (policies) that maximize "Bayesian

model evidence" expected under a policy, where model evidence is the (marginal)

likelihood that particular sensory inputs would be observed under a given model.

In real-world settings, directly computing Bayesian model evidence is

generally intractable. Thus, some approximation is necessary. Active Inference

proposes that the brain computes a quantity called "variational free energy" that

provides a bound on model evidence, such that minimization of free energy

indirectly maximizes model evidence (this is exactly the same functional used in

machine learning where it is known as an evidence lower bound or ELBO). In this

case, decision-making will be approximately (Bayes) optimal if it infers (and enacts)

the policy that will minimize expected free energy (i.e., free energy with respect to a

policy, where one takes expected future observations into account). Technically,

expected free energy is the average free energy under the posterior predictive

density over policy-specific outcomes.

Expected free energy can be decomposed in different ways that reflect

uncertainty and prior preferences, respectively (e.g., epistemic and instrumental

affordance or ambiguity and risk). This formulation means that any agent that

minimizes expected free energy engages initially in exploratory behavior to

minimize uncertainty in a new environment. Once uncertainty is resolved, the agent

then exploits that environment to fulfil its prior preferences. The formal basis for

Active Inference has been thoroughly detailed elsewhere (KJ Friston, FitzGerald,

Rigoli, Schwartenbeck, & Pezzulo, 2017), and the reader is referred there for a full

mathematical treatment.

When the generative model is formulated as a partially observable Markov

decision process (a mathematical framework for modeling decision-making in cases

where some outcomes are under the control of the agent and others are not, and

where states of the world are not directly known but must be inferred from

observations), active inference takes a particular form. Here, the generative model is

specified by writing down plausible or allowable policies, hidden states of the world

(that must be inferred from observations), and observable outcomes, as well as a

number of matrices that define the probabilistic relationships between these

quantities (see right panel of figure1).

Figure 1. Left: Illustration of the trial structure performed by the agent. At the first time
point, the agent is exposed to one of 8 possible animals that are each characterized by a
unique combination of visual features. At the 2nd time point, the agent would then report
which animal concept matched that feature combination. The agent could report a specific
category (e.g., pigeon, hawk, minnow, etc.) or a general category (i.e., bird or fish) if
insufficiently certain about the specific category. See the main text for more details. Right:
Illustration of the Markov decision process formulation of active inference used in the
simulations described in this paper. The generative model is here depicted graphically, such
that arrows indicate dependencies between variables. Here observations (**o**) depend on
hidden states (**s**), as specified by the **A** matrix, and those states depend on both previous
states (as specified by the **B** matrix,  or the initial states specified by the **D** matrix) and the
policies (**π**) selected by the agent. The probability of selecting a particular policy in turn
depends on the expected free energy (**G**) of each policy with respect to the prior preferences
(**C**) of the agent. The degree to which expected free energy influences policy selection is also
modulated by a prior policy precision parameter (**γ**), which is in turn dependent on beta (*β*)
–where higher values of beta promote more randomness in policy selection (i.e., less
influence of the differences in expected free energy across policies). For more details
regarding the associated mathematics, see (KJ Friston, Lin, et al., 2017; KJ Friston, Parr, et
al., 2017).
235

The 'A' matrix indicates which observations are generated by each

combination of hidden states (i.e., the likelihood mapping specifying the probability

that a particular set of observations would be observed given a particular set of

hidden states). The 'B' matrix is a transition matrix, indicating the probability that

one hidden state will evolve into another over time. The agent, based on the selected

policy, controls some of these transitions (e.g., those that pertain to the positions of

its limbs). The 'D' matrix encodes prior expectations about the initial hidden state

the agent will occupy. Finally, the 'C' matrix specifies prior preferences over

observations; it quantifies the degree to which different observed outcomes are

rewarding or punishing to the agent. In these models, observations and hidden

states can be factorized into multiple outcome *modalities* and hidden state *factors*.

This means that the likelihood mapping (the 'A' matrix) can also model the

interactions among different hidden states when generating outcomes

(observations).

**From principles to process theories**

One potential empirical advantage of the present approach stems from the

fact that active inference models have a plausible biological basis that affords

testable neurobiological predictions. Specifically, these models have well-articulated

companion micro-anatomical neural process theories, based on commonly used

message-passing algorithms (KJ Friston, FitzGerald, et al., 2017; Parr & Friston,

2018; Parr, Markovic, Kiebel, & Friston, 2019). In these process theories, for

example, the activation level of different neural populations (typically portrayed as

consisting of different cortical columns) can encode posterior probability estimates

over different hidden states. These activation levels can then be updated by synaptic

inputs with particular weights that convey the conditional probabilities encoded in

the 'A' and 'B' (among other) matrices described above, where active learning then

corresponds to associative synaptic plasticity. Phasic dopamine responses also play

a particular role in these models, by reporting changes in policy precision (i.e., the

degree of confidence in one policy over others) upon new observations (see Figure 2

and the associated legend for more details). In what follows, we describe how the

type of generative model – that underwrites these processes – was specified to

perform concept inference/learning.

Figure 2. This figure illustrates the mathematical framework of active inference and associated neural process theory used in the simulations described in this paper. The differential equations in the left panel approximate Bayesian belief updating within the graphical model depicted in the right panel of Figure 1 via a gradient descent on free energy (**F**). The right panel also illustrates the proposed neural basis by which neurons making up cortical columns could implement these equations. The equations have been expressed in terms of two types of prediction errors. State prediction errors ($\boldsymbol{\varepsilon}$) signal the difference between the (logarithms of) expected states (**s**) under each policy and time point—and the corresponding predictions based upon outcomes/observations (**A** matrix) and the (preceding and subsequent) hidden states (**B** matrix, and, although not written, the **D** matrix for the initial hidden states at the first time point). These represent prior and likelihood terms respectively – also marked as messages 2, 3, and 4, which are depicted as being passed between neural populations (colored balls) via particular synaptic connections in the right panel. These (prediction error) signals drive depolarization (**v**) in those neurons encoding hidden states (**s**), where the probability distribution over hidden states is then obtained via a softmax (normalized exponential) function ($\boldsymbol{\sigma}$). Outcome prediction errors ($\boldsymbol{\varsigma}$) instead signal the difference between the (logarithms of) expected observations (**o**) and those predicted under prior preferences (**C**). This term additionally considers the expected ambiguity or conditional entropy (**H**) between states and outcomes as well as a novelty term (**W**) reflecting the degree to which beliefs about how states generate outcomes would change upon observing different possible state-outcome mappings (computed from the **A** matrix). This prediction error is weighted by the expected observations to evaluate the expected free energy (**G**) for each policy ($\boldsymbol{\pi}$), conveyed via message 5. These policy-specific free energies are then integrated to give the policy expectations via a softmax function, conveyed through message 1. Actions at each time

point (**u**) are then chosen out of the possible actions under each policy (**U**) weighted by the
value (negative expected free energy) of each policy. In our simulations, the model learned
associations between hidden states and observations (**A**) via a process in which counts
were accumulated (**a**) reflecting the number of times the agent observed a particular
outcome when she believed that she occupied each possible hidden state. Although not
displayed explicitly, learning prior expectations over initial hidden states (**D**) is similarly
accomplished via accumulation of concentration parameters (**d**). These prior expectations
reflect counts of how many times the agent believes it previously occupied each possible
initial state. Concentration parameters are converted into expected log probabilities using
digamma functions ($\psi$). The way in which Bayesian model reduction was performed in this
paper is also written in the lower left (where B indicates a beta function, and **m** is the
posterior probability of each model). Here, the posterior distribution over initial states (**d**)
is used to assess the difference in the evidence ($\Delta$**F**) it provides for the number of hidden
states in the current model and other possible models characterized by fewer hidden states.
Prior concentration parameters are shown in italics, posterior in bold, and those priors and
posteriors associated with the reduced model are equipped with a tilde (**~**). As already
stated, the right panel illustrates a possible neural implementation of the update equations
in the left panel. In this implementation, probability estimates have been associated with
neuronal populations that are arranged to reproduce known intrinsic (within cortical area)
connections. Red connections are excitatory, blue connections are inhibitory, and green
connections are modulatory (i.e., involve a multiplication or weighting). These connections
mediate the message passing associated with the equations in the left panel. Cyan units
correspond to expectations about hidden states and (future) outcomes under each policy,
while red states indicate their Bayesian model averages (i.e., a "best guess" based on the
average of the probability estimates for the states and outcomes across policies, weighted
by the probability estimates for their associated policies. Pink units correspond to (state
and outcome) prediction errors that are averaged to evaluate expected free energy and
subsequent policy expectations (in the lower part of the network). This (neural) network
formulation of belief updating means that connection strengths correspond to the
parameters of the generative model described in the text. Learning then corresponds to
changes in the synaptic connection strengths. Only exemplar connections are shown to
avoid visual clutter. Furthermore, we have just shown neuronal populations encoding
hidden states under two policies over three time points (i.e., two transitions), whereas in
the task described in this paper there are greater number of allowable policies. For more
information regarding the mathematics and processes illustrated in this figure, see (KJ
Friston, Lin, et al., 2017; KJ Friston, Parr, et al., 2017).
334

**A model of concept inference and learning**

To model concept inference, we constructed a simple task for an agent to

perform (see figure 1, left panel). In this task, the agent was presented with different

animals on different trials and asked to answer a question about the type of animal

that was seen. As described below, in some simulations the agent was asked to

report the type of animal that was learned previously; in other simulations, the

agent was instead asked a question that required conceptual generalization.

Crucially, to answer these questions the agent was required to observe different

animal features, where the identity of the animal depended on the combination of

features. There were three feature categories (size, color, and species-specific;

described further below) and two discrete time points in a trial (observe and

report).

To simulate concept learning (based on the task described above) we needed

to specify an appropriate generative model. Once this model has been specified, one

can use standard (variational) message passing to simulate belief updating and

behavior in a biologically plausible way: for details, please see (KJ Friston,

FitzGerald, et al., 2017; KJ Friston, Parr, et al., 2017). In our (minimal) model, the

first hidden state factor included (up to) eight levels, specifying four possible types

of birds and four possible types of fish (Figure 3A). The outcome modalities

included: a feature space including two size features (big, small), two color features

(gray, colorful), and two species-differentiating features (wings, gills). The 'A' matrix

specified a likelihood mapping between features and animal concepts, such that

each feature combination was predicted by an animal concept (Hawk, Pigeon,

Parrot, Parakeet, Sturgeon, Minnow, Whale shark, Clownfish). This model was

deliberately simple to allow for a clear illustration, but it is plausibly scalable to

include more concepts and a much larger feature space. The 'B' matrix for the first

hidden state factor was an identity matrix, reflecting the belief that the animal

identity was conserved during each trial (i.e., the animals were not switched out

mid-trial).

**Generative model**

B *Concepts*

A *Concepts*

**Hidden State Factor 1: Concept Inference**

ANIMAL CONCEPTS

Parakeet  Parrot  Pigeon  Hawk  Clownfish  Whale Shark  Minnow  Sturgeon

Large  Small    Colorful  Gray    Wings  Gills

Size    Color    Species-specific

Observable Features

Outcome Factors 1, 2, 3

A

364
365

Figure 3. (A) Illustration of the first hidden state factor containing columns (levels) for 8 different animal concepts. Each of these 8 concepts generated a different pattern of visual feature observations associated with the outcome modalities of size, color, and species-specific features. The B matrix was an identity matrix, indicating that the animal being observed did not change within a trial (white = 1, black = 0). The A matrix illustrates the specific mapping from animal concepts to feature combinations. As depicted, each concept corresponded to a unique point in a 3-dimensional feature space. (B) illustration of the 2nd hidden state factor corresponding to the verbal reports the agent could choose in response to her observations. These generated feedback as to whether her verbal report was accurate with respect to a basic category report or a specific category report. As illustrated in the C matrix, the agent most preferred to be correct about specific categories, but least preferred being incorrect. Thus, reporting the basic categories was a safer choice if the agent was too uncertain about the specific identity of the animal.

The second hidden state factor was the agent's report. That this is assumed to factorise from the first hidden state factor means that there is no prior constraint that links the chosen report to the animal generating observations. The agent could report each of the eight possible specific animal categories, or opt for a less specific report of a bird or a fish. Only one category could be reported at any time. Thus, the

agent had to choose to report only bird vs. fish or to report a more specific category.

In other words, the agent could decide upon the appropriate level of coarse-graining

of her responses (figure 3B).

During learning trials, the policy space was restricted such that the agent

could not provide verbal reports or observe corrective feedback (i.e., all it could do

is "stay still" in its initial state and observe the feature patterns presented). This

allowed the agent to learn concepts in an unsupervised manner (i.e. without being

told what the true state was or whether it was correct or incorrect). After learning,

active reporting was enabled, and the 'C' matrix was set so that the agent preferred

to report correct beliefs. We defined the preferences of the agent such that she

preferred correctly reporting specific category knowledge and was averse to

incorrect reports. This ensured that she only reported the general category of bird

vs. fish, unless sufficiently certain about the more specific category.

In the simulations reported below, there were two time points in each trial of

categorisation or conceptual inference. At the first time point, the agent was

presented with the animals features, and always began in a state of having made no

report (the "start" state). The agent's task was simply to observe the features, infer

the animal identity, and then report it (i.e., in reporting trials). Over 32 simulations

(i.e., 4 trials per animal), we confirmed that, if the agent already started out with full

knowledge of the animal concepts (i.e., a fully precise 'A' matrix), it would report the

specific category correctly 100% of the time. Over an additional 32 simulations, we

also confirmed that, if the agent was only equipped with knowledge of the

distinction between wings and gills (i.e., by replacing the rows in the 'A' matrix

corresponding to the mappings from animals to size and color with flat

distributions), it would report the generic category correctly 100% of the time but

would not report the specific categories.[2] This was an expected and straightforward

consequence of the generative model – but provides a useful example of how agents

trade off preferences and different types of uncertainty.

415                 **Simulating concept learning and the acquisition of expertise**

417             Having confirmed that our model could successfully recognize animals if

418     equipped with the relevant concepts (i.e., likelihood mappings) – we turn now to

419     concept learning.

421     **Concept acquisition**

422             We first examined our model's ability to acquire concept knowledge in two

423     distinct ways. This included 1) the agent's ability to "expand" (i.e., fill in an unused

424     column within) its state space and add new concepts, and 2) the agent's ability to

425     increase the granularity of its conceptual state space and learn more specific

426     concepts, when it already possessed broader concepts.

428     *Adding Concepts*

---

[2] However, "risky" reporting behavior could be elicited by manipulating the strengths of the agent's preferences such that she placed a very high value on reporting specific categories correctly (i.e., relative to how much she disliked reporting incorrectly).

To assess whether our agent could expand her state space by acquiring a new

concept, we first set one column of the previously described model's 'A' matrix

(mapping an animal concept to its associated features) to be a uniform distribution[3];

creating an imprecise likelihood mapping for one concept – essentially, that concept

predicted all features with nearly equal probability. Here, we chose sturgeon (large,

gray, gills) as the concept for which the agent had no initial knowledge (see Figure

4A, right-most column of left-most 'pre-learning' matrix). We then generated 2000

observations based on the outcome statistics of a model with full knowledge of all

eight animals (the "generative process"), to test whether the model could learn the

correct likelihood mapping for sturgeon (note: this excessive number of

observations was used for consistency with later simulations, in which more

concepts had to be learned, and also to evaluate how performance improved as a

function of the number of observations the agent was exposed to; see figure 4B).

In these simulations, learning was implemented via updating (concentration)

parameters for the model's 'A' matrix after each trial. For details of these free energy

minimizing learning processes, please see (KJ Friston et al., 2016) as well as the left

panel of Figure 2 and associated legend. An intuitive way to think about this belief

updating process is that the strength of association between a concept and an

observation is quantified simply by counting how often they are inferred to co-

occur. This is exactly the same principle that underwrites Hebbian plasticity and

long-term potentiation (Brown, Zhao, & Leung, 2010). Crucially, policies were

---

[3] To break the symmetry of the uniform distribution, we added small amounts of Gaussian noise (with a variance of .001) to avoid getting stuck in local free energy minima during learning.

restricted during learning, such that the agent could not select reporting actions;

thus, learning was driven entirely by repeated exposure to different feature

combinations. We evaluated successful learning in two ways. First, we compared the

'A' matrix learned by the model to that of the generative process. Second, we

disabled learning after various trial numbers (i.e., such that concentration

parameters no longer accumulated) and enabled reporting. We then evaluated

reporting accuracy with 20 trials for each of the 8 concepts.

As shown in Figure 4A, the 'A' matrix (likelihood) mapping – learned by the

agent – and the column for sturgeon in particular, strongly resembled that of the

generative process. When first evaluating reporting, the model was 100 % accurate

across 20 reporting trials, when exposed to a sturgeon (reporting accuracy when

exposed to each of the other animals also remained at 100%) and first reached this

level of accuracy after around 50 exposures to all 8 animals (with equal probability)

(figure 4B). The agent also always chose to report specific categories (i.e., it never

chose to only report bird or fish). Model performance was stable over 8 repeated

simulations.

Crucially, during learning, the agent was not told which state was generating

its observations. This meant that it had to solve both an inference and a learning

problem. First, it had to infer whether a given feature combination was better

explained by an existing concept, or by a concept that predicts features uniformly. In

other words, it had to decide that the features were sufficiently different – from

things it had seen before – to assign it a new hypothetical concept. Given that a novel

state is only inferred when another state is not a better explanation, this precludes

learning 'duplicate' states that generate the same patterns of observations. The

second problem is simpler. Having inferred that these outcomes are caused by

something new, the problem becomes one of learning a simple state-outcome

mapping through accumulation of Dirichlet parameters.

To examine whether this result generalized, we repeated these simulations

under conditions in which the agent had to learn more than one concept. When the

model needed to learn one bird (parakeet) and one fish (minnow), the model was

also able to learn the appropriate likelihood mapping for these 2 concepts (although

note that, because the agent did not receive feedback about the state it was in during

learning, the new feature mappings were often not assigned to the same columns as

in the generative process; see figure 4A). Reporting also reached 100% accuracy,

but required a notably greater number of trials. Across 8 repeated simulations, the

mean accuracy reached by the model after 2000 trials was 98.75% (SD = 2%).

A

B

Figure 4. (A) illustration of representative simulation results in which the agent successfully
learned 1, 2, or 4 new animal concept categories with no prior knowledge beforehand. The
generative process is shown in the upper right, illustrating the feature combinations to be
learned. Pre-learning, either 1, 2 or 4 columns in the likelihood mapping began as a flat
distribution with a slight amount of Gaussian noise. The agent was then provided with 2000
observations of the 8 animals with equal probability. Crucially, the agent was prevented
from providing verbal reports during these 2000 trials and thus did not receive feedback
about the true identity of the animal. Thus, learning was driven completely by repeated
exposure in an unsupervised manner. Also note that, while the agent was successful at
learning the new concepts, it did not always assign the new feature patterns to the same
columns as illustrated in the generative process. This is to be expected given that the agent
received no feedback about the true hidden state that generated her observations. (B)
illustration of how reporting accuracy, and the proportion of basic category and specific
category responses, changed as a function of repeated exposures. This was accomplished by
taking the generative model at a number of representative trials and then testing it with 20
observations of each animal in which reporting was enabled. As can be seen, maximal
accuracy was achieved much more quickly when the agent had to learn fewer concepts.
When it had learned 4 concepts, it also began by reporting the general categories and then
subsequently became sufficiently confident to report the more specific categories.
507

When the model needed to learn all 4 birds, performance varied somewhat

more when the simulations were repeated. The learned likelihood mappings after

2000 trials always resembled that of the generative process, but with variable levels

of precision; notably, the model again assigned different concepts to different

columns relative to the generative process, as would be expected when the agent is

not given feedback about the state she is in. Over 8 repeated simulations, the model

performed well in 6 (92.50 % – 98.8 % accuracy) and failed to learn one concept in

the other 2 (72.50 % accuracy in each) due to overgeneralization (e.g., mistaking

parrot for Hawk in a majority of trials; i.e., the model simply learned that there are

large birds). Figure 4A and 4B illustrate representative results when the model was

successful (note: the agent never chose to report basic categories in the simulations

where only 1 or 2 concepts needed to be learned).

To further assess concept learning, we also tested the agent's ability to

successfully avoid state duplication. That is, we wished to confirm that the model

would only learn a new concept if actually presented with a new animal for which it

did not already have a concept. To do so, we equipped the model with knowledge of

seven out of the eight concept categories, and then repeatedly exposed it only to the

animals it already knew over 80 trials. We subsequently exposed it to the eighth

animal (Hawk) for which it did not already have knowledge over 20 additional

trials. As can be seen in figure 5, the unused concept column was not engaged during

the first 80 trials (bottom left and middle). However, in the final 20 trials, the agent

correctly inferred that her current conceptual repertoire was unable to explain her

new pattern of observations, leading the unused concept column to be adumbrated

and the appropriate state-observation mapping to be learned (bottom right). We

repeated these simulations under conditions in which the agent already had

knowledge of six, five, or four concepts. In all cases, we observed that unused

concept columns were never engaged inappropriately.

536
Figure 5. Illustration of representative simulation results when the agent had to avoid
inappropriately learning a new concept (i.e., avoid state duplication) after only being
exposed to animals for which it already had knowledge. Here the agent began with prior
knowledge about seven concept categories and was also equipped with an eighth column
that could be engaged to learn a new concept category (bottom left). The agent was then
presented with several instances of each of the seven animals that she already knew (80
trials in total). In this simulation, the agent was successful in assigning each stimulus to an
animal concept she had already acquired and did not engage the unused concept 'slot'
(bottom middle). Finally, the agent was presented with a new animal (a hawk) that she did
not already know over 20 trials. In this case, the agent successfully engaged the additional
column (i.e., she inferred that none of the concepts she possessed could account for her new
observations) and learned the correct state-observation mapping (bottom right).
549

Crucially, these simulations suggest that adaptive concept learning needs to

be informed by existing knowledge about other concepts, such that a novel concept

should only be learned if observations cannot be explained with existing conceptual

knowledge. Here, this is achieved via the interplay of inference and learning, such

that agents initially have to infer whether to assign an observation to an existing

concept, and only if this is not possible is an 'open slot' employed to learn about a

novel concept.

*Increasing granularity*

Next, to explore the model's ability to increase the granularity of its concept

space, we first equipped the model with only the distinction between birds and fish

(i.e., the rows of the likelihood mapping corresponding to color and size features

were flattened in the same manner described above). We then performed the same

procedure used in our previous simulations. As can be seen in Figure 6A (bottom

left), the 'A' matrix learned by the model now more strongly resembled that of the

generative process. Figure 6B (bottom) also illustrates reporting accuracy and the

proportion of basic and specific category reports as a function of trial number. As

can be seen, the agent initially only reported general categories, and became

sufficiently confident to report specific categories after roughly 50 – 100 trials, but

her accuracy increased gradually over the next 1000 trials (i.e., the agent reported

specific categories considerably before its accuracy improved). Across 8 repeated

simulations, the final accuracy level reached was between 93% – 98% in 7

simulations, but the model failed to learn one concept in the 8th case, with 84.4%

overall accuracy (i.e., a failure to distinguish between pigeon and parakeet, and

therefore only learned a broader category of "small birds").

To assess whether learning basic categories first was helpful in subsequently

learning specific categories, we also repeated this simulation without any initial

knowledge of the basic categories. As exemplified in figure 6A and 6B, the model

tended to perform reasonably well, but most often learned a less precise likelihood

mapping and reached a lower reporting accuracy percentage after 2000 learning

trials (across 8 repeated simulations: mean = 81.21%, SD = 6.39%, range from

68.80% – 91.30%). Thus, learning basic concept categories first appeared to

facilitate learning more specific concepts later.

A

583
584

B
## Increasing Granularity:
### Reporting Accuracy as a Function of Trial Number (without feedback)

**No Prior Knowledge**

**After Learning Basic Categories**

Figure 6. (A, left) Illustration of representative simulation results when the agent had to learn to increase the granularity of her concept space. Here the agent began with prior knowledge about the basic concept categories (i.e., she had learned the broad categories of "bird" and "fish") but had not learned the feature patterns (i.e., rows) that differentiate different types of birds and fish. Post learning (i.e., after 2000 exposures), the agent did successfully learn all of the more granular concept categories, although again note that specific concepts were assigned to different columns then depicted in the generative process due to the unsupervised nature of the learning. (A, right) illustration of the analogous learning result when the agent had to learn all 8 specific categories without prior knowledge of the general categories. Although moderately successful, learning tended to be more difficult in this case. (B) Representative plots of reporting accuracy in each of the 2 learning conditions as a function of the number of exposures. As can be seen, when the model starts out with prior knowledge about basic categories, it slowly becomes sufficiently confident to start reporting the more specific categories, and its final accuracy is high. In contrast, while the agent that did not start out with any prior knowledge of the general categories also grew confident in reporting specific categories over time, her final accuracy levels tended to be lower. In both cases, the agent began reporting specific categories before she achieved significant accuracy levels, therefore showing some initial overconfidence.

Overall, these findings provide a proof of principle that this sort of active

inference scheme can add concepts to a state space in an unsupervised manner (i.e.,

without feedback) based purely on (expected) free energy minimization. In this

case, it was able to accomplish this starting from a completely uninformative

likelihood distribution. It could also learn more granular concepts after already

acquiring more basic concepts, and our results suggest that learning granular

concepts may be facilitated by first learning basic concepts (e.g., as in currently

common educational practices).

The novel feature of this generative model involved 'building in' a number of

"reserve" hidden state levels. These initially had uninformative likelihood mappings;

yet, if a new pattern of features was repeatedly observed, and the model could not

account for this pattern with existing (informative) state-observation mappings,

these additional hidden state levels could be engaged to improve the model's

explanatory power. This approach therefore accommodates a simple form of

structure learning (i.e., model expansion).

**Integrating model expansion and reduction**

We next investigated ways in which model expansion could be combined with

Bayesian model reduction (KJ Friston, Lin, et al., 2017) – allowing the agent to adjust

her model to accommodate new patterns of observations, while also precluding

unnecessary conceptual complexity (i.e., over-fitting). To do so, we again allowed

the agent to learn from 2000 exposures to different animals as described in the

previous section – but also allowed the model to learn its 'D' matrix (i.e., accumulate

concentration parameters reflecting prior expectations over initial states). This

allowed an assessment of the agent's probabilistic beliefs about which hidden state

factor levels (animals) she had been exposed to. In different simulations, the agent

was only exposed to some animals and not others. We then examined whether a

subsequent model reduction step could recover the animal concepts presented

during the simulation; eliminating those concepts that were unnecessary to explain

the data at hand. The success of this 2-step procedure could then license the agent to

"reset" the unnecessary hidden state columns after concept acquisition, which

would have accrued unnecessary likelihood updates during learning. Doing so

would allow the optimal ability for those "reserve" states to be appropriately

engaged, if and when the agent was exposed to truly novel stimuli.

The 2nd step of this procedure was accomplished by applying Bayesian model

reduction to the 'D' matrix concentration parameters after learning. This is a form of

post-hoc model optimization (K. J. Friston et al., 2016; Karl Friston, Parr, & Zeidman,

2018) that rests upon estimation of a 'full' model, followed by analytic computation

of the evidence that would have been afforded to alternative models (with

alternative, 'reduced', priors) had they been used instead. Mathematically, this

procedure is a generalization of things like automatic relevance determination (Karl

Friston, Mattout, Trujillo-Barreto, Ashburner, & Penny, 2007; Wipf & Rao, 2007) or

the use of the Savage Dickie ratio in model comparison (Cornish & Littenberg,

2007). It is based upon straightforward probability theory and, importantly, has a

simple physiological interpretation; namely, synaptic decay and the elimination of

unused synaptic connections. In this (biological) setting, the concentration

parameters of the implicit Dirichlet distributions can be thought of as synaptic tags.

For a technical description of Bayesian model reduction techniques and their

proposed neural implementation, see (KJ Friston, Lin, et al., 2017; Hobson & Friston,

2012; Hobson, Hong, & Friston, 2014); see the left panel of Figure 2 for additional

details).

The posterior concentration parameters were compared to the prior

distribution for a full model (i.e., a flat distribution over 8 concepts) and prior

distributions for possible reduced models (i.e., which retained different possible

combinations of some but not all concepts; technically, reduced models were

defined such that the to-be-eliminated concepts were less likely than the to-be-

retained concepts). If Bayesian model reduction provided more evidence for one or

more reduced models, the reduced model with the most evidence was selected.

Note: an alternative would be to perform model reduction on the 'A' matrix, but this

is more complex due to a larger space of possible reduced models; it also does not

address the question of the number of hidden state levels to retain in a

straightforward manner.

In our first simulation, we presented our agent with all animals except for

parakeets with equal probability over 2000 trials. When compared to the full model,

the winning model corresponded to the correct 7-animal model matching the

generative process in 6/8 cases (log evidence differences ranged from -3.12 to -8.3),

and in 2/8 cases it instead selected a 6-animal model due to a failure to distinguish

between 2 specific concepts during learning (log evidence differences = -5.30, -

7.70). Figure 7 illustrates the results of a representative successful case). In the

successful cases, this would correctly license the removal of changes in the model's

'A' and 'D' matrix parameters for the 8th animal concept during learning in the

previous trials.  Similar results were obtained whenever any single animal type was

absent from the generative process.

Figure 7. Representative illustrations of simulations in which the agent performed Bayesian
model reduction after learning. In these simulations, the agent was first exposed to 2000
trials in which either 7, 6, or 5 animals were actually presented (i.e., illustrated in the top
row, where only the white columns had nonzero probabilities in the generative process). In
each case, model reduction was often successful at identifying the reduced model with the
correct number of animal types presented (bottom row, where black columns should be
removed) based on how much evidence it provided for the posterior distribution over
hidden states learned by the agent (2nd row). This would license the agent to reset
the unneeded columns in its likelihood mapping (3rd row) to their initial state (i.e., a
flat distribution over features) such that they could be engaged if/when new types
of animals began to be observed (i.e., as in the simulations illustrated in the previous
sections).

In a second simulation, the generative process contained 2 birds and all 4

fish. Here, the correct reduced model was correctly selected in 6/8 simulations (log

evidence differences range from -.96 to -8.24, with magnitudes greater than -3 in

5/6 cases), whereas it incorrectly selected the 5-animal model in 2 cases (log

evidence differences = -3.54, -4.50). In a third simulation, the generative process

contained 1 bird and all 4 fish. Here, the correct reduced model had the most

evidence in only 3/8 simulations (log evidence differences = -4.10, -4.11, -5.48),

whereas a 6-animal model was selected in 3/8 cases and a 3-animal and 7-animal

model were each selected once (log evidence differences > -3.0). Figure 7 also

illustrates representative examples of correct model recovery in these 2nd and 3rd

simulations.

While we have used the terms 'correct' and 'incorrect' above to describe the

model used to generate the data, we acknowledge that 'all models are wrong' (Box,

Hunter, & Hunter, 2005), and that the important question is not whether we can

recover the 'true' process used to generate the data, but whether we can arrive at

the simplest but accurate explanation for these data. The failures to recover the

'true' model highlighted above may reflect that a process other than that used to

generate the data could have been used to do so in a simpler way. Simpler here

means we would have to diverge to a lesser degree, from our prior beliefs, in order

to explain the data under a given model, relative to a more complex model. It is

worth highlighting the importance of the word *prior* in the previous sentence. This

means that the simplicity of the model is sensitive to our prior beliefs about it. To

illustrate this, we repeated the same model comparisons as above, but with precise

beliefs in an 'A' matrix that complies with that used to generate the data. Specifically,

we repeated the three simulations above but only enabled 'D' matrix learning (i.e.,

the model was already equipped with the 'A' matrix of the generative process). In

each case, Bayesian model reduction now uniquely identified the correct reduced

model in 100% of cases.

These results demonstrate that – after a naïve model has expanded its hidden

state space to include likelihood mappings and initial state priors for a number of

concept categories – Bayesian model reduction can subsequently be used to

eliminate any parameter updates accrued for *one or two* redundant concept

categories. In practice, the 'A' and 'D' concentration parameters for these redundant

categories could be reset to their default pre-learning values – and could then be re-

engaged if new patterns of observations were repeatedly observed in the future.

However, when three concepts should have been removed, Bayesian model

reduction was much less reliable. This appeared to be due to imperfect 'A' matrix

learning, when occurring simultaneously with the (resultingly noisy) accumulation

of prior expectations over hidden states – as a fully precise 'A' matrix led to correct

model reduction in every case tested (i.e., suggesting that this type of model

reduction procedure could be improved by first allowing state-observation learning

to proceed alone, then subsequently allowing the model to learn prior expectations

over hidden states, which could then be used in model reduction).

**Can concept acquisition allow for generalization?**

One important ability afforded by concept learning is generalization. In a

final set of simulations, we asked if our model of concept knowledge could account

for generalization. To do so, we altered the model such that it no longer reported

what it saw, but instead had to answer a question that depended on generalization

from particular cross-category feature combinations. Specifically, the model was

shown particular animals and asked: "could this be seen from a distance?" The

answer to this question depended on both size and color, such that the answer was

yes only for colorful, large animals (i.e., assuming small or gray animals would blend

in with the sky or water and be missed).

Crucially, this question was asked of animals that the model had not been

exposed to, such that it had to generalize from knowledge it already possessed (see

Figure 8). To simulate and test for this ability, we equipped the model's 'A' matrix

with expert knowledge of 7 out of the 8 animals (i.e., as if these concepts had been

learned previously, as in our simulations above). The $8^{th}$ animal was unknown to the

agent, in that it's likelihood mapping was set such that the $8^{th}$ animal state "slot"

predicted all observations equally (i.e., with a small amount of Gaussian noise, as

above).  In one variant, the model possessed all concepts except for "parrot," and it

knew that the answer to the question was yes for "whale shark" but not for any

other concept it knew. To simulate one-shot generalization, learning was disabled

and a parrot (which it had never seen before) was presented 20 times to see if it

would correctly generalize and answer "yes" in a reliable manner. In another

variant, the model had learned all concepts except "minnow" and was tested the

same way to see if it would reliably provide the correct "no" response.

Here, we observed that in both of these cases (as well as all others we tested)

the model generalized remarkably well. It answered "yes" and "no" correctly in

100% of trials. Thus, the agent did not simply possess concepts to explain things it

saw. It instead demonstrated generalizable knowledge and could correctly answer

questions when seeing a novel stimulus.

Figure 8. Depiction of simulations in which we tested the agent's ability to generalize from
prior knowledge and correctly answered questions about new animals to which she had not
previously been exposed. In the simulations, the generative model was modified so that the
agent instead chose to report either "yes" or "no" to the question: "could this animal be seen
from a distance?" Here, the answer was only yes if the animal was both large and colorful.
We observed that when the agent started out with no knowledge of parrots it still correctly
answered this question 100% of the time, based only on its knowledge of other animals.
Similarly, when it started with no knowledge of minnows, it also correctly reported "no"
100% of the time. Thus, the agent was able to generalize from prior knowledge with no
additional learning.

**Open questions and relation to other theoretical accounts of concept learning**

As our simulations show, this model allows for learning novel concepts (i.e.,

novel hidden states) based on assigning one or more 'open slots' that can be utilised

to learn novel feature combinations. In a simple example, we have shown that this

setup offers a potential computational mechanism for 'model expansion'; i.e., the

process of expanding a state space to account for novel instances in perceptual

categorisation. We also illustrated how this framework can be combined with model

reduction, which may be a mechanism for 're-setting' these open slots based on

recent experience.

This provides a first step towards understanding how agents flexibly expand

or reduce their model to adapt to ongoing experience. Yet, several open questions

remain, which have partly been addressed in previous work. For example, the

proposed framework resonates with previous similarity-based accounts of concept

learning. Previous work has proposed a computational framework for arbitrating

between assigning an observation to a previously formed memory or forming a

novel (hidden) state representation (S. J. Gershman, Monfils, Norman, & Niv, 2017),

based on evidence that this observation was sampled from an existing or novel

latent state. This process is conceptually similar to our application of Bayesian

model reduction over states. In the present framework, concept learning relies on a

process based on inference and learning. First, agents have to *infer* whether ongoing

observations can be sufficiently explained by existing conceptual knowledge – or

speak to the presence of a novel concept that motivates the use of an 'open slot'.

This process is cast as inference on (hidden) states. Second, if the agent infers that

there is a novel concept that explains current observations, it has to *learn* about the

specific feature configuration of that concept (i.e., novel state). This process

highlights the interplay between inference, which allows for the acquisition of

knowledge on a relatively short timescale, and learning, which allows for knowledge

acquisition on a longer and more stable timescale.

810            Similar considerations apply to the degree of 'similarity' of observations. In

the framework proposed here, we have assumed that the feature space of

observations is already learned and fixed. However, these feature spaces have to be

learned in the first place, which implies learning the underlying components or

feature dimensions that define observations. This relates closely to notions of

structure learning as dimensionality reduction based on covariance between

observations, as prominently discussed in the context of spatial navigation (Behrens

et al., 2018; Dordek, Soudry, Meir, & Derdikman, 2016; Stachenfeld et al., 2016;

Whittington, Muller, Mark, Barry, & Behrens, 2018).

819            Another important issue is how such abstract conceptual knowledge is

formed across different contexts or tasks. For example, the abstract concept of a

'bird' will be useful for learning about the fauna in a novel environment, but specific

types of birds – tied to a previous context – might be less useful in this regard. This

speaks to the formation of abstract, task-general knowledge that results from

training across different tasks, as recently discussed in the context of meta-

reinforcement learning (Ritter, Wang, Kurth-Nelson, & Botvinick, 2018; J X Wang et

al., 2016) with a putative link to the prefrontal cortex (Jane X. Wang et al., 2018). In

the present framework, such task-general knowledge would speak to the formation

of a hierarchical organisation that allows for the formation of conceptual knowledge

both within and across contexts. Also note that our proposed framework depends

on a pre-defined state space, including a pre-defined set of 'open slots' that allow for

novel context learning. The contribution of the present framework is to show how

these 'open slots' can be used for novel concept learning and be re-set based on

model reduction. It will be important to extend this approach towards learning the

structure of these models in the first place, including the appropriate number of

'open slots' (i.e., columns of the A-matrix) for learning in a particular content

domain and the relevant feature dimensions of observations (i.e., rows of A-matrix).

(Note: In addition to ontogenetic learning, in some cases structural priors regarding

the appropriate number of open slots [and relevant feature inputs for learning a

given state space of open slots] might also reflect inherited [i.e.,

genetically/developmentally pre-specified] patterns of structural neuronal

connectivity – based on what was adaptive within the evolutionary niche of a given

species – which could then be modified based on subsequent experience.)

This corresponds to a potentially powerful and simple application of

Bayesian model reduction, in which candidate models (i.e., reduced forms of a full

model) are readily identifiable based upon the similarity between the likelihoods

conditioned upon different hidden states. If two or more likelihoods are sufficiently

similar, the hidden states can be merged (by assigning the concentration

parameters accumulated during experience-dependent learning to one or other of

the hidden states). The ensuing change in model evidence scores the reduction in

complexity. If this reduction is greater than the loss of accuracy – in relation to

observations previously encountered – Bayesian model reduction will, effectively,

merge one state into another; thereby freeing up a state for the learning of new

concepts. We will demonstrate this form of structure learning via Bayesian model

reduction in future work.

We must also highlight here that cognitive science research on concept and

category learning has a rich empirical and theoretical history, including many

previously proposed formal models. While our primary focus has been on using

concept learning as an example of a more general approach by which state space

expansion and reduction can be implemented within future active inference

research, it is important to recognize this previous work and highlight where it

overlaps with the simulations we've presented. For example, our results suggesting

that first learning general categories facilitates the learning of more specific

categories relates to both classic and contemporary findings showing that children

more easily acquire "basic" and "superordinate" (e.g., dog, animal) concepts before

learning "subordinate" (e.g., chihuahua) concepts (Mervis & Rosch 1981; Murphy

2016), and that this may involve a type of "bootstrapping" process (Beck 2017).

Complementary work has also highlighted ways in which learning new words

during development can invoke a type of "placeholder" structure, which then

facilitates the acquisition of a novel concept – which bears some resemblance to our

notion of blank "concept slots" that can subsequently acquire meaningful semantics

(Gelman & Roberts 2017).

There is also a series of previously proposed formalisms within the literature

on category learning. For example, two previously proposed models – the "rational

model" (Anderson 1991; Sanborn et al. 2010) and the SUSTAIN model (Love et al.

2004) – both describe concept acquisition as involving cluster creation mechanisms

that depend on statistical regularities during learning and that use probabilistic

updating. The updating mechanisms within SUSTAIN are based on

surprise/prediction-error in the context of both supervised and unsupervised

learning. This model also down-weights previously created clusters when their

associated regularities cease to be observed in recent experience. Although not built

in intentionally, this type of mechanism emerges naturally within our model in two

ways. First, when a particular hidden state ceases to be inferred, concentration

parameters will accumulate to higher values for other hidden states in the **D** matrix,

reflecting relatively stronger prior expectations for hidden states that continue to be

inferred – which would favor future inference of those states over those absent from

recent experience. Second, if one pattern of observations were absent from recent

experience (while other patterns continued to be observed), concentration

parameters in the **A** matrix would also accumulate to higher values for patterns that

continued to be observed – resulting in relatively less confidence in the state-

outcome mapping for the less-observed pattern. (However, with respect to this

latter mechanism, so long as this mapping was sufficiently precise and distinct from

others [i.e., it had previously been observed many times farther in the past], this

would not be expected to prevent successful inference if this pattern were observed

again.)

It is also worth highlighting that, as our model is intended primarily as a

proof of concept and a demonstration of an available model expansion/reduction

approach that can be used within active inference research, it does not explicitly

incorporate some aspects – such as top-down attention – that are of clear

importance to cognitive learning processes, and that have been implemented in

previous models. For example, the adaptive resonance theory (ART) model

(Grossberg 1987) was designed to incorporate top-down attentional mechanisms

and feedback mechanisms to address a fundamental knowledge acquisition problem

– the temporal instability of previously learned information that can occur when a

system also remains sufficiently plastic to learn new (and potentially overlapping)

information. While our simulations do not explicitly incorporate these additional

complexities, there are clear analogues to the top-down and bottom-up feedback

exchange in ART within our model (e.g., the prediction and prediction-error

signaling within the neural process theory associated with active inference). ART

addresses the temporal instability problem primarily through mechanisms that

learn top-down expectancies that guide attention and match them with bottom-up

input patterns – which is quite similar to the prior expectations and likelihood

mappings used within active inference.

As an emergent property of the "first principles" approach in active

inference, our model therefore naturally incorporates the top-down effects in ART

simulations, which have been used to account for known context effects on

categorical perception within empirical studies (McClelland & Rumelhart 1981).

This is also consistent with more recent work on cross-categorization (Shafto et al.

2011), which has shown that human category learning is poorly accounted for by

both a purely bottom-up process (attempting to explain observed features) and a

purely top-down approach (involving attention-based feature selection) – and has

instead used simulations to show that a Bayesian joint inference model better fits

empirical data.

Other proposed Bayesian models of concept learning have also had

considerable success in predicting human generalization judgments (Goodman et al.

2008). The proof of concept model presented here has not been constructed to

explicitly compete with such models. It will be an important direction for future

work to explore the model's ability to scale up to handle more complex concept

learning problems. Here we simply highlight that the broadly Bayesian approach

within our model is shared with other models that have met with considerable

success – supporting the general plausibility of using this approach within active

inference research to model and predict the neural basis of these processes (see

below).

934                    **Potential advantages of the approach**

The present approach may offer some potential theoretical and empirical

advantages in comparison to previous work. One theoretical advantage corresponds

to the parsimony of casting this type of structure learning as an instance of Bayesian

model selection. When integrated with other aspects of the active inference

framework, this entails that perceptual inference, active learning, and structure

learning are all expressions of the same principle; namely, the minimization of

variational free energy, over three distinct timescales. A second, related theoretical

advantage is that, when this type of structure learning is cast as Bayesian model

selection/reduction, there is no need to invoke additional procedures or schemes

(e.g., nonparametric Bayes or 'stick breaking' processes; (S. Gershman & Blei,

2012)). Instead, a generative model with the capacity to represent a sufficiently

complex world will automatically learn causal structure in a way that contextualizes

active inference within active learning, and active learning within structure

learning.

Based on the process theories summarized in Figure 2, the present model

would predict that the brain contains "reserve" cortical columns and synapses (most

likely within secondary sensory and association cortices) available to capture new

patterns in observed features. To our knowledge, no direct evidence supporting the

presence of unused cortical columns in the brain has been observed, although the

generation of new neurons (with new synaptic connections) is known to occur in

the hippocampus (Chancey et al., 2013). "Silent synapses" have also been observed

in the brain, which does appear consistent with this prediction; such synapses can

persist into adulthood and only become activated when new learning becomes

necessary (e.g., see (Chancey et al., 2013; Funahashi, Maruyama, Yoshimura, &

Komatsu, 2013; Kerchner & Nicoll, 2008)). One way in which this idea of "spare

capacity" or "reserve" cortical columns might be tested in the context of

neuroimaging would be to examine whether greater levels of neural activation –

within conceptual processing regions – are observed after learning additional

concepts, which would imply that additional populations of neurons become

capable of being activated. In principle, single-cell recording methods might also test

for the presence of neurons that remain at baseline firing rates during task

conditions, but then become sensitive to new stimuli within the relevant conceptual

domain after learning.

Figure 9 provides a concrete example of two specific empirical predictions

that follow from simulating the neural responses that should be observed within our

concept learning task under these process theories. In the left panel, we plot the

firing rates (darker = higher firing rate) and local field potentials (rate of change in

firing rates) associated with neural populations encoding the probability of the

presence of different animals that would be expected across a number of learning

trials. In this particular example, the agent began with knowledge of the basic

categories of 'bird' and 'fish,' but needed to learn the eight more specific animal

categories over 50 interleaved exposures to each animal (only 10 equally spaced

learning trials involving the presentation of a parakeet are shown for simplicity). As

can be seen, early in learning the firing rates and local field potentials remain at

baseline levels; in contrast, as learning progresses, these neural responses take a

characteristic shape with more and more positive changes in firing rate in the

populations representing the most probable animal, while other populations drop

further and further below baseline firing rates.

The right panel depicts a similar simulation, but where the agent was

allowed to self-report what it saw on each trial (for clarity of illustration, we here

show 12 equally spaced learning trials for parakeet over 120 total trials). Enabling

policy selection allowed us to simulate expected phasic dopamine responses during

the task, corresponding to changes in the precision of the probability distribution

over policies after observing a stimulus on each trial. As can be seen, during early

trials the model predicts small firing rate increases when the agent is confident in its

ability to correctly report the more general animal category after observing a new

stimulus, and firing rate decreases when the agent becomes less confident in one

policy over others (i.e., as confidence in reporting the specific versus general

categories becomes more similar). Larger and larger phasic dopaminergic responses

are then expected as the agent becomes more and more confident in her ability to

correctly report the specific animal category upon observing a new stimulus. It will

be important for future neuroimaging studies to test these predictions in this type of

concept learning/stimulus categorization task.

1000

Figure 9. Simulated neuronal firing rates, local field potentials, and dopaminergic responses across learning trials based on the neural process theory associated with active inference that is summarized in Figure 2. The top left panel displays the predicted firing rates (darker = higher firing rate) of neural populations encoding the probability of each hidden state over 50 interleaved exposures to each animal (only 10 equally spaced learning trials involving the presentation of a parakeet are shown for simplicity) in the case where the agent starts out with knowledge of the basic animal categories but must learn the more specific categories. As can be seen, initially each of the four neural populations encoding possible bird categories (i.e., one row per possible category) have equally low firing rates (gray); as learning continues, firing rates increase for the 'parakeet' population and decrease for the others. The bottom left panel illustrates the predicted local field potentials (based on the rate of change in firing rates) that would be measured across the task. The top right panel displays the predicted firing rates of neural populations in an analogous simulation in which reporting policies were enabled (for clarity of illustration, we here show 12 equally spaced learning trials for parakeet over 120 total trials). Enabling policy selection allowed us to simulate the phasic dopaminergic responses (reporting changes in the precision of the probability distribution over policies) predicted to occur across learning trials; here the agent first becomes confident in her ability to correctly report the general animal category upon observing a stimulus, then becomes unsure about reporting specific versus general categories, and then becomes confident in her ability to report the specific categories.

1023

1024                                **Discussion**

1025

The Active Inference formulation of concept learning presented here

demonstrates a simple way in which a generative model can acquire both basic and

highly granular knowledge of the hidden states/causes in its environment. In

comparison to previous theoretical work using active inference (e.g., (M. Mirza,

Adams, Mathys, & Friston, 2016; Parr & Friston, 2017; Schwartenbeck, FitzGerald,

Mathys, Dolan, & Friston, 2015)), the novel aspect of our model was that it was

further equipped with "reserve" hidden states initially devoid of content (i.e., these

states started out with uninformative likelihood mappings that predicted all

outcomes with roughly equal probability). Over multiple exposures to different

stimuli, these hidden states came to acquire conceptual content that captured

distinct statistical patterns in the features of those stimuli. This was accomplished

via the model's ability to infer when its currently learned hidden states were unable

to account for a new observation, leading an unused hidden state column to be

engaged that could acquire a new state-observation mapping.

Crucially, the model was able to start with some concepts and then expand its

representational repertoire to learn others – but would only do so when a new

stimulus was observed. This is conceptually similar to nonparametric Bayesian

learning models, such as the "Chinese Room" process and the "Indian Buffet"

process, that can also infer the need to invoke additional hidden causes with

additional data (S. Gershman & Blei, 2012). These statistical learning models do not

need to build in additional "category slots" for learning as in our model and can, in

principle, entertain infinite state spaces. On the other hand, it is less clear at present

how the brain could implement this type of learning. An advantage of our model is

that learning depends solely on biologically plausible Hebbian mechanisms (for a

possible neural implementation of model reduction, see (KJ Friston, Lin, et al., 2017;

Hobson & Friston, 2012; Hobson et al., 2014)).

The distinction between nonparametric Bayesian learning and the current

active learning scheme may be important from a neurodevelopmental perspective

as well. In brief, structure learning in this paper starts with a generative model with

a type of structural prior reflecting a specific amount of built in 'spare capacity',

where uncommitted or naive conceptual 'slots' are used to explain the sensorium,

during optimization of free energy or model evidence. In contrast, nonparametric

Bayesian approaches add new slots when appropriate. One might imagine that

neonates are equipped with brains with 'spare capacity' (Baker & Tenenbaum,

2014) that is progressively leveraged during neurodevelopment, much in the spirit

of curriculum learning (Al-Muhaideb & Menai, 2011). This suggestion appears

consistent with previous work demonstrating varying levels of category learning

ability across the lifespan, which has previously been formally modeled as an

individual difference in values of a parameter constraining the ability to form new

clusters in response to surprising events (Love & Gureckis 2007) – which bears

similarity to the idea of capacity limitations arising from finite numbers of concept

slots in our model.

In this sense, the current approach to structure learning may be better

considered as active learning with generative models that are equipped with a large

number of available hidden states capable of acquiring content, which are then

judiciously reduced/reset – via a process of Bayesian model reduction.

Furthermore, as in the acquisition of expertise, our model can also begin with broad

category knowledge and then subsequently learn finer-grained within-category

distinctions, which has received less attention from the perspective of the

aforementioned models. Reporting broad versus specific category recognition is

also a distinct aspect of our model – driven by differing levels of uncertainty and an

expectation (preference) not to incorrectly report a more specific category.

Our simulation results also demonstrated that, when combined with

Bayesian model reduction, the model can guard against learning too many

categories during model expansion – often retaining only the number of hidden

causes actually present in its environment – and to keep "reserve" hidden states for

learning about new causes if or when they appear. With perfect "expert" knowledge

of the possible animal types it could observe (i.e., fully precise likelihood mappings

matching the generative process) this was true in general. Interestingly, however,

with an imperfectly learned likelihood mapping, model reduction only succeeded

when the agent had to remove either 1 or 2 concepts from her model; when 3

potential categories needed to be removed, the correct reduced model was

identified less than half the time. It would be interesting to empirically test whether

similar learning difficulties are present in humans.

Neurobiological theories associated with Active Inference also make

predictions about the neural basis of this process (Hobson & Friston, 2012; Hobson

et al., 2014). Specifically, during periods of rest (e.g., daydreaming) or sleep, it is

suggested that, because sensory information is down-weighted, learning is driven

mainly by internal model simulations (e.g., as appears to happen in the phenomenon

of hippocampal replay; (Feld & Born, 2017; Lewis, Knoblich, & Poe, 2018; Pfeiffer &

Foster, 2013)); this type of learning can accomplish a model reduction process in

which redundant model parameters are identified and removed to prevent model

over-fitting and promote selection of the most parsimonious model that can

successfully account for previous observations.  This is consistent with work

suggesting that, during sleep, many (but not all) synaptic strength increases

acquired in the previous day are attenuated (Tononi & Cirelli, 2014). The role of

sleep and daydreaming in keeping "reserve" representational resources available

for model expansion could therefore be especially important to concept learning –

consistent with the known role of sleep in learning and memory (Ackermann &

Rasch, 2014; Feld & Born, 2017; Perogamvros & Schwartz, 2012; Stickgold, Hobson,

Fosse, & Fosse, 2001; Walker & Stickgold, 2010).

In addition, an emergent feature of our model was its ability to generalize

prior knowledge to new stimuli to which it had not previously been exposed. In fact,

the model could correctly generalize upon a single exposure to a new stimulus – a

type of "one-shot learning" capacity qualitatively similar to that observed in humans

(Landau, Smith, & Jones, 1988; E. Markman, 1989; Xu & Tenenbaum, 2007b). While

it should be kept in mind that the example we have provided is very simple, it

demonstrates the potential usefulness of this novel approach. Some other

prominent approaches in machine-learning (e.g., deep learning) tend to require

larger amounts of data (Geman et al., 1992; Hinton et al., 2012; LeCun et al., 2015;

Lecun et al., 1998; Mnih et al., 2015), and do not learn the rich structure that allows

humans to use concept knowledge in a wide variety of generalizable functions

(Barsalou, 1983; Biederman, 1987; Feldman, 1997; Jern & Kemp, 2013; A. B.

Markman & Makin, 1998; Osherson & Smith, 1981; Ward, 1994; Williams &

Lombrozo, 2010). Other recent hierarchical Bayesian approaches in cognitive

science have made progress in this domain, however, by modeling concepts as types

of probabilistic programs (Ghahramani, 2015; Goodman, Tenenbaum, &

Gerstenberg, 2015; Lake et al., 2015).

It is important to note that this model is deliberately simple and is meant

only to represent a proof of principle that categorical inference and conceptual

knowledge acquisition can be modeled within this particular neurocomputational

framework, and to present this approach as a potentially useful tool in future active

inference research. We chose a particular set of feature combinations to illustrate

this, but it remains to be demonstrated that learning in this model would be equally

successful with a larger feature space and set of learnable hidden causes. Due to

limited scope, we have also not thoroughly addressed all other overlapping lines of

research. For example, work on exemplar models of concepts has also led to other

computational approaches. As one example, the EBRW model (Nosofsky & Palmeri

1997) has demonstrated ways of linking exemplar learning to drift diffusion models.

Another model within this line of research is the ALCOVE model (Nosofsky et al.

1994) – an error-driven connectionist model of exemplar-based category learning

that employs selective attention and learns attentional weights (this model also

built on earlier work; see (Nosofsky 2011)). Yet another connectionist model with

some conceptual overlap to our own is the DIVA model, which learns categories by

recoding observations as task-constrained principle components and uses model fit

for subsequent recognition (Kurtz 2007). It will be important in future work to

examine the strengths and limitations of a scaled-up version of our approach in

relation to these other models.

Finally, another topic for future work would be the expansion of this type of

model to context-specific learning (e.g., with an additional hidden state factor for

encoding distinct contexts). In such cases, regularities in co-occurring features differ

in different contexts and other cues to context may not be directly observable (e.g.,

the same species of bird could be a slightly different color or size in different parts

of the world that otherwise appear similar) – creating difficulties in inferring when

to update previously learned associations and when to instead acquire competing

associations assigned to new contexts. At present, it is not clear whether the

approach we have illustrated would be successful at performing this additional

function, although the process of inferring the presence of a new hidden state level

in a second hidden state factor encoding context would be similar to what we have

shown within a single state factor (for related work on context-dependent

contingency learning, see (S. J. Gershman et al., 2017; S. Gershman, Jones, Norman,

Monfils, & Niv, 2013)). Another point worth highlighting is that we have made

particular choices with regard to various model parameters and the number of

observations provided during learning. Further investigations of the space of these

possible parameter settings will be important. With this in mind, however, our

current modelling results could offer additional benefits. For example, the model's

simplicity could be amenable to empirical studies of saccadic eye movements

toward specific features during novel category learning (e.g. following the approach

of (M. B. Mirza, Adams, Mathys, & Friston, 2018)). This approach could also be

combined with measures of neural activity in humans or other animals, allowing

more direct tests of the neural predictions highlighted above. In addition, the

introduction of exploratory, novelty-seeking, actions could be used to reduce the

number of samples required for learning, with agents selecting those data that are

most relevant.

In conclusion, the Active Inference scheme we have described illustrates

feature integration in the service of conceptual inference: it can successfully

simulate simple forms of concept acquisition and concept differentiation (i.e.

increasing granularity), and it spontaneously affords one-shot generalization.

Finally, it speaks to empirical work in which behavioral tasks could be designed to

fit such models, which would allow investigation of individual differences in concept

learning and its neural basis. For example, such a model can simulate (neuronal)

belief updating to predict neuroimaging responses as we illustrated above; i.e., to

identify the neural networks engaged in evidence accumulation and learning

(Schwartenbeck et al., 2015). In principle, the model parameters (e.g., 'A' matrix

precision) can also be fit to behavioral choices and reaction times – and thereby

phenotype subjects in terms of the priors under which they infer and learn

(Schwartenbeck & Friston, 2016). This approach could therefore advance

neurocomputational approaches to concept learning in several directions.

1184

**Software note**

Although the generative model – specified by the various matrices described in this

paper – changes from application to application, the belief updates are generic and

can be implemented using standard routines (here **spm_MDP_VB_X.m**). These

routines are available as Matlab code in the DEM toolbox of the most recent version

of SPM academic software: http://www.fil.ion.ucl.ac.uk/spm/. The simulations in

this paper can be reproduced (and customized) via running the Matlab code

included here is supplementary material (**Concepts_model.m**).

1193

1194

1195                                   **References**

1196

Ackermann, S., & Rasch, B. (2014). Differential Effects of Non-REM and REM Sleep

on Memory Consolidation? *Current Neurology and Neuroscience Reports*, *14*(2),

430. https://doi.org/10.1007/s11910-013-0430-8

Al-Muhaideb, S., & Menai, M. E. B. (2011). Evolutionary computation approaches to

the Curriculum Sequencing problem. *Natural Computing*, *10*(2), 891–920.

https://doi.org/10.1007/s11047-010-9246-5

Anderson, J. R. (1991). The Adaptive Nature of Human Categorization. *Psychological*

*Review*, *98*(3), 409–429. https://doi.org/10.1037/0033-295X.98.3.409

Baker, C., & Tenenbaum, J. (2014). Modeling human plan recognition using Bayesian

theory of mind. In G. Sukthankar, C. Geib, H. Dui, D. Pynadath, & R. Goldman

(Eds.), *Plan, activity, and intent recognition* (pp. 177–204). Boston: Morgan

Kaufmann.

Barsalou, L. W. (1983). Ad hoc categories. *Memory & Cognition*, *11*(3), 211–227.

https://doi.org/10.3758/bf03196968

Behrens, T. E. J., Muller, T. H., Whittington, J. C. R., Mark, S., Baram, A. B., Stachenfeld,

K. L., & Kurth-Nelson, Z. (2018). What Is a Cognitive Map? Organizing

Knowledge for Flexible Behavior. *Neuron*, *100*(2), 490–509.

https://doi.org/10.1016/J.NEURON.2018.10.002

Biederman, I. (1987). Recognition-by-components: a theory of human image

understanding. *Psychological Review*, *94*(2), 115–147.

https://doi.org/10.1037/0033-295X.94.2.115

Botvinick, M. M., Niv, Y., & Barto, A. C. (2009). Hierarchically organized behavior and

its neural foundations: A reinforcement learning perspective. *Cognition*, *113*(3),

262–280. https://doi.org/10.1016/J.COGNITION.2008.08.011

Box, G. E., Hunter, J. S., & Hunter, W. G. (2005). *Statistics for experimenters. Wiley*

*Series in Probability and Statistics.* Hoboken, NJ.: Wiley.

Brown, T. H., Zhao, Y., & Leung, V. (2010). Hebbian plasticity. In *Encyclopedia of*

*Neuroscience* (pp. 1049–1056). https://doi.org/10.1016/B978-008045046-

9.00796-8

Chancey, J., Adlaf, E., Sapp, M., Pugh, P., Wadiche, J., & Overstreet-Wadiche, L. (2013).

GABA depolarization is required for experience-dependent synapse unsilencing

in adult-born neurons. *The Journal of Neuroscience : The Official Journal of the*

*Society for Neuroscience*, *33*(15), 6614–6622.

https://doi.org/10.1523/JNEUROSCI.0781-13.2013

Conant, C., & Ashbey, W. (1970). Every good regulator of a system must be a model

of that system. *International Journal of Systems Science*, *1*(2), 89–97.

https://doi.org/10.1080/00207727008920220

Cornish, N. J., & Littenberg, T. B. (2007). Tests of Bayesian model selection

techniques for gravitational wave astronomy. *Physical Review D*, *76*(8), 083006.

https://doi.org/10.1103/PhysRevD.76.083006

Dordek, Y., Soudry, D., Meir, R., & Derdikman, D. (2016). Extracting grid cell

characteristics from place cell inputs using non-negative principal component

analysis. *ELife*, *5*(MARCH2016), 1–36. https://doi.org/10.7554/eLife.10094

Feld, G., & Born, J. (2017). Sculpting memory during sleep: concurrent consolidation

and forgetting. *Current Opinion in Neurobiology*, *44*, 20–27.

https://doi.org/10.1016/J.CONB.2017.02.012

Feldman, J. (1997). The Structure of Perceptual Categories. *Journal of Mathematical*

*Psychology*, *41*(2), 145–170. https://doi.org/10.1006/jmps.1997.1154

Friston, K. J., Litvak, V., Oswal, A., Razi, A., Stephan, K. E., van Wijk, B. C. M., …

Zeidman, P. (2016). Bayesian model reduction and empirical Bayes for group

(DCM) studies. *NeuroImage*, *128*, 413–431.

https://doi.org/10.1016/J.NEUROIMAGE.2015.11.015

Friston, Karl, Mattout, J., Trujillo-Barreto, N., Ashburner, J., & Penny, W. (2007).

Variational free energy and the Laplace approximation. *NeuroImage*, *34*(1),

220–234. https://doi.org/10.1016/J.NEUROIMAGE.2006.08.035

Friston, Karl, Parr, T., & Zeidman, P. (2018). Bayesian model reduction. Retrieved

from http://arxiv.org/abs/1805.07092

Friston, Karl, & Penny, W. (2011). Post hoc Bayesian model selection. *NeuroImage*,

*56*(4), 2089–2099. https://doi.org/10.1016/J.NEUROIMAGE.2011.03.062

Friston, KJ. (2010). The free-energy principle: a unified brain theory? *Nature*

*Reviews. Neuroscience*, *11*(2), 127–138. https://doi.org/10.1038/nrn2787

Friston, KJ, FitzGerald, T., Rigoli, F., Schwartenbeck, P., O Doherty, J., & Pezzulo, G.

(2016). Active inference and learning. *Neuroscience and Biobehavioral Reviews*,

*68*, 862–879. https://doi.org/10.1016/j.neubiorev.2016.06.022

Friston, KJ, FitzGerald, T., Rigoli, F., Schwartenbeck, P., & Pezzulo, G. (2017). Active

Inference: A Process Theory. *Neural Computation*, *29*(1), 1–49.

https://doi.org/10.1162/NECO_a_00912

Friston, KJ, Lin, M., Frith, C., Pezzulo, G., Hobson, J., & Ondobaka, S. (2017). Active

Inference, Curiosity and Insight. *Neural Computation*, *29*(10), 2633–2683.

https://doi.org/10.1162/neco_a_00999

Friston, KJ, Parr, T., & de Vries, B. (2017). The graphical brain: Belief propagation

and active inference. *Network Neuroscience*, *1*(4), 381–414.

https://doi.org/10.1162/NETN_a_00018

Funahashi, R., Maruyama, T., Yoshimura, Y., & Komatsu, Y. (2013). Silent synapses

persist into adulthood in layer 2/3 pyramidal neurons of visual cortex in dark-

reared mice. *Journal of Neurophysiology*, *109*(8), 2064–2076.

https://doi.org/10.1152/jn.00912.2012

Geman, S., Bienenstock, E., & Doursat, R. (1992). Neural Networks and the

Bias/Variance Dilemma. *Neural Computation*, *4*(1), 1–58.

https://doi.org/10.1162/neco.1992.4.1.1

Gershman, S., & Blei, D. (2012). A tutorial on Bayesian nonparametric models.

*Journal of Mathematical Psychology*, *56*(1), 1–12.

https://doi.org/10.1016/J.JMP.2011.08.004

Gershman, S. J., Monfils, M.-H., Norman, K. A., & Niv, Y. (2017). The computational

nature of memory modification. *ELife*, *6*. https://doi.org/10.7554/eLife.23763

Gershman, S. J., & Niv, Y. (2010). Learning latent structure: carving nature at its

joints. *Current Opinion in Neurobiology*, *20*(2), 251–256.

https://doi.org/10.1016/J.CONB.2010.02.008

Gershman, S., Jones, C., Norman, K., Monfils, M., & Niv, Y. (2013). Gradual extinction

prevents the return of fear: implications for the discovery of state. *Frontiers in*

*Behavioral Neuroscience*, *7*, 164. https://doi.org/10.3389/fnbeh.2013.00164

Ghahramani, Z. (2015). Probabilistic machine learning and artificial intelligence.

*Nature*, *521*(7553), 452–459. https://doi.org/10.1038/nature14541

Goodman, N. D., Tenenbaum, J. B., & Gerstenberg, T. (2015). *Concepts: New*

*Directions*. (E. Margolis & S. Laurence, Eds.). Cambridge, MA: MIT Press.

Hinton, G., Deng, L., Yu, D., Dahl, G., Mohamed, A., Jaitly, N., … Kingsbury, B. (2012).

Deep Neural Networks for Acoustic Modeling in Speech Recognition: The

Shared Views of Four Research Groups. *IEEE Signal Processing Magazine*, *29*(6),

82–97. https://doi.org/10.1109/MSP.2012.2205597

Hobson, J., & Friston, K. (2012). Waking and dreaming consciousness:

neurobiological and functional considerations. *Progress in Neurobiology*, *98*(1),

82–98. https://doi.org/10.1016/j.pneurobio.2012.05.003

Hobson, J., Hong, C.-H., & Friston, K. (2014). Virtual reality and consciousness

inference in dreaming. *Frontiers in Psychology*, *5*, 1133.

https://doi.org/10.3389/fpsyg.2014.01133

Jern, A., & Kemp, C. (2013). A probabilistic account of exemplar and category

generation. *Cognitive Psychology*, *66*(1), 85–125.

https://doi.org/10.1016/j.cogpsych.2012.09.003

Kemp, C., Perfors, A., & Tenenbaum, J. (2007). Learning overhypotheses with

hierarchical Bayesian models. *Developmental Science*, *10*(3), 307–321.

https://doi.org/10.1111/j.1467-7687.2007.00585.x

Kerchner, G., & Nicoll, R. (2008). Silent synapses and the emergence of a

postsynaptic mechanism for LTP. *Nature Reviews. Neuroscience*, *9*(11), 813–

825. https://doi.org/10.1038/nrn2501

Lake, B. B. M., Salakhutdinov, R., & Tenenbaum, J. J. B. (2015). Human-level concept

learning through probabilistic program induction. *Science (New York, N.Y.)*,

*350*(6266), 1332–1338. https://doi.org/10.1126/science.aab3050

Landau, B., Smith, L. B., & Jones, S. S. (1988). The importance of shape in early lexical

learning. *Cognitive Development*, *3*(3), 299–321.

https://doi.org/10.1016/0885-2014(88)90014-7

LeCun, Y., Bengio, Y., & Hinton, G. (2015). Deep learning. *Nature*, *521*(7553), 436–

444. https://doi.org/10.1038/nature14539

Lecun, Y., Bottou, L., Bengio, Y., & Haffner, P. (1998). Gradient-based learning

applied to document recognition. *Proceedings of the IEEE*, *86*(11), 2278–2324.

https://doi.org/10.1109/5.726791

Lewis, P., Knoblich, G., & Poe, G. (2018). How Memory Replay in Sleep Boosts

Creative Problem-Solving. *Trends in Cognitive Sciences*, *22*(6), 491–503.

https://doi.org/10.1016/j.tics.2018.03.009

Love, B. C., Medin, D. L., & Gureckis, T. M. (2004). SUSTAIN: a network model of

category learning. *Psychological Review*, *111*(2), 309–332.

https://doi.org/10.1037/0033-295X.111.2.309

MacKay, D. J. C., & Peto, L. C. B. (1995). A hierarchical Dirichlet language model.

*Natural Language Engineering*, *1*(03), 289–308.

https://doi.org/10.1017/S1351324900000218

Markman, A. B., & Makin, V. S. (1998). Referential communication and category

acquisition. *Journal of Experimental Psychology. General*, *127*(4), 331–354.

https://doi.org/10.1037/0096-3445.127.4.331

Markman, E. (1989). *Categorization and Naming in Children*. Cambridge, MA: MIT

Press.

McKay, R., & Dennett, D. (2009). The evolution of misbelief. *Behavioral and Brain*

*Sciences*, *32*(06), 493. https://doi.org/10.1017/S0140525X09990975

McNicholas, P. (2016). Model-Based Clustering. *Journal of Classification*, *33*(3), 331–

373. https://doi.org/10.1007/s00357-016-9211-9

Mirza, M., Adams, R., Mathys, C., & Friston, K. (2016). Scene Construction, Visual

Foraging, and Active Inference. *Frontiers in Computational Neuroscience*, *10*, 56.

https://doi.org/10.3389/fncom.2016.00056

Mirza, M. B., Adams, R. A., Mathys, C., & Friston, K. J. (2018). Human visual

exploration reduces uncertainty about the sensed world. *PLOS ONE*, *13*(1),

e0190429. https://doi.org/10.1371/journal.pone.0190429

Mnih, V., Kavukcuoglu, K., Silver, D., Rusu, A. A., Veness, J., Bellemare, M. G., …

Hassabis, D. (2015). Human-level control through deep reinforcement learning.

*Nature*, *518*(7540), 529–533. https://doi.org/10.1038/nature14236

Osherson, D. N., & Smith, E. E. (1981). On the adequacy of prototype theory as a

theory of concepts. *Cognition*, *9*(1), 35–58. https://doi.org/10.1016/0010-

0277(81)90013-5

Parr, T., & Friston, K. (2017). Working memory, attention, and salience in active

inference. *Scientific Reports*, *7*(1), 14678. https://doi.org/10.1038/s41598-

017-15249-0

Parr, T., & Friston, K. (2018). The Anatomy of Inference: Generative Models and

Brain Structure. *Frontiers in Computational Neuroscience*, *12*, 90.

https://doi.org/10.3389/fncom.2018.00090

Parr, T., Markovic, D., Kiebel, S., & Friston, K. (2019). Neuronal message passing

using Mean-field, Bethe, and Marginal approximations. *Scientific Reports*, *9*(1),

1889. https://doi.org/10.1038/s41598-018-38246-3

Perfors, A., Tenenbaum, J., Griffiths, T., & Xu, F. (2011). A tutorial introduction to

Bayesian models of cognitive development. *Cognition*, *120*(3), 302–321.

https://doi.org/10.1016/j.cognition.2010.11.015

Perogamvros, L., & Schwartz, S. (2012). The roles of the reward system in sleep and

dreaming. *Neuroscience & Biobehavioral Reviews*, *36*(8), 1934–1951.

https://doi.org/10.1016/J.NEUBIOREV.2012.05.010

Pfeiffer, B., & Foster, D. (2013). Hippocampal place-cell sequences depict future

paths to remembered goals. *Nature*, *497*(7447), 74–79.

https://doi.org/10.1038/nature12112

Ritter, S., Wang, J., Kurth-Nelson, Z., & Botvinick, M. (2018). Episodic Control as

Meta-Reinforcement Learning. *BioRxiv*, 360537.

https://doi.org/10.1101/360537

Salakhutdinov, R., Tenenbaum, J. B., & Torralba, A. (2013). Learning with

hierarchical-deep models. *IEEE Transactions on Pattern Analysis and Machine*

*Intelligence*, *35*(8), 1958–1971. https://doi.org/10.1109/TPAMI.2012.269

Sanborn, A. N., Griffiths, T. L., & Navarro, D. J. (2010). Rational Approximations to

Rational Models: Alternative Algorithms for Category Learning. *Psychological*

*Review*, *117*(4), 1144–1167. https://doi.org/10.1037/a0020511

Schmidhuber, J. (2006). Developmental robotics, optimal artificial curiosity,

creativity, music, and the fine arts. *Connection Science*, *18*(2), 173–187.

https://doi.org/10.1080/09540090600768658

Schwartenbeck, P., FitzGerald, T., Mathys, C., Dolan, R., & Friston, K. (2015). The

Dopaminergic Midbrain Encodes the Expected Certainty about Desired

Outcomes. *Cerebral Cortex*, *25*(10), 3434–3445.

https://doi.org/10.1093/cercor/bhu159

Schwartenbeck, P., & Friston, K. (2016). Computational Phenotyping in Psychiatry: A

Worked Example. *ENeuro*, *3*(4), ENEURO.0049-0016.2016.

https://doi.org/10.1523/ENEURO.0049-16.2016

Sharot, T. (2011). The optimism bias. *Current Biology*, *21*(23), R941–R945.

https://doi.org/10.1016/J.CUB.2011.10.030

Stachenfeld, K. L., Botvinick, M. M., & Gershman, S. J. (2016). The hippocampus as a

predictive map.

Stickgold, R., Hobson, J., Fosse, R., & Fosse, M. (2001). Sleep, learning, and dreams:

off-line memory reprocessing. *Science*, *294*(5544), 1052–1057.

https://doi.org/10.1126/science.1063530

Tervo, D. G. R., Tenenbaum, J. B., & Gershman, S. J. (2016). Toward the neural

implementation of structure learning. *Current Opinion in Neurobiology*, *37*, 99–

105. https://doi.org/10.1016/J.CONB.2016.01.014

Tononi, G., & Cirelli, C. (2014). Sleep and the Price of Plasticity: From Synaptic and

Cellular Homeostasis to Memory Consolidation and Integration. *Neuron*, *81*(1),

12–34. https://doi.org/10.1016/J.NEURON.2013.12.025

Walker, M., & Stickgold, R. (2010). Overnight alchemy: sleep-dependent memory

evolution. *Nature Reviews. Neuroscience*, *11*(3), 218; author reply 218.

https://doi.org/10.1038/nrn2762-c1

Wang, J X, Kurth-Nelson, Z., Tirumala, D., Soyer, H., Leibo, J. Z., Munos, R., …

Botvinick, M. (2016). Learning to reinforcement learn. *ArXiv:1611.05763*.

Wang, Jane X., Kurth-Nelson, Z., Kumaran, D., Tirumala, D., Soyer, H., Leibo, J. Z., …

Botvinick, M. (2018). Prefrontal cortex as a meta-reinforcement learning

system. *Nature Neuroscience*, *21*(6), 860–868.

https://doi.org/10.1038/s41593-018-0147-8

Ward, T. B. (1994). Structured Imagination: the Role of Category Structure in

Exemplar Generation. *Cognitive Psychology*, *27*(1), 1–40.

https://doi.org/10.1006/cogp.1994.1010

Whittington, J. C. R., Muller, T. H., Mark, S., Barry, C., & Behrens, T. E. J. (2018).

Generalisation of structural knowledge in the hippocampal-entorhinal system.

Williams, J. J., & Lombrozo, T. (2010). The role of explanation in discovery and

generalization: evidence from category learning. *Cognitive Science*, *34*(5), 776–

806. https://doi.org/10.1111/j.1551-6709.2010.01113.x

Wipf, D. P., & Rao, B. D. (2007). An Empirical Bayesian Strategy for Solving the

Simultaneous Sparse Approximation Problem. *IEEE Transactions on Signal*

*Processing*, *55*(7), 3704–3716. https://doi.org/10.1109/TSP.2007.894265

Xu, F., & Tenenbaum, J. (2007a). Sensitivity to sampling in Bayesian word learning.

*Developmental Science*, *10*(3), 288–297. https://doi.org/10.1111/j.1467-

7687.2007.00590.x

Xu, F., & Tenenbaum, J. (2007b). Word Learning as Bayesian Inference. *Psychological*

*Review*, *114*(2), 245–272.

1427

1428