Boredom begets creativity or why predictive coding is not enough to explain intelligent behavior

Jaime Gomez-Ramirez*1 and Tommaso Costa†2

¹ The Hospital for Sick Children, Department of Neuroscience and Mental Health, University of Toronto, Bay St. 686, Toronto, (Canada) ² Koelliker Hospital, Department of Psychology, University of Turin, Via Verdi, 10, 10124 Turin (Italy)

Abstract

Here, we investigate whether systems that minimize prediction error e.g., predictive coding, can also show creativity, or on the contrary, prediction error minimization unqualifies for the design of systems that respond in creative ways to non recurrent problems. We argue that there is a key ingredient that has been overlooked by researchers and needs to be incorporated to build creative artificial systems. This ingredient is boredom. We propose a mathematical model based on the Black-Scholes equation which provides mechanistic insights into the interplay between pain (boredom) and pleasure (prediction) as the key drivers of behavior.

1 Introduction

The value in building artificial systems with optimal predictive power is beyond question. Robots in real world missions, without the capacity to build accurate predictions of the state of the world are unreliable and doomed to a short existence. In biological systems, the idea that organisms organize sensory data into an internal model of the outside world, goes back to the early days of experimental psychology. In Helmholtz's Handbook of Physiological Optics published in 1867, it is argued that the brain unconsciously adjusts itself to produce a coherent experience. According to this view, our perceptions of ex-

ternal objects are images or better said, symbols, that do not resemble the referenced objects. Helmholtz's insight had an enormous impact in a variety of fields, including cybernetics (Ashby, 2015), cognitive psychology (Neisser, 2014), machine learning (Neal and Hinton, 1998). Helmoltz's theory of perception as a process of probabilistic inference, in which sensory causes need to be inferred based upon changes of body states, has become a major tenet in computational neuroscience (Dayan and Abbott, 2002). A recent incarnation of this approach is the Helmholtz's machine postulated by Dayan, Hinton and Zemel (Dayan et al., 1995), (Dayan and Hinton, 1996). The brain is here conceptualized as a statistical

inference engine whose function is to infer the causes of sensory input.

Predictive coding is a form of differential coding where the encoded signal is the difference between the actual signal and its prediction. Predictive coding exploits the fact that under stationary and ergodic assumptions ¹, the value of one data point e.g., a pixel, regularly predicts the value of its nearest neighbors. Accordingly, the variance of the difference signal can be much smaller than that of the original signal, making differential coding a very efficient way to compress information (Shi and Sun, 1999). Elaborations of this same idea abound under different nomenclature and uses. For example, in digital signal processing, current signal values are estimated with Kalman Filters, a recursive algorithm that yield estimates of the current state variables, and update those estimates without assuming that the estimation errors are Gaussian (Kalman, 1960).

Predictive coding techniques aims at reducing redundancy for signal transmission efficiency and it is been proposed as a unifying mathematical framework for understanding information processing in the nervous system (Friston, 2010), (Huang and Rao, 2011). Specifically, predictive coding has been used to model spatial redundancy in the visual system (Srinivasan et al., 1982), temporal redundancy in the auditory system (Baldeweg, 2006) and the mirror neuron system (Kilner et al., 2007). Interestingly, this approach extends Barlow's redundancy reduction hypothesis, a theoretical model for sensory coding in the brain (Barlow, 1972). It ought to be

noted that Barlow himself has pointed out that the initial emphasis in the efficient coding theory in compressive coding needs to be amended, by thinking of neural representations not as efficient encoding of stimuli but as estimates of the probable truth of hypotheses about the environment (Barlow, 2001).

In the predictive coding framework, the workings of the brain encode Bayesian principles. Due in part to the ever increasing computational power of computers, Bayesian approaches alike to the Helmholtz's machine have become the workhorse for studying how the nervous system operates in situations of uncertainty (Rao and Ballard, 1999), (Knill and Pouget, 2004), (Friston, 2012). The main rationale is that the nervous system maintains internal probabilistic models informed by sensory information. The models are continuously updated in the light of their performance in predicting the upcoming suite of cues. In a general sense, predictive coding is a Bayesian approach to brain function in which the brain is conceived as a device trained to do error correction.

Predictive coding provides a mathematical description of optimal behavior but crucially, it does not prescribe how Bayesian optimal perception, sensorimotor integration or decisionmaking under uncertainty, materializes (Friston and Price, 2001), (Friston, 2009), (Friston, 2012). Agents that minimize surprise or the free energy in Friston's account are Bayes optimal, but this is not the same thing as behaving optimally. For example, in evolutionary terms, an optimal behavior would require to increment the organism's offspring and life span, maximizing hedonic pleasure or its reverse, reducing pain. Critics of predictive coding have often missed this point. They argue that if biological systems behave in the way that free energy min-

¹A signal is stationary when its defining probabilities are fixed in time. A signal is ergodic when can be constructed as a generalization of the law of large numbers (long term averages can be closely approximated by averages across the probability space)

imization prescribes, they would have a bland and uneventful existence, because they will inevitably seek the most predictable habitat, for example, a corner in a dark room, and they will stay there ad infinitum. This is being called the "dark-room problem" (Friston et al., 2012). Critics of predictive coding fail to recognize that the minimization free energy is an overarching principle and not a normative theory of biological behavior.

However, Friston's way out of the "dark-room problem" is unconvincing. The arguments goes as follow, probabilities are always conditional to the system's prior information, thus, a system equipped with a generative model (priors) that dislikes dark-rooms or similar dull environments will not be stuck in a corner minimizing prediction error, but will walk away in order to sample the external world according to its own priors. But where the priors come from and how they are shaped by the environment is never said. This is indeed the crux of the matter in Bayesian statistics. The translation of subjective prior beliefs into mathematically formulated prior distributions is an ill-defined problem (Gomez-Ramirez and Sanz, 2013). And yet, the minimization of surprise is a sufficient condition for keeping the system within an admissible set of states. A bacterium, a cockroach, a bird and a human being all have in common that in order to persevere in their actual forms, they must limit their possible physiological states, that is, organisms constrain their phenotype in order to resist disorder. Friston goes even further to claim that the physiology of biological systems can be reduced almost entirely to their homeostasis (Friston). Homeostasis is the control mechanism in charge of keeping the organism's internal conditions stable and within bounds. Survival depends on the organism's capacity to maintain its physiology within an optimal homeostatic range (Damasio and Carvalho, 2013).

Here is the conundrum that this paper addresses. On the one hand, free energy, surprise or surprisal ² minimization is conducive to achieving the homeostatic balance necessary for the organism's survival and well-being and on the other hand, surprise minimization can not possibly be the unique modus operandi of biological systems. Organisms that minimize prediction error would never engage in exploration, risk-taking or creativity, for the simple reason that these behaviors might increase the prediction error. In consequence, surprise or free energy can not be used as the unique necessary factor to explain choices under uncertainty conditions. We argue that the actual quantity that is maximized is the difference between prediction error and boredom.

The crucial intuition behind our model is strikingly simple. A system that minimizes prediction error is not only attentive to homeostasis and the vital maintenance functions of the body, but it also maximizes pleasure. For example, the reward effect in the appreciation of aesthetic work might come from the transition from a state of uncertainty to a state of increased predictability (Cruys and Wagemans, 2011). However, this is until the signal error becomes stationary, or in the art work example, the art work has not anymore the potential of surprising us, in that case boredom kicks in, reducing the overall value of the subjective experience.

Boredom is an aversive (negative valence) emotion. Thus, boredom creates the conditions to start exploring new hypothesis by sampling

²See the Appendix for the technical definition of surprisal and notes on predictive coding and free energy minimization

the environment in new and creative ways, or put in other words, boredom begets creativity. Until very recently, the function of boredom has been considered of little of no interest for understanding human functioning. This situation is rapidly changing, recent studies in human psychology shows that the experience of boredom might be accompanied by stress and increases levels of arousal to ready the person for alternatives (Posner et al., 2009) (Bench and Lench, 2013).

We are only just starting to understand the physiological signatures of boredom. Boredom compared with sadness shows rising heart rate, decreased skin conductance level, and increased cortisol levels (Merrifield and Danckert, 2014). Boring environments can generate stress, impulsivity, lowered levels of positive affect and risky behavior. Furthermore, in people with addiction, episodes of boredom are one of the most common predictors of relapse or risky behavior (Blaszczynski et al., 1990).

In the next section (2) we introduce a mathematical model that extends and complement predictive coding. In essence, predictive coding claims that a sufficient account of biological behavior is surprisal (entropy) minimization. We argue, on the contrary, that surprisal minimization in any of its equivalent forms such as free energy minimization and marginal likelihood maximization, is not a sufficient but a necessary explanans of biological behavior. Section 3 discusses the simulations of the model to help have an intuitive grasping of the mathematical model based on the Black-Scholes equation of option pricing. Section 4 provides a discussion about the limitations of maximum likelihood methods in relation with the previous results. An appendix with technical definitions of some of the concepts employed is included at the end of the paper

2 Methods

In this section we build a mathematical model to explain intelligent (biological or technical) behavior as the maximization of the subjective experience. The subjective experience consists on two terms with opposed valence, prediction and boredom. Prediction is a positive or hedonic state and boredom is a negative emotional state. In short, what organisms do is to maximize subjective experience, and in order to achieve that objective they tend to minimize surprise as predictive coding correctly claims, while at the same time diminishing boredom, a negative emotion that arises during monotonous tasks or in environments with low entropy. The model thus, extends the prediction error minimization by incorporating the boredom component in the utility function.

We start by defining the utility function that agents maximize. The rationale behind this is that organisms maximize subjective experience by making prediction pleasure as large as possible while keeping boredom to tolerable levels ³. A utility function is a mathematical description of subjective value that is constructed from choices under incomplete information conditions. We postulate that value-based decision do not only maximize prediction, rather agents maximize the difference between prediction and boredom. In this view, organisms do not exclusively operate in prediction mode, sooner or later, depending on the intrinsic agent's motivations and how they match with the environ-

³Note that prediction pleasure is the inverse of prediction error, therefore maximize prediction pleasure is the same as minimize prediction error.

ment, the marginal utility of prediction will decrease and the organism will switch to exploration mode, that is, the organism will become less concerned with predicting its current state, and they will be more prone to visit surprising states that overall increase its well being as encoded by the experience value. The utility function is defined as

$$v = p - k \tag{1}$$

where v is the subjective experience and p and b represent prediction and boredom, respectively. Equation 1 simply states that the subjective experience has two components with opposed valences, prediction pleasure and boredom. It seems clear that the larger the prediction pleasure (p) the greater the value of the subjective experience limited by the boredom (k) that prediction can bring in. When the prediction pleasure is greater than the boredom the subjective experience is overall positive or pleasant, on the contrary, when the boredom exceeds the prediction pleasure, experience is overall negative or painful.

The instantaneous subjective experience v_t is calculated as the difference between the instantaneous pleasure p_t and the negative pain b_t , which in our model is assumed to be constant $(b_t = k)$. The boredom constant k represents the agent's disposition to get bored and is thus, an inherent property of the system or causa sui. Prediction pleasure, on the other hand, is directly calculated from the prediction error. Prediction pleasure at time t, p_t , is the reciprocal of prediction error, s_t , that is, $p_t = \frac{1}{s_t}$. Accordingly, the value of the experience at time t is the difference between the prediction pleasure at t minus the

boredom component.

$$v_t = \frac{1}{s_t} - k = p_t - k \tag{2}$$

We need now to be more precise in the formulation of the terms included in equation 2. A reasonable assumption is that the prediction error describes a Brownian geometric model. The random variable prediction pleasure is the inverse of the prediction error and describes a generalized Wiener process. Under this assumption, the prediction pleasure p, in the limit as $\Delta t \to 0$, can be modeled as the following stochastic process,

$$dp = \mu p dt + \sigma s dz$$
$$\frac{dp}{p} = \mu dt + \sigma dz$$

where p is a random variable that represents the prediction pleasure, μ is the drift or the mean change per unit time, σ the variance per unit time and dz is a Wiener process with zero drift and 1.0 variance rate. Since the drift is equal to zero, the expected value of z is zero, that is, at any future time z is expected to be equal to its current value. The variance rate of 1.0 means that the variance of the change in z in a time interval of length T equals i.e., the variance rate grows proportionally to the maturity time T. The variable μ can be seen as the expected percentage gain/loss of prediction pleasure. For example, $\mu = 0.1$ means that prediction pleasure is expected to increment by a 10%. The variable σ is the volatility of the prediction pleasure. It is expected that σ in a world with high entropy will be larger than in a world with low entropy, ceteris paribus. For example, a surprising world with a large number of objects and events that are hard to predict will yield a large σ while a predictable environment, for example, an empty room will yield a low value of σ .

We are interested in quantifying the subjective experience as a function of the underlying prediction pleasure and time. To model subjective experience based on the underlying prediction pleasure and time, we borrow from the noted Black-Scholes model (Black and Scholes, 1973) used in mathematical finance for option pricing. In a seemingly way as an option price is a derivative of a stock price, a subjective experience value is referred to the underlying prediction pleasure at a given time t within a time horizon T, t < T. The Black-Scholes model will thus, help us establishing a working analytical framework to study the interplay between prediction and boredom.

In the rest of the section we derive the Black–Scholes equation from the Itô lemma (Ito) ⁴. Those not interested in the steps previous to the obtention of the model can directly jump to and the Results section having in mind equation 13.

2.1 Derivation of the Black-Scholes model

A random variable x follows a Itô process if

$$x = a(x, t)dt + b(x, t)dz$$

where dz is a Wiener process with a drift rate a and a variance rate b^2 both functions of x and t. The Itô lemma shows that a function f of x and t follows the stochastic process described in equation 3. The demonstration can be found elsewhere (Shreve).

$$df = \left(\frac{\partial f}{\partial x}a + \frac{\partial f}{\partial t} + \frac{1}{2}\frac{\partial^2 f}{\partial x^2}b^2\right)dt + \frac{\partial f}{\partial x}bdz \quad (3)$$

We can now relate equation 3 with the utility function defined in equation 1 in which the function v represents the subjective experience which is contingent on the prediction pleasure p. Then, according to the Itô lemma, the stochastic process of a function of p and t is simply obtained by substituting the drift rate $a = \mu p$ and the standard deviation rate $b = \sigma p$ into equation 3, resulting

$$df = \left(\frac{\partial f}{\partial p}\mu p + \frac{\partial f}{\partial t} + \frac{1}{2}\frac{\partial^2 f}{\partial p^2}(\sigma p)^2\right)dt + \frac{\partial f}{\partial p}\sigma p dz$$
(4)

Note that f also follows a Itô process with a drift rate of

$$\frac{\partial f}{\partial p}\mu p + \frac{\partial f}{\partial t} + \frac{1}{2}\frac{\partial^2 f}{\partial p^2}(\sigma p)^2 \tag{5}$$

and a variance rate

$$\left(\frac{\partial f}{\partial p}\right)^2 (\sigma p)^2 \tag{6}$$

It is possible to use the Itô lemma to characterize, for example, the process $\ln p$, where p is the prediction pleasure

$$f = \ln p \tag{7}$$

From equation 4 we obtain a generalized Wiener process with constant drift $\mu - \frac{\sigma^2}{2}$ and constant variance σ^2

$$df = \left(\mu - \frac{\sigma^2}{2}\right)dt + \sigma dz$$

The change in $\ln p$ between instant time 0 and final time T is therefore normally distributed with mean $(\mu - \frac{\sigma^2}{2})T$ and variance $\sigma^2 T$

$$\ln p_T - \ln p_0 \sim N\left(\left(\mu - \frac{\sigma^2}{2}\right)T, \sigma^2 T\right)$$

$$\ln p_T \sim N\left(\ln p_0 + \left(\mu - \frac{\sigma^2}{2}\right)T, \sigma^2 T\right)$$
(8)

 $^{^4}$ Black-Scholes can also be derived from a bionamial tree, see pages 298-300 in (Hull, 2011)

According to equation 8 the random variable $\ln p$ is normally distributed, therefore the variable prediction pleasure p follows a lognormal distribution.

The lognormal property of p can be used to study the probability distribution of the rate r of the prediction pleasure percentage earned/loss between two instants. The relationship between the prediction pleasure between times 0 and t = T is given by the equation

$$p_t = p_0 e^{rt}$$

Solving for r we have

$$r = \frac{1}{t} \ln \frac{p_t}{p_0}$$

and from Equation 8

$$r \sim N\left(\mu - \frac{\sigma^2}{2}, \frac{\sigma^2}{T}\right)$$
 (9)

The discount factor r can be understood as a prediction rate, which in essence, represents how much structure there is in the outside world. For example, in an external world in which information can not be compressed at all, r will be zero because a structure-less world entirely lacks predictability. In the other extreme of the spectrum, a very predictable world will have a large value of r. The prediction rate r can thus, be seen as a proxy for the structure of the outside world. The larger the prediction rate r, the more structure there is in the world to be discover by an agent equipped with the proper perceptual, motoric and cognitive capabilities.

Consider now that we are interested in studying the behavior of a system with a boredom constant k over a period of time T. The expected experience value at time t (v_t) is its expected

value at time T discounted at the rate r.

$$v_t = e^{-r(T-t)} \hat{E}(p_t - k)$$

= $e^{-r(T-t)} \hat{E}(p_t) - ke^{-r(T-t)}$ (10)

The value of the subjective experience at time t (t < T), v_t , is thus, equal to the expected prediction pleasure minus the boredom at the expiration time T, discounted at a discount rate r. Substituting equation 2 into equation 10 gives

$$v_t = p_t - ke^{-r(T-t)} \tag{11}$$

If the expiration time T is very far in the future, then the value of the subjective experience will be very similar to the prediction pleasure, on the other hand, if the expiration date is near, $(T-t\sim 0)$, the subjective experience is equal to prediction pleasure minus the boredom constant.

Equation 14 assumes that both prediction and boredom mode are equally likely. However, a more realistic model will weight the prediction and boredom terms by their respective probabilities. We borrow from the Black-Scholes model to define the subjective experience relative to the prediction pleasure constrained by the boredom component. The Black-Scholes formula to calculate the price of a call option (buying) for an underlying stock price s, strike price s, maturity s and risk free interest rate s is

$$c(s_t, k, t, \sigma, r, T) = s_t N(d_1) - k e^{-r(T-t)} N(d_2)$$
(12)

where s_t is the price of the underlying stock at time t defined as a generalized Wiener process, k is the strike price of the option, r is the constant riskless used to discount the value of the option back to time t from the maturity time T. The terms $N(d_1)$ and $N(d_2)$ are cumulative standard normal distributions, $N(d_i) = P(x > d_i)$. In particular, $N(d_2)$ is the probability that the option will be exercised, this will occur when the strike price k is only paid if the option is in the money. The interpretation of $N(d_1)$ is less straightforward but simplifying, it represents the probability that the stock price is less in value than the strike price, which is counted as zero in the calculus of the option price. For a more in depth discussion on the Black-Scholes model, the reader might want to consult the seminal paper (Black and Scholes, 1973) and two excellent textbooks (Hull, 2005) and (Duffie, 2001).

In a call option (equation 20), the buyer will be interested in exercise the option at time T, that is, buy the underlying stock, only if "is in the money", that is, $s_t N(d_1) > ke^{-r(T-t)} N(d_2)$. The discount factor $e^{-r(T-t)}$ reflects the need to take into account how much will cost to the buyer to borrow the money at the current time t in order to exercised the option.

Now that we have showed how to derive the Black-Scholes model for option price, we can go further with our analogy and quantify the experience value. The subjective experience is a function of the underlying prediction pleasure and boredom. Continuing with the financial analogy, in the Black-Scholes option pricing model (equation 19), the option is exercised only when the payoff is positive, in our model, on the other hand, the subjective experience is always "exercised". This means that the experience is what it is, positive when the prediction component is larger than the boredom term and negative the boredom exceeds the prediction pleasure.

Finally, we can define the value of the experience as the difference between the prediction and boredom discounted and weighted by the probability of being in each mode,

$$v_t = p_t N(d_1) - k e^{-r(T-t)} N(d_2)$$
 (13)

where the first term in the right side of equation 13 represents prediction pleasure factored by the probability of being in predictive mode, $N(d_1)$, and the second term quantifies the pain trigger by a boring experience in a world with with complexity r discounted a time t and factored by the probability of being in boredom mode, $N(d_2)$. $N(d_1)$ and $N(d_2)$ are cumulative probability distribution functions for the variables d_1 and d_2 defined as

$$d_{1} = \frac{\log \frac{p_{t}}{k} + (r_{t} + \frac{\sigma^{2}}{2})(T - t)}{\sigma \sqrt{T - t}}$$

$$d_{2} = \frac{\log \frac{k}{p_{t}} + (r_{t} - \frac{\sigma^{2}}{2})(T - t)}{\sigma \sqrt{T - t}}$$
(14)

A simple intuitive understanding of equation 13 comes from realizing that the agent transitions between two dynamic regimes -prediction and boredom- and the probability of being in prediction mode, that is, having more pleasure than pain is given by the probability $N(d_1)$ and the probability of being in boredom mode $N(d_2)$. The variables d_1 and d_2 are identical except for two things, i) the first term in the numerator is $\frac{\log s_t}{k}$ in d_1 and its inverse, $\frac{\log k}{s_t}$, in d_2 and ii) when prediction pleasure is equal to the boredom constant, s=k, the probability of being in prediction mode increments with the variability and decrements by the same amount in the boredom case.

3 Results

The underlying assumption for our model of prediction pleasure inspired in the Black-Scholes model is that both the Markov and the Martingale in stock price change also hold for prediction error. For that we need to assume that the prediction error is a stochastic process with

no memory, that is, the conditional probability distribution of the future states only depends on the current state and is therefore independent of any previous state (Markov property) and that knowledge of the past will be of no use in better predicting the future (Martingale property). These assumptions are compatible with the free energy principle, which is intended to explain biological systems behavior in changing a environment, under ergodic assumptions. Crucially, the ergodic assumption is what allows the system to minimize sensory entropy by means of surprise minimization at all times (Friston et al., 2010). Intuitively, the ergodic theorem states that for a random variable, in the long run, the time average is equal to the space average (Birkhoff, 1931).

We run simulations of the model described in equation 13 in different scenarios, according to the different setting of the four parameters of the model, namely the initial prediction pleasure (p_0) , the boredom constant (k), the expected rate of variation of the prediction pleasure (μ) and the variance of the prediction pleasure (σ) . The parameters prediction pleasure (p_0) and boredom constant (k) can be seen as the priors. For example, all things being equal, an agent with a large ratio k/p_0 will likely have a predominantly boredom experience compared to another agent with a large p_0/k which, on the contrary, will likely have a overall positive experience. In addition to the bias or predisposition of the agent to get bored, the expected rate of return r represents the environment's complexity and is directly specified by parameters (μ) and σ (equation 9). Remind that the parameter μ is the expected increase in the prediction pleasure at the maturity time and σ is the variability.

Figure 1 displays an agent's behavior under different agent-environment couplings, specified by the expected rate of return $\hat{r} = \mu - \frac{\sigma^2}{2}$. A pos-

itive value of \hat{r} denotes that it is likely that the agent will predict the world consistently. Thus, for two agents, a_1 and a_2 with $\hat{r_1} = \mu_1 - \frac{\sigma_1^2}{2}$ and $\hat{r_2} = \mu_2 - \frac{\sigma_2^2}{2}$ and $\hat{r_1} > \hat{r_2}$ we expect that agent a_1 will have larger prediction pleasure than agent a_2 , all things being equal. If $r_1 = r_2 = 0$ we are agnostic about the predictive power of both agents in their respective environments.

To get a grasping of the workings of the model and to show that the model has the right general properties, we consider what happens when some of its parameters take extreme values in equation 14.

If prediction pleasure is very large compared to boredom, $p_t >> k$, d_1 will have a very large value and d_2 will be very little, therefore $N(d_1) \simeq 1$ and $N(d_2) \simeq 0$. In this situation, the overall experience will be positive. On the contrary, when the ratio between $p_t/k \sim 0$ the overall experience will be negative or dominated by boredom. The rationale behind this is that if the world is very predictable, it is very likely that the agent will experience prediction pleasure or boredom depending on the own agent's bias specified by its preference to predict measured by the initial prediction pleasure p_0 or to get bored quantified with the k constant.

Figure 1 shows the simulation of the model when we are agnostic about the capacity of the agent to capture the structure of the world. We codify this case with $\mu = \frac{\sigma^2}{2}$, r = 0. When the agent does not have any particular predisposition of being in prediction or boredom mode $(p_0 = k)$ (figure 1 a), prediction pleasure decays and boredom starts to rise after a sufficient amount of time. If the agent has a predisposition to get bored $(k = 10p_0)$ (figure 1 b), prediction pleasure decays at much faster rate and boredom rises faster and earlier than in the previous case

than in the previous case. When the agent has a predisposition to enjoy prediction as opposed to get bored $(p_0 = 10k)$ (figure 1 c), both prediction pleasure and boredom remain stable over time.

Figure 2 shows the simulation of the model when we are optimistic about the capacity of the agent to capture the structure of the world. We codify this case with $\mu > \frac{\sigma^2}{2}, \ r > 0$. Thus, there is a structure of the outside world and the agent is equipped with the perceptual, motoric and cognitive capacities to predict the sensorial input. When the agent does not have any particular predisposition of being in prediction or boredom mode $(p_0 = k)$ (figure 2 a), prediction remains stable and so does boredom but at the end boredom will rise. Although the agent is predicting the world and having prediction pleasure, being consistently successful at predicting the world has the side effect of getting bored reducing the overall experience value. If the agent has a predisposition to get bored $(k = 10p_0)$ (figure 2 b), the overall experience value will be markedly negative at the end of the period. When the agent has a predisposition to enjoy prediction as opposed to get bored $(p_0 = 10k)$ (figure 1 c), both prediction pleasure and boredom remain stable over time, keeping the the overall experience value at a constant positive value.

In both figures (r = 0 and r > 0) the boredom component after an initial period of stability ends up raising reducing the overall experience value. Only when the agent has a clear predisposition to predict boredom stays stable and the experience value is explained with the prediction error (figures 1 and 2 c). This result is in agreement with the intuition that agents with a predisposition to like to predict will look for a quiet corner to predict optimally. On the other hand, agents that do not show any particular predisposition to predict, after a period of prediction allowing them to get acquainted to the environment will inevitably start getting bored, diminishing the overall experience value, triggering a risk prone behavior e.g.,look for the way out of the dark room, to counter decrease in experience value motivated by the increase in boredom.



4 Discussion

In the predictive coding framework the brain tries to infer the causes of the body sensations based on a generative model of the world. This inverse problem is famously formalized by the Bayes rule. When incoming sensorial data fully agree with beliefs, an exhaustion of prediction occurs in which prediction error signal becomes stationary. Thus, the system reaches an equilibrium characterized by sampling data from the environment in such a way that the system is never surprised. The idea behind this model is that somewhere in the brain there is a decision signal that encodes hypothesis about the sensorial information that is being processed.

Figure 3 schematically shows the process of decision making implemented as reaction time signal for two different hypothesis. Each signal is defined by two parameters: the prior probability of he hypothesis and the latency time which can be model as the time derivative or growth rate at which the signal reaches the threshold. The motivation in using this toy model is twofold. First, to provide a comprehensible account of surprisal minimization in a simple but explanatory powerful scenario consisting of only two parameters and second, indicate the limitations of surprisal minimization according to the utility model defined in section 2.

Let us see with an example how we can infer the hidden causes of the sensory input using likelihood maximization. A camper is sitting in front of a bonfire in the woods. It is a chilly and windy day. He hears a noise whose source can not recognize. The camper has two hypothesis to explain the noise, i) the noise is just the breeze moving the leaves or ii) the noise is caused by a black bear approaching the camp. Let A be the breeze signal and B the bear signal. Initially,

since there are only a few bears in those woods and the it is particularly windy, the camper gives more weight to the hypothesis A -the noise is caused by the wind- than to hypothesis B -it is a bear. Formally, the ratio of the logarithm of the likelihood that evidence E is caused by A versus B is bigger than 1.

$$\frac{\log p(E|A)}{\log p(E|B)} > 1 \tag{15}$$

The course of action -stay if the breeze hypothesis is true or go if the bear hypothesis is true- is given by the likelihood, or as Figure 4 explains by the likelihood factored by the priors. This decision process can be seen as a handicapped race between the two hypothesis that compete to reach the necessary evidence that would trigger the action. The race is handicapped because the competing hypothesis start with different prior probabilities. For example, if there are very few bears in the area, the bear hypothesis will be rarely selected. The discontinuous red line in Figure 4 represents the threshold at the crossing point of the two distributions encoding the competing hypothesis. However, if we take into account that being attacked by a bear is very rare, the prior probability of breeze is larger than that of a bear and the threshold can be shifted to the right as in Figure 4 b achieving more sensible responses to the properties external world.

But let us imagine now that after a long uneventful period of time and the consequent boredom, the agent would like to take the risk of getting into the woods to explore the surrounding area. How can surprisal minimization or the analogous likelihood maximization explain this new behavior? By readjust the prior and/or time derivative of the likelihood in such a way that the bear hypothesis reaches the threshold before the breeze hypothesis the agent would leave the place to escape from the upcoming danger, rather than staying. But this is not the same as exploring. The likelihood maximization based decision process depicted in Figure 4 is unable to explain why the organism would engage in such a behavior. Even if we are as indulgent with the use of words as to equate exploring with escaping the model would not tell us why the organism reaches to the observed behavior. Crucially, the evaluation of the priors is outside the model. The crux of the matter is to realize that likelihood maximization in all its forms and denominations i.e., surprise minimization or free energy minimization, can not explain behaviors that are not dedicated to minimize prediction error.

The mathematical model proposed in section 2 extends predictive coding into an unifying and coherent approach. From an evolutionary perspective, subjective experience exists to facilitate learning of conditions responsible for homeostatic imbalances and of their corrective responses. Our model is in agreement with the well established fact that the brain deliberately randomizes reaction times bin order to have action variability (Carpenter, 1999). If we always react in the same way to common stimuli e.g., staying if the noise is caused by the breeze, life will be boring and there would be no incentive to explore, discover and wander. By randomizing the rate of rise of the signal towards the threshold it is possible to randomize choice or increasing its variability. There is an evolutionary advantage in doing surprising actions. For example, in a prey-predator game, both prey and the predator, will have a better change to succeed if they behave surprisingly rather than in predictable ways.

The tension between adaptation and satiation is taken care by the homeostatic control mechanism that keeps the organism's internal conditions within admissible bounds. The exhaustion of prediction disrupts the homeostatic balance, boredom leads to variety seeking to restore the homeostatic balance. This idea exists in popular parlance in the idiom "die of success", minimizing prediction error would make the system to seek for facile environments to predict, neglecting exploration and over valuing risk, which would make the system maladapted for prospering and survive in more complex or realistic environments. Our model provides a overarching principle for behavioral modeling, extending the predictive coding framework to a more explanatory framework. Biological systems do not just minimize free energy, rather free energy or surprise is one dependent variable, the other is boredom, and the interplay between both pleasure (prediction) and pain (boredom) defines the independent variable, subjective experience, which is the quantity that systems, all things being equal, maximize.

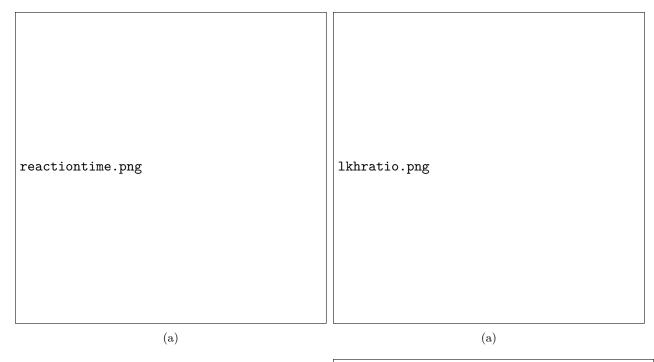


Figure 3: Figure a shows the handicapped race between the two hypothesis in a single trial. When we are certain that the generative model has created the observed data, the action is taken. There are two hypothesis -the sound is produced by the breeze or by a beareach with its corresponding action -stay or go. The competition between the two hypothesis is handicapped in the sense that the likelihood is weighted by the prior. Thus, for the bear hypothesis to reach the threshold before the breeze hypothesis does it, it would require that the likelihood of the former grows at a faster rate.

lkhratio-2.png

(b)

Figure 4: Figure a depicts the distribution of the responses of a neuron or neurons of interest in 14the auditory cortex encoding the stimulus. The x-axis represents the number of spikes that the neuron(s) fire per time unit. The intuition is that the larger the number of spikes, s the most likely that the cause of the noise being a bear. The probability of response E (stay) given that the cause was the breeze is p(E|A) and p(E|B) for the bear causing the response (go). If we want

References

- Ashby, W. R. (2015). An Introduction to Cybernetics. Martino Fine Books.
- Baldeweg, T. (2006). Repetition effects to sounds: evidence for predictive coding in the auditory system. *Trends in Cognitive Sciences*, 10(3):93–94.
- Barlow, H. (1972). Single units and sensation: A neuron doctrine for perceptual psychology? *Perception*, 1:371–394.
- Barlow, H. (2001). Redundancy reduction revisited. *Network (Bristol, England)*, 12(3):241–253.
- Barto, A., Mirolli, M., and Baldassarre, G. (2013). Novelty or Surprise? *Cognitive Science*, 4:907.
- Bench, S. W. and Lench, H. C. (2013). On the function of boredom. *Behavioral Sciences* (*Basel, Switzerland*), 3(3):459–472.
- Birkhoff, G. D. (1931). Proof of the ergodic theorem. *Proceedings of the National Academy of Sciences*, 17(12):656–660.
- Black, F. and Scholes, M. S. (1973). The pricing of options and corporate liabilities. *Journal of Political Economy*, 81(3):637–54.
- Blaszczynski, A., McCONAGHY, N., and Frankova, A. (1990). Boredom proneness in pathological gambling. *Psychological Reports*, 67(1):35–42.
- Carpenter, R. (1999). A neural mechanism that randomises behaviour. *Journal of Consciousness Studies*, 6(1):13–13.

- Cruys, S. V. d. and Wagemans, J. (2011). Putting reward in art: A tentative prediction error account of visual art. 2(9):1035–1062.
- Damasio, A. and Carvalho, G. B. (2013). The nature of feelings: evolutionary and neurobiological origins. *Nature Reviews Neuroscience*, 14(2):143–152.
- Dayan, P. and Abbott, L. (2002). Theoretical neuroscience: computational and mathematical modeling of neural systems. MIT Press.
- Dayan, P. and Hinton, G. E. (1996). Varieties of Helmholtz Machine. *Neural Networks*, 9(8):1385–1403.
- Dayan, P., Hinton, G. E., Neal, R. M., and Zemel, R. S. (1995). The Helmholtz machine. *Neural Computation*, 7(5):889–904.
- Duffie, D. (2001). Dynamic Asset Pricing Theory, Third Edition. Princeton University Press, third edition edition.
- Friston, K. The free-energy principle: a unified brain theory? 11(2):127–138.
- Friston, K. (2009). The free-energy principle: a rough guide to the brain? *Trends in Cognitive Sciences*, 13(7):293–301.
- Friston, K. (2010). The free-energy principle: a unified brain theory? *Nature Reviews Neuroscience*, 11:127–138.
- Friston, K. (2012). The history of the future of the Bayesian brain. *Neuroimage*, 62-248(2):1230–1233.
- Friston, K., Schwartenbeck, P., FitzGerald, T., Moutoussis, M., Behrens, T., and Dolan, R. J.

- (2013). The anatomy of choice: active inference and agency. Frontiers in Human Neuroscience, 7.
- Friston, K., Thornton, C., and Clark, A. (2012). Free-energy minimization and the dark-room problem. *Perception Science*, 3:130.
- Friston, K. J., Daunizeau, J., Kilner, J., and Kiebel, S. J. (2010). Action and behavior: a free-energy formulation. *Biological Cybernet*ics, 102(3):227–260.
- Friston, K. J. and Price, C. J. (2001). Dynamic representations and generative models of brain function. *Brain research bulletin*, 54(3):275–85.
- Gomez-Ramirez, J. (2013). Don't blame the economists it is an inverse problem! *European Journal of Futures Research*, 1(1):1–7.
- Gomez-Ramirez, J. and Sanz, R. (2013). On the limitations of standard statistical modeling in biological systems: A full bayesian approach for biology. *Progress in Biophysics and Molecular Biology*.
- Huang, Y. and Rao, R. P. N. (2011). Predictive coding. Wiley Interdisciplinary Reviews. Cognitive Science, 2(5):580-593.
- Hull, C. J. (2011). Options, Futures, and Other Derivatives. Prentice Hall, 9 edition edition.
- Hull, J. C. (2005). Options, Futures and Other Derivatives. Prentice Hall, 6 edition edition.
- Ito, K. On Stochastic Differential Equations.
 American Mathematical Society.
- Kalman, R. (1960). A New Approach to Linear Filtering and Prediction Problems. *Transac-*

- tions of the ASME Journal of Basic Engineering, 0(82 (Series D)):35–45.
- Kilner, J. M., Friston, K. J., and Frith, C. D. (2007). Predictive coding: an account of the mirror neuron system. *Cognitive Processing*, 8(3):159–166.
- Knill, D. C. and Pouget, A. (2004). The Bayesian brain: the role of uncertainty in neural coding and computation. *Trends in Neurosciences*, 27(12):712–719.
- Merrifield, C. and Danckert, J. (2014). Characterizing the psychophysiological signature of boredom. *Experimental Brain Research*, 232(2):481–491.
- Neal, R. and Hinton, G. E. (1998). A View Of The Em Algorithm That Justifies Incremental, Sparse, And Other Variants. In *Learning* in *Graphical Models*, pages 355–368. Kluwer Academic Publishers.
- Neisser, U. (2014). Cognitive Psychology: Classic Edition. Psychology Press.
- Palm, G. (2012). Novelty, Information and Surprise. Springer Science & Business Media.
- Posner, J., Russell, J. A., Gerber, A., Gorman, D., Colibazzi, T., Yu, S., Wang, Z., Kangarlu, A., Zhu, H., and Peterson, B. S. (2009). The Neurophysiological Bases of Emotion: An fMRI Study of the Affective Circumplex Using Emotion-Denoting Words. *Human brain mapping*, 30(3):883–895.
- Rao, R. P. and Ballard, D. H. (1999). Predictive coding in the visual cortex: a functional interpretation of some extra-classical receptivefield effects. *Nature Neuroscience*, 2(1):79–87.

Shi, Y. Q. and Sun, H. (1999). Image and Video Compression for Multimedia Engineering: Fundamentals, Algorithms, and Standards. CRC Press, 1 edition edition.

Shreve, S. Stochastic Calculus for Finance II: Continuous-Time Models. Springer.

Srinivasan, M. V., Laughlin, S. B., and Dubs, A. (1982). Predictive coding: a fresh view of inhibition in the retina. Proceedings of the Royal Society of London. Series B, Biological Sciences, 216(1205):427–459.

Tribus, M. (1961). Thermostatics and Thermodynamics. D. Van Nostrand Company, 1st edition edition.

Appendix

4.1 Surprisal

The self information or surprisal associated with an outcome x is defined as

$$S = -\log p(x) \tag{16}$$

Surprisal represents the surprise of seeing the outcome x. The more likely the outcome is, the less surprising is and therefore lower surprisal value Tribus (1961), Barto et al. (2013). For example, in a fair dice the surprisal associated with having a 4 is $-\log p(x=4) = -\log \frac{1}{6} = 1.79$ bits. Having any other outcome is less surprising and therefore the surprisal is lower, $-\log p(xneq4) = -\log(\frac{5}{6}) = 0.18$.

However, if we follow the Bayesian interpretation of probability, surprisal is the log likelihood of outcomes x, marginalized over their causes or model m

$$S = -\log p(x|m) \tag{17}$$

The Bayesian approach is necessary if we want to explain why if, for example, playing the state lottery, the winner sequence of numbers 1,2,3,4,5 seems more surprising than the sequence 45,11,23,15,67 despite the fact that the probability of both outcomes $x_a = \{1, 2, 3, 4, 5\}$ and $x_b = \{45, 11, 23, 15, 67\}$ are identical Palm (2012). When the generative model m is incorporated into the equation of surprisal (Equation 17) the probability of seeing x_a can be considered larger than the probability of seeing x_b if the model m assumes that the most likely outcome needs more bits to be described, that is, the model m may assume incompressibility of the outcome. It follows then that to evaluate surprise it is necessary to marginalize over the hidden causes of outcomes, that is to say, we need to calculate the likelihood of the outcomes given the causes or p(x|m). Knowing the causes of observations is obviously not always possible (Gomez-Ramirez, 2013).

4.2 Free energy minimization and predictive coding

The Helmoltz machine addresses this problem by using variational free energy as a proxy, more specifically, an upper bound on surprise for surprise. Under this view, an agent that minimizes the free energy is also minimizing surprise and most importantly maximizing the model evidence, that is, the likelihood of outcomes Dayan et al. (1995). The rationale is that although agents might not know the causes of their observations they can infer them by minimizing the free energy Friston et al. (2013). Predictive Coding is an unifying framework to understand redundancy reduction and efficient coding (economy of thought) in the nervous system. By transmitting only the unpredicted parts

of the messages predictive coding allows to reduce redundancy. According to predictive coding, agents try to minimize the dispersion of the sensory state, that is to say, the agent samples the world to minimize its surprise or surprisal which is defined as, $-\log p(s|m)$, where s represents the probability of sensory outcome given a generative model, m. Since the agent can not possibly know the sensory outcome before it actually occurs, it is not possible to directly minimize this quantity. However, what we can do is to minimize an upper bound of the surprisal, namely, the free energy F. This bound is created by simply adding a cross entropy or Kullback-Leibler divergence which is always non negative. Accordingly, we can indirectly minimize surprise by minimizing the free energy,

$$F(s, \theta, \phi) = -\log p(s|\theta) + D_{KL}(Q(\phi, s), P(\theta, s))$$
(18)

where F is the free energy, $H = -\log p(s|\theta)$ is the surprisal or the log probability of generating a particular sample, s, from a model with parameters θ and $D_{KL}(Q, P)$ is the divergence between the recognition distribution Q and the generative distribution, P. Note that the recognition and the generative distributions have their own parameters ϕ and θ , respectively, which are optimized at the same time to maximize the overall fit function, F. The important point to keep in mind here is that the free energy F is minimized by maximizing the marginal likelihood, $p(s|\theta)$, or identically said, minimizing the entropy, $H = -\log p(s|\theta)$. In essence, Equation 18 defines a Bayesian evidence model in which minimizing the free energy corresponds to maximizing the likelihood or evidence upon the agent's model of the world.

Black Scholes formula and option price

The most important result in the valuation of options is due to Black, Scholes and Merton (Black and Scholes, 1973). An option is a security giving the right to sell or buy an asset within a specified period of time. The Black-Scholes formula calculates the price for both the call option (buying) and the put option (selling) at a maturity T with strike price. An "European option" gives the right to buy the asset for the striking price, thus, if the the asset's price at maturity is larger than the strike price the option is exercised. The price of a call option is therefore $max(s_T - k, 0)$, that is, the price for this option is the difference between the actual price and the strike price when $s_T - k > 0$ or 0 otherwise, because if the asset's price is less than the strike price $(s_T < k)$ we are not obligated to buy the asset. The Black-Scholes model for a call option is

$$c(s_t, k, t, \sigma, r, T) = s_t N(d_1) - k e^{-r(T-t)} N(d_2)$$
(19)

and for a put option is

$$p(s_t, k, t, \sigma, r, T) = ke^{-r(T-t)}N(-d_2) - s_tN(-d_1)$$
(20)

Assuming that the stock price changes follows a binomial distribution (ups and downs in value) we can derive the values of d_1 and d_2 as a binomial. For more details see about how these results are obtained, see(Hull, 2011).

$$d_1 = \frac{\log \frac{S_t}{K} + (r + \frac{\sigma^2}{2})(T - t)}{\sigma} \sqrt{T - t} \qquad (21)$$

and

$$d_2 = d_1 - \sigma\sqrt{T - t} \tag{22}$$

 $N(d_2)$ is the risk neutral probability of the outflow K that is the risk neutral probability that the option finish in the money. The interpretation of $N(d_1)$ is more complicated, see () for a comprehensible account see (Hull, 2005) and (Duffie, 2001). Form Equation 22 it is straightforward to see that for zero variability σ , $d_1 = d_2$, for large variability and time, then $N(d_2) \sim 0$.

We build on the analogy that subjective experience can be studied as a derivative or option of the prediction pleasure, that is, just as options price are calculated via the underlying stock price, it is possible establish approach to calculate subjective value referred to prediction pleasure. In this vein, given the distribution of the prediction pleasure P which consists on N samples $N=T/\Delta T$

$$V = P - B \tag{23}$$

The subjective experience V at time 0 is defined as

$$V_0 = P_0 N(d_1) - Be^{-r(T)} N(d_2)$$
 (24)

where P represents the prediction pleasure at each moment in time t = 0, B represents the propensity of the agent to get bored, r is the drift or how fast prediction pleasure decays over time and the term $N(d_2)$ is the cumulative standard normal distribution that yields the probability $N(d_2) = P(x > d_2)$. Both d_1 and d_2 have been adjusted to the needs of our problem. Based on the variable d_2 , which according to the Black-Scholes-Merton model is defined as,

$$d_2 = d_1 - \sigma \tag{25}$$

where,

$$d_1 = \frac{\log \frac{P_t}{B} + (r_t + \frac{\sigma^2}{2})T}{\sigma\sqrt{T}}$$
 (26)

One major difference between our model and the option pricing model is that the subjective experience is always what it is, while the option is only executed if P > B. It follows that d_1 and d_2 needs to be accordingly changed (Equation 14).