Expectancy-based rhythmic entrainment as continuous Bayesian inference 2 Jonathan Cannon March 5, 2021 4 Department of Brain and Cognitive Science, Massachusetts Institute of Technology, Cambridge, MA, USA 5 Tel.: +314-749-6902 jcan@mit.edu Abstract When presented with complex rhythmic auditory stimuli, humans are able to track underlying temporal structure (e.g., a "beat"), both covertly and with their movements. This capacity goes far beyond 10 that of a simple entrained oscillator, drawing on contextual and enculturated timing expectations and 11 adjusting rapidly to perturbations in event timing, phase, and tempo. Previous modeling work has 12 described how entrainment to rhythms may be shaped by event timing expectations, but sheds little 13 light on any underlying computational principles that could unify the phenomenon of expectation-based 14 entrainment with other brain processes. Inspired by the predictive processing framework, we propose 15 that the problem of rhythm tracking is naturally characterized as a problem of continuously estimat-16 ing an underlying phase and tempo based on precise event times and their correspondence to timing 17 expectations. We present two inference problems formalizing this insight: PIPPET (Phase Inference 18 from Point Process Event Timing) and PATIPPET (Phase and Tempo Inference). Variational solutions 19 to these inference problems resemble previous "Dynamic Attending" models of perceptual entrainment, 20 but introduce new terms representing the dynamics of uncertainty and the influence of expectations in 21 the absence of sensory events. These terms allow us to model multiple characteristics of covert and 22 motor human rhythm tracking not addressed by other models, including sensitivity of error corrections 23 to inter-event interval and perceived tempo changes induced by event omissions. We show that positing 24 these novel influences in human entrainment yields a range of testable behavioral predictions. Guided 25 by recent neurophysiological observations, we attempt to align the phase inference framework with a 26 specific brain implementation. We also explore the potential of this normative framework to guide the 27

- ²⁸ interpretation of experimental data and serve as building blocks for even richer predictive processing and
- ²⁹ active inference models of timing.
- 30 Keywords: Bayesian Inference, Active Inference, Timing, Rhythm, Entrainment

31 **Introduction**

The human brain is remarkably proficient at identifying and exploiting temporal structure in its environment, especially in the auditory domain. This phenomenon is most easily observed in the case of auditory stimuli with underlying periodicity: humans adeptly and often spontaneously synchronize their movements with such auditory rhythms [1], and human brain activity in auditory and motor regions aligns to auditory stimulus periodicity even in the absence of movement [2]. Both of these phenomena are cases of "entrainment" (sensorimotor and neural, respectively), where we define "entrainment" as in [3]: the temporal alignment of a biological or behavioral process with the regularities in an exogenously occurring stimulus.

A simple sinusoidal phase oscillator can entrain to a periodic stimulus; however, it is difficult to discuss the 39 flexible entrainment of human behavior and cognitive processes to variable and sometimes aperiodic patterns 40 such as speech without invoking the cognitive concept of "temporal expectation." Expectations for event 41 timing can be used to achieve a range of behavioral goals. They can help us hone our sensory detection, our 42 sensory discrimination, and our response time for behaviorally important stimuli at the anticipated time [4, 43 5, 6]. In some situations, temporal expectations attenuate neural responses [7], which may help to conserve 44 neural resources. And timing expectations bias our perception of time, allowing us to use prior experience 45 to supplement noisy sensory data as we make temporal judgments [8]. 46

Entrainment in humans involves an interplay of stimulus and temporal expectation [9]. Nowhere is 47 this clearer than in interaction with music, humankind's playground for auditory temporal expectation and 48 entrainment [10]. But the precise nature of this interplay is an open question. The framework of Dynamic 49 Attending Theory characterizes temporal expectancy as pulses of "attentional energy" issued by entrained 50 neural oscillators, and mathematical models based on these ideas describe bidirectional interactions between 51 temporal expectation and entrainment that reproduce aspects of human behavior and perception [11, 12]. But 52 although the behavior of these models may be satisfying in certain applications, the groundwork underlying 53 them is less so: key high-level concepts like the "attentional pulse" are difficult to define mechanistically or 54 computationally, so the implementations of these concepts in models remain impressionistic. 55

An alternative approach to modeling the role of expectations in the brain is the "predictive processing" framework [13]. This framework posits that the brain engages in a continuous process of inferring the hidden causes of sensory events based on a learned understanding of how those causes produce sensation. Unlike the

terms in Dynamic Attending Theory models, the terms in predictive processing models are directly linked to the formal inference problem being solved: the solution to the problem demands that certain quantities be computed, giving us reason to expect to find those quantities represented in the brain. In particular, "precision" or certainty plays a key role, determining how new sensory information is weighted relative to existing beliefs about the hidden causes.

Here, we apply the predictive processing approach to the process of expectancy-based entrainment by formalizing it as an inference problem: namely, the problem of inferring the state of the exogenous process giving rise to a series of events in time. We use the mathematical tool of point processes to formulate a model of precise event timing. We derive an optimal solution to the inference problem, which we hypothesize corresponds with the brain's mechanisms for entrainment. The resulting models resemble Dynamic Attending Theory models, but introduce two key novel elements:

Dynamically estimated phase uncertainty moderates the balance between top-down and bottom-up
 influences on estimated phase.

⁷² 2. Event expectations influence estimated phase even in the absence of actual events.

These elements allow them to reproduce aspects of human entrainment unaddressed by existing models,including:

Failure to track phase through excessive syncopation (events occurring at weakly expected times but
 omitted at strongly expected times).

2. Illusory contraction of intervals when expected events are omitted.

Near-linear corrections to phase after event timing perturbations, with larger (and even over-) corrections for stimulus trains with longer inter-onset intervals.

They are also significantly more flexible than Dynamic Attending Theory models in their descriptive power, allowing us to describe entrainment based on either periodic or aperiodic expectation patterns, and, as predictive processing models, they recast entrainment in a formal language that links it a the wide range of other cognitive phenomena.

In the next section, we formulate three versions of the problem of expectancy-based entrainment that are amenable to precise solutions, which we refer to collectively as the "phase inference framework." In the first, "Phase Inference from Point Process Event Timing" (PIPPET), a hidden phase variable advances steadily with added noise, and the observer is tasked with continuously inferring the phase based on the observation of events emitted probabilistically at certain phases with certain degrees of precision. In the second version,

"Phase And Tempo Inference from Point Process Event Timing" (PATIPPET), the rate of phase advance 89 (tempo) is also a dynamic variable with drift, and the solution simultaneously estimates phase, tempo, and 90 certainty about both. The third version (mPIPPET) generalizes the first two to incorporate the observation 91 of multiple types of events, each with distinct characteristic phases and precisions, into the inference process. 92 We present variational filtering equations that approximate perfect Bayesian solutions to these problems. 93 In the Results section, we simulate these filters, drawing on music as a rich source of intuitive examples 94 of entrainment informed by expectation. In doing so, we provide intuition into the range of behaviors of 95 these solutions, and show how novel features introduced by the normative framework reproduce key aspects 96 of human entrainment behavior that are not explained by other models. In the Discussion, we discuss the 97 potential contributions of PIPPET and PATIPPET to the analysis of experimental data, to richer and more 98 detailed models, and to our understanding of entrainment in the brain. 99

¹⁰⁰ 2 Mathematical framework

Predictive processing should be a natural modeling framework for understanding rhythmic expectation and 101 entrainment [14, 15, 16]. However, existing predictive coding models that operate in continuous time are 102 structured to perform inference based on continuous observation, characterizing prediction errors in terms 103 of deviation between a true level of input and a mean expected level [17, 18]. In other words, they describe 104 predictions about "what" rather than "when." They are therefore ill-suited to characterizing moment-by-105 moment errors in *timing* prediction, which are made sporadically and separated by intervals mostly devoid 106 of informative prediction error. This may be a fundamental shortcoming in modeling inference in the brain: 107 behavior and neurophysiology suggests that information about "when" is carried by its own distinctive 108 pathways and represented separately from "what," both in perceptual and motor tasks [19, 6, 10]. Bayesian 109 methods have been applied to describe inferences about timing in the brain [20, 21, 22], but in these cases 110 the problem the brain solves has been formulated as discrete inferences about consecutive intervals rather 111 than a continuous inference process. 112

Here, we use event timing to inform a continuous variational inference process by first creating a generative model describing the probabilistic generation of precisely timed events and then variationally inverting that model. To model event generation, we use the mathematical tool of point processes.

¹¹⁶ 2.1 Phase Inference from Point Process Event Timing (PIPPET)

PIPPET is the problem of dynamically estimating a hidden noisy phase variable based on the timing of events generated as a point process whose rate is modulated as a specific function of phase. The generative

model consists of a phase $\phi \in \mathbb{R}$ that advances as a drift-diffusion process:

$$d\phi = dt + \sigma dW_t \tag{1}$$

and an inhomogeneous point process that generates events with probability $\lambda(\phi)$. This function is known to the observer. We will refer to $\lambda(\phi)$ as an "expectation template" because it describes the temporal structure of the observer's event expectations, though it can also be understood as a hazard rate for events. To achieve both analytical tractability and flexible descriptive power, we assume that $\lambda(\phi)$ is a sum of a constant λ_0 and a set of scaled Gaussian peaks indexed by i = 1, 2, ... etc. Each Gaussian peak i is centered at a mean phase ϕ_i with variance v_i and scale λ_i :

$$\lambda(\phi) = \lambda_0 + \sum_i \lambda_i \varphi(\phi | \phi_i, v_i) \tag{2}$$

where $\varphi(\cdot|m, v)$ denotes the pdf of a Gaussian distribution with mean m and variance v.

• Each Gaussian mean ϕ_i represents a phase at which an event is expected;

- λ_i represents the strength of that expectation;
- and v_i^{-1} is the temporal precision of that expectation.

• $\lambda_0 > 0$ represents the rate of events being generated as part of a uniform noise background unrelated to phase.

The point process with rate described by (2) can be understood as a sum of independent point processes i, one for each expectation peak and one for the uniform background process with rate λ_0 , whose events are indistinguishable. The mathematics of updating a phase estimate at an event can be understood to involve a causal inference on which of these processes caused each event.

 $\lambda(\phi)dt$ is the likelihood function over ϕ associated with the occurrence of an event, so $\lambda(\phi)$ is a rescaled likelihood function. See Figure 1A for illustration.

¹³⁸ Note that ϕ is assumed to be on the real line, not the circle. This design decision allows PIPPET to ¹³⁹ entrain to temporally patterned expectations with or without periodic structure by choosing a periodic or ¹⁴⁰ aperiodic expectation template λ .

Given a series of event times $\{t_n\}$ tallied by an event-counting function $N_t : \mathbb{R} \to \mathbb{Z}^{0+}$, an expectation template $\lambda(\phi)$, and a prior distribution $p_0(\phi)$ describing the distribution of phase at time t = 0, the observer's goal is to infer a posterior distribution $p_t(\phi) = p(\phi|N_{\tau < t})$ describing an estimate of phase ϕ at any time tbased on the event history up to t.

In [23], Snyder derives an exact PDE for the evolution of this posterior distribution over time. Following 145 the predictive processing ansatz of maintaining Gaussian posterior distributions (the Laplace assumption), 146 which provides both computational tractability and neurophysiological plausibility by reducing the repre-147 sentation of the posterior to a mean and a variance, we project the posterior onto a Gaussian at each dt148 time-step. We do this by moment-matching: we use Snyder's solution to determine the evolution of the mean 149 and variance of the posterior, and then replace the true posterior with a Gaussian with the same mean and 150 variance. This choice of Gaussian is the choice with minimum KL divergence from the true posterior [24], 151 and therefore also minimizes the free energy of the solution within the family of possible Gaussian posteriors 152 in accordance with the Free Energy Principle [25]. 153

The result of this derivation is a generalization of a Kalman-Bucy filter with Poisson observation noise. 154 Eden and Brown [26] have derived an explicit form for this filter, but it relies on a local approximation of 155 the rate function λ that hides some of the interesting effects of events expected at nearby time points. For 156 λ a mixture of Gaussians, we derive a filter directly from Snyder's solution in [23] that more accurately 157 approximates the optimal (Bayesian) solution. The derivation is presented in Appendix 6.2. 158

Solution: the PIPPET filter At any time t, let μ_t denote the mean and V_t denote the variance of the 159 Gaussian posterior. At each event time t, we let μ_t and V_t equal the left-hand limits of μ and V before 160 the event, and we write μ_{t+} and V_{t+} to denote their right-hand limit values after the event (μ and V are 161 left-continuous). Let dN_t denote the increment in the event-counting process at time t, which is either 0 or 162 1 with probability one. μ_t and V_t evolve according to the stochastic differential equation: 163

$$\begin{cases} d\mu = dt + (\hat{\mu} - \mu)(dN_t - \Lambda dt) \\ dV = \sigma^2 dt + (\hat{V} - V)(dN_t - \Lambda dt) \end{cases}$$
(3)

or, equivalently, they evolve between events according to the ODE: $\begin{cases} \dot{\mu} = & 1 - \Lambda(\hat{\mu} - \mu) \\ & \\ \dot{V} = & \sigma^2 - \Lambda(\hat{V} - V) \end{cases}$ and reset at

each event to $\mu_{t+} = \hat{\mu}$ and $V_{t+} = \hat{V}$, where we define

$$\hat{\mu} := \frac{\lambda_0}{\Lambda} \mu_t + \sum_{i=1,\cdots} \frac{\Lambda_i}{\Lambda} \hat{\mu}_i$$
$$\hat{V} := \frac{\lambda_0}{\Lambda} \left(V_t + (\mu_t - \mu_{t+})^2 \right) + \sum_{i=1,\cdots} \frac{\Lambda_i}{\Lambda} \left(\hat{V}_i + (\hat{\mu}_i - \mu_{t+})^2 \right)$$

(Note that in this formulation, μ_{t+} must be calculated before V_{t+} .)

$$\hat{\mu}_{i} := \frac{V_{t}^{-1} \mu_{t} + v_{i}^{-1} \phi_{i}}{V_{t}^{-1} + v_{i}^{-1}} \quad \text{and} \quad \hat{V}_{i} := \frac{1}{V_{t}^{-1} + v_{i}^{-1}}$$
$$\Lambda_{i} := \lambda_{i} \varphi(\mu_{t} | \phi_{i}, v_{i} + V_{t}) \quad \text{and} \quad \Lambda := \sum_{i} \Lambda_{i}$$

¹⁶⁴ These terms are illustrated in Figure 1. Intuitively,

- Λ (implicitly a function of μ_t and V_t) is the degree to which an event is anticipated at t while taking into account uncertainty about underlying phase, also known as the "subjective hazard rate". Λ_i is the degree to which an event is anticipated from peak i (the "conditional subjective hazard rate").
- At each event time t, $\lambda(\phi)$ serves as a (rescaled) likelihood function for phase, and the role of prior 168 is played by the phase distribution p_t , a Gaussian with mean μ_t and variance V_t . Each peak i of λ is 169 a possible "cause" of the event, as is the background event rate λ_0 . Each peak is associated with a 170 "candidate posterior" with mean $\hat{\mu}_i$ and variance \hat{V}_i – this would be the posterior on phase if the event 171 were known to be caused by peak i. $\hat{\mu}_i$ is a weighted sum of the current mean estimated phase μ_t and 172 the center ϕ_i of expectation peak *i*, weighted by their respective precisions. Note that, following the 173 predictive processing ansatz, this is the phase that minimizes precision-weighted prediction error with 174 respect to predicted event timing and predicted phase. 175
- At an event, the phase distribution resets to a Gaussian with mean $\hat{\mu}$ and variance \hat{V} . These are weighted sums of the influences of each candidate posterior, each weighted by conditional subjective hazard rate Λ_i . The expression for \hat{V} contains additional terms $(\hat{\mu}_i - \mu_{t+})^2$ and $(\mu_t - \mu_{t+})^2$, which cause the variance of the posterior to increase if the cause of the event is ambiguous.
- The background rate λ_0 acts as an alternative possible cause for any event. It serves to weight the posterior phase distribution toward the prior distribution before the event, and gives rise to causal ambiguity for any event and a resulting increase in posterior variance.
- Between events, each dt time step is taken as a Bayesian inference with likelihood $1 \lambda(\phi)dt$ and with a Gaussian prior consisting of the posterior of the previous time step carried forward by dt according to the Fokker-Planck evolution associated with equation (1). This prior causes μ_t to increase steadily and V_t to grow at rate σ^2 . The likelihood pushes μ and V away from $\hat{\mu}$ and \hat{V} with a strength proportionate to subjective hazard rate Λ . Thus, the absence of an event continuously pushes the posterior in the opposite direction as would the occurrence of an event.

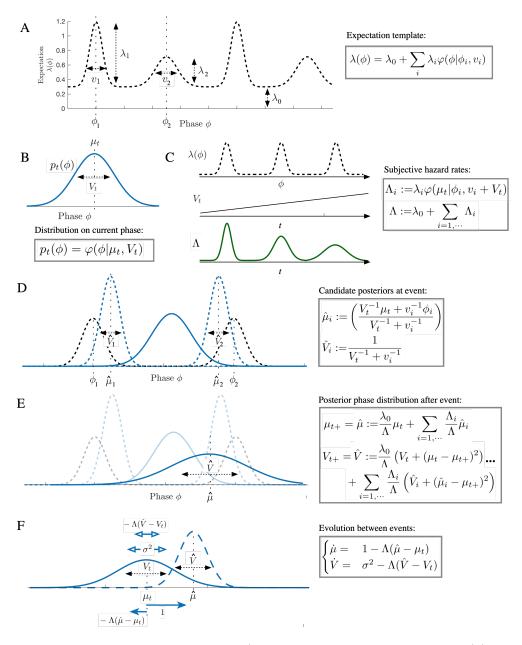


Figure 1: Illustration of the PIPPET filter. A) In the PIPPET generative model, $\lambda(\phi)$ represents the instantaneous rate of events occurring when the underlying temporal process is at phase ϕ . This is assumed to be a sum of Gaussian-shaped functions with means ϕ_i representing the phases at which specific events are expected, variances v_i representing (the inverse of) the temporal precision of the expectations, and scales λ_i representing the strength of the expectations. A constant λ_0 is also added, representing the instantaneous rate of events unrelated to phase. B) At any time t, the filter's estimate of current phase $p_t(\phi)$ is forced to be a Gaussian with mean μ_t (the estimated phase at time t) and variance V_t (the level of uncertainty about the phase estimate). D) These allow us to define a subjective hazard rate Λ (implicitly a function of time) representing the degree to which an event is anticipated at t, and conditional subjective hazard rates Λ_i representing the degree to which an event is anticipated from peak i. These hazard rates become less precise as phase uncertainty V_t increases. D) Each peak i of λ is associated with a "candidate posterior" with mean $\hat{\mu}_i$ and variance \hat{V}_i – this would be the posterior on phase if the event were known to come from peak i. E) At an event, the phase distribution resets to a Gaussian with mean $\hat{\mu}$ and variance \hat{V} . These incorporate the influences of each candidate posterior, and \hat{V} can increase if the cause of the event is ambiguous (as dramatically illustrated above). F) Between events, μ_t increases at rate 1 and V_t grows at rate σ^2 . Additionally, μ and V are pushed away from $\hat{\mu}$ and \hat{V} with a strength proportionate to subjective hazard rate Λ .

2.2 Phase And Tempo Inference from Point Process Event Timing (PATIP PET)

PATIPPET extends PIPPET by making the rate of phase advancement itself a noisy dynamic variable subject to ongoing inference. The dynamic state of the system is now a two-dimensional vector $\mathbf{x} = \begin{pmatrix} \phi \\ \theta \end{pmatrix}$, where ϕ is the phase as above, θ is the rate of phase advancement (or tempo), and σ and σ_{θ} are the levels of phase and tempo noise, respectively:

$$d\mathbf{x} = \begin{pmatrix} \theta \\ 0 \end{pmatrix} dt + \begin{pmatrix} \sigma dW_t \\ \sigma_\theta dW_t^\theta \end{pmatrix}$$
(4)

As above, an inhomogeneous point process generates events with probability based on an expectation template λ , which in this case is a function of both phase ϕ and tempo θ . In this formulation, we want events to occur with a certain probability in each $d\phi$ phase bin regardless of tempo, which we can accomplish by scaling the event rate by θ :

$$\lambda(\phi,\theta) = \theta\left(\lambda_0 + \sum_i \lambda_i \varphi(\phi|\phi_i, v_i)\right)$$
(5)

¹⁹⁹ Note that this is the same as the PIPPET expression for event rate if we set $\theta = 1$.

As before, the observer's goal is to infer a posterior distribution at any time t using preceding event times; now the distribution $p_t(\mathbf{x})$ describes an estimate of both phase and tempo. A similar derivation provides a point-process Kalman-Bucy filter that optimally serves this function within the constraint of Gaussian posteriors, providing a running estimate of a mean phase and tempo μ_t and a phase/tempo covariance matrix \mathbf{V}_t . The solution is presented in 6.1 and its derivation is presented in 6.2.

The resulting PATIPPET filter generalizes the PIPPET filter, and is identical if the initial tempo distribution is set to a delta distribution at $\theta = 1$ and σ_{θ} is set to zero. At each event, the distribution of phase and tempo is discontinuously updated to a 2D Gaussian posterior, which evolves continuously between events. This scheme is similar to [27], which estimates phase and tempo by updating a 2D Gaussian posterior, but is updated in continuous time and is significantly more flexible in its capacity to track phase based on arbitrary expectation templates.

211 2.3 PIPPET with multiple event streams (mPIPPET)

Finally, we generalize PIPPET to include multiple types of events (indexed by j), each generated as point processes with rates determined by functions $\lambda^{j}(\phi)$ of a single underlying phase:

$$d\phi = dt + \sigma dW_t \tag{6}$$

214

$$\lambda^{j}(\phi) = \lambda_{0}^{j} + \sum_{i} \lambda_{i}^{j} \varphi(\phi | \phi_{i}^{j}, v_{i}^{j})$$

$$\tag{7}$$

The Kalman-Bucy estimate of phase for this model is described by mean μ and variance V evolving according to the ODE

$$\begin{cases} \dot{\mu} = 1 - \sum_{j} \Lambda^{j} (\hat{\mu}^{j} - \mu) \\ \dot{V} = \sigma^{2} - \sum_{j} \Lambda^{j} (\hat{V}^{j} - V) \end{cases}$$

$$\tag{8}$$

and resetting to $\mu_{t+} = \hat{\mu}^j$ and $V_{t+} = \hat{V}^j$ when an event occurs in stream j, where we define Λ^j , $\hat{\mu}^j$, and \hat{V}^j as we defined Λ , $\hat{\mu}$, and \hat{V} above but in reference only to event stream j.

The same adjustment can be made to the PATIPPET generative model, and the PATIPPET filter can be similarly generalized to account for multiple event streams.

221 **3** Results

In this section we conduct a series of simulations to illustrate how the novel terms representing dynamic tracking of uncertainty and the influence of expectations in the absence of events allow the PIPPET and PATIPPET filters to reproduce perceptual and behavioral observations during human entrainment to auditory rhythms. Parameters for these simulations are listed in Appendix 6.3.

²²⁶ 3.1 Updating posterior in response to events

We simulated the PIPPET filter with a single expectation peak and varied parameters to illustrate its basic 227 behavior (Figure 2). Figure Figure 2, column i illustrates the effect of an event on the phase estimate as a 228 function of initial estimated phase μ_t . Events occurring when μ_t is near an expected event phase ϕ_1 caused 229 μ to shift linearly toward ϕ_1 . When we set the uniform rate of background events $\lambda_0 > 0$, events occurring 230 far from the expected event phase ϕ_1 were attributed to the background and therefore caused negligible 231 adjustment to the phase estimate. Phase uncertainty V_t decreased at events except when λ_0 was positive 232 and μ was not sufficiently close to ϕ_1 ; in this case, V_t increased due to causal ambiguity, or stayed the same 233 if the cause was unambiguously the uniform background source. 234

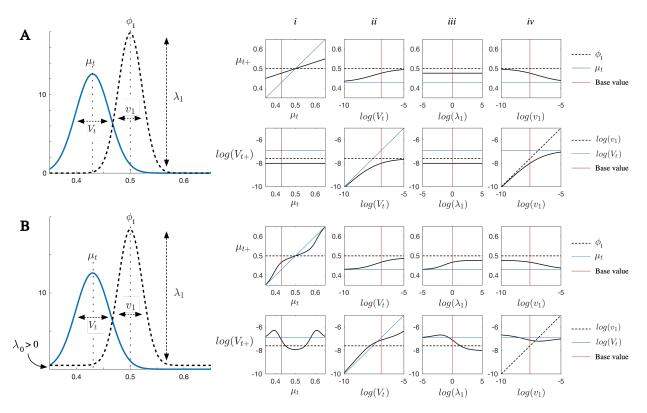


Figure 2: Characterizing PIPPET's behavior at events. A) An event is expected at phase $\phi_1 = 0.5$ with variance v_1 and expectation strength λ_1 . The expected background event rate is set to $\lambda_0 = 0$. An event occurs when the phase estimate is at μ_t with uncertainty V_t . Panels in columns *i-iv* show the resulting mean μ_{t+} and variance V_{t+} of the posterior on phase as the parameters μ_t , V_t , λ_1 , and v_1 are varied. *i*) μ is corrected linearly toward ϕ_1 , while V decreases uniformly regardless of initial phase. *ii*) Corrections to μ are more thorough when V_t is large. *iii*) These corrections do not depend on λ_1 . *iv*) These corrections are more thorough for smaller v_1 . B) The same simulations are carried out with background event rate $\lambda_0 = 0.5$. *i*) If μ_t is close to ϕ_1 , it is linearly corrected toward ϕ_1 and V_t decreases; if it is far, no correction is made. In the liminal zone, V_t increases due to the ambiguity of whether the event was related to the expectation peak or due to the background source. *ii*) V_{t+} is larger due to the effect of ambiguity as to whether the event is associated with ϕ_1 or with the background rate. *iii*) Now the correction depends on λ_1 : stronger expectations make this peak the favored cause relative to the background source. *iv*) Note that if the expectation peak is extremely narrow, V_{t+} may still be large after the event and μ_t may not fully reset to ϕ_1 due to the aforementioned causal ambiguity.

²³⁵ 3.2 Tracking complex rhythms with uneven subdivision

The PIPPET framework describes entrainment to rhythms in which each expected event phase may or may not be populated by an event. It is formulated in sufficient generality to describe entrainment to rhythms based on timing expectations with complex, non-isochronous stress patterns [28] and with noninteger duration ratios using suitably constructed (presumably learned) expectation templates $\lambda(\phi)$. Such rhythmic patterns have been shown to support highly precise synchronization in musicians with appropriate training and enculturated expectations [29], and should therefore be accounted for by models of human entrainment.

As an example of entrainment to a complex rhythm based on a temporal structure with non-integer 243 duration ratios, we simulated entrainment to a swing rhythm. The rhythm is based on an underlying grid of 244 "swung" eighth notes, where the first event of every pair is followed by a slightly longer inter-event duration 245 than the second. Though the "swing" feel is often caricatured using eighth note pairs with a 2:1 duration 246 ratio, this value has been shown to vary by with style and tempo and is certainly not limited to small 247 integer ratios [30]. We used an expectation template with a swing ratio of 3:2 (though the exact ratio is not 248 important) and associated the first eighth note in each pair with a stronger expectation than the second. 249 The PIPET filter entrained to a complex, syncopated rhythm based on this template, drawing on the timing 250 of both strongly and weakly expected events (Figure 3A). It corrected its phase estimate when an event 251 timing shift or a phase shift was introduced into the rhythm (Figure 3B and 3C). 252

²⁵³ 3.3 Failure mode: too much syncopation

The phase inference framework can account for human failures to track perfectly timed rhythms, i.e., rhythms in which every event falls at a peak of the expectation template. A prime example of this failure mode in human rhythm tracking is tracking overly syncopated rhythms (rhythms with a predominance of events at time points with weaker expectations). Listeners tend to "re-hear" such rhythms by attributing events to metrical positions where events are more strongly expected [31, 32].

In PIPPET, these failures consist of inferring the presence of phase noise where none actually occurred. Such behavior is a necessary consequence of Bayesian optimality: a given stimulus may be generated by different combinations of phase noise and point process event generation noise, and the inference process is concerned only with the most likely explanation for the stimulus, which may include phase noise even if the stimulus was actually generated without it.

Using the expectation template with a swing grid as in the previous section, we simulated a strongly syncopated rhythm (Figure 4A). The rhythm's phase was not tracked successfully due to a convergence of two

bioRxiv preprint doi: https://doi.org/10.1101/2020.11.05.369603; this version posted March 6, 2021. The copyright holder for this preprint (which was not certified by peer review) is the author/funder, who has granted bioRxiv a license to display the preprint in perpetuity. It is made available under aCC-BY 4.0 International license.

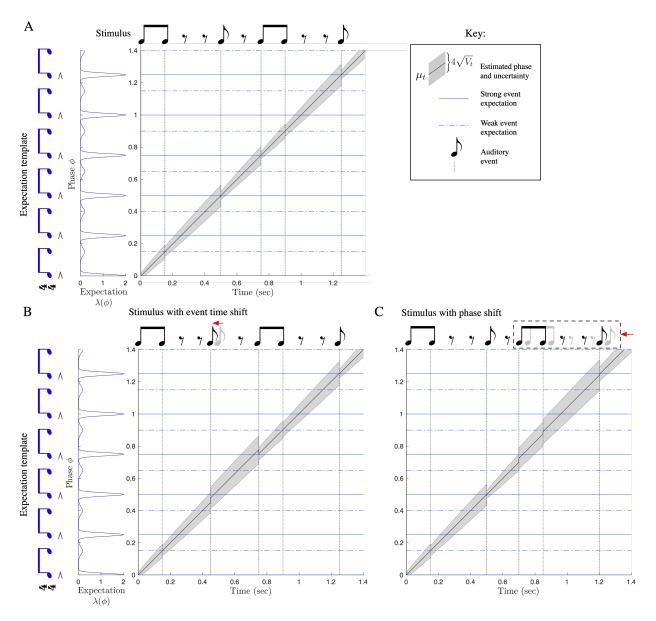


Figure 3: Tracking phase through swung rhythms. PIPPET is given a pattern of expectations representing "swung" eighth notes, with alternating longer and shorter inter-event durations and stronger, more precise expectations on the first of every pair. Dotted lines correspond to weaker expectations and solid lines correspond to stronger expectations. A) Phase is successfully tracked over the course of a rhythmic stimulus, with phase uncertainty growing between events and contracting at events. B) One event in the rhythm is shifted earlier in time. Estimated phase μ_t adjusts partially to compensate for the timing shift, and then adjusts back at the subsequent event. Uncertainty V_t is not as effectively reined in by these unpredictablytimed events, but decreases as later events corroborate the corrected phase estimate. C) A phase shift is introduced into the rhythm, moving all subsequent events earlier in time. When the first early event arrives, uncertainty increases. Estimated phase is corrected over the first few events after the shift, and V_t decreases most substantially when the estimate μ_t is corroborated by a strongly expected event happening at the appropriate estimated phase.

- ²⁶⁶ factors: the disproportionate influence of the higher peaks of the expectation template, and the accumulation
- $_{267}$ of phase uncertainty V_t . Phase uncertainty was only slightly reduced by events occurring at weakly expected

phases, so it accumulated over the course of the rhythm, and especially during the long silence. Once V_t was large, indicating the possibility of substantial phase noise having accumulated, the higher expectation peaks ϕ_i became the most likely explanations for events that were actually perfectly timed to coincide with nearby lower peaks – since precise event timing was no longer a reliable indicator of the source of an event, local peak height became the best indicator, and higher peaks won out. Thus, at each event, the estimated phase was adjusted to better align the higher peaks with the events.

The same rhythm could be successfully tracked in two alternate conditions. First, it was successfully 274 tracked when we decreasing the rate of accumulation of phase uncertainty σ^2 (Figure 4B), demonstrating the 275 key role of uncertainty in making the system susceptible to the disruptive effect of syncopation. Second, it 276 was successfully tracked when an additional stream of sensory input was added by simulating an isochronous 277 finger tap (Figure 4C). We used mPIPPET to create a second expectation template for tapping. As phase 278 tracking was simulated, we planned new tap events just before μ reached expected tap phases by extrapolating 279 μ forward. When taps occurred, phase uncertainty decreased, reducing the disruptive effects of syncopation. 280 Note that planning actions specifically to fulfill sensory expectations and using this sensory feedback to 281 inform inference about the outside world is an example of "active inference", the principle framework for 282 understanding action in the literature on predictive processing [25]. 283

²⁸⁴ **3.4** Tempo inference

We simulated the PATIPPET filter with basic metronomic expectations to observe its capacity to infer phase and tempo at once. We gave the model a wide initial range of possible tempi and a simple metronomic stimulus with actual tempo near the upper end of that range. In these conditions and with the parameter set we chose, the model established the appropriate tempo and phase to within a tight range over the course of the first two events (Figure 5).

In addition to its value as a model of human rhythmic cognition, the PATIPPET filter shows promise as a general-purpose tempo tracking algorithm for musical applications. This would require a principled method of choosing values for the various free parameters of the generative model, which might be done a priori based on a labeled corpus, adaptively over the course of listening, or through some combination of the two. We leave a more thorough exploration of the relative performance of this model to future work.

²⁹⁵ 3.5 Period-dependent corrections

In sensorimotor entrainment literature, finger taps entrained to a metronome generally shift to correct a certain fraction of an event timing perturbation on the next tap. This fraction is called α . In human

bioRxiv preprint doi: https://doi.org/10.1101/2020.11.05.369603; this version posted March 6, 2021. The copyright holder for this preprint (which was not certified by peer review) is the author/funder, who has granted bioRxiv a license to display the preprint in perpetuity. It is made available under aCC-BY 4.0 International license.

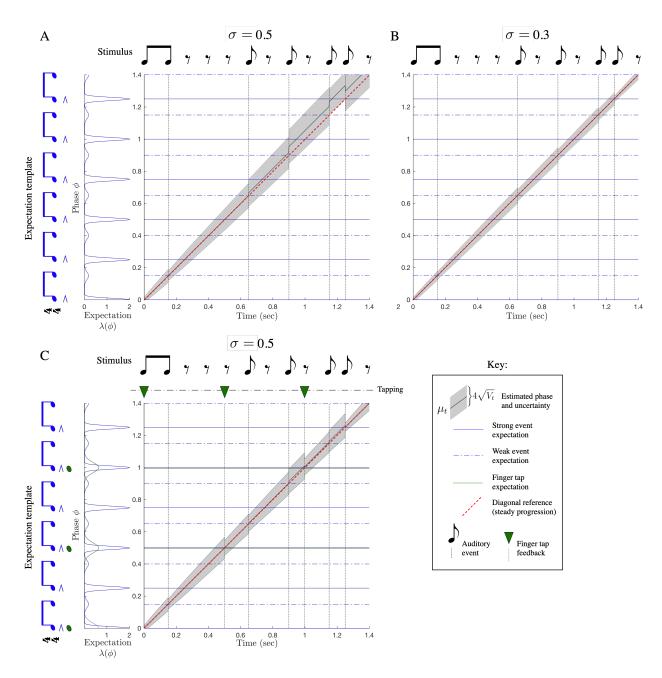


Figure 4: Too much syncopation causes rhythm tracking failure. A predominance of events associated with weak expectations combined with accumulated phase uncertainty can lead to a failure to track phase accurately. A) In this example, phase uncertainty V increases over a long silence. At the next event, this high uncertainty leads the model to partially attribute a weakly expected event to the nearby phase at which an event is strongly expected. As a result, the model ends up aligning the fifth event with a strong phase rather than a weak one, and overestimating phase at the final event (correct phase marked with yellow dot). B) When the rate of accumulation of phase uncertainty (i.e., the expected phase noise σ^2) is decreased, phase is tracked correctly. C) Alternatively, phase can be tracked successfully by inserting an isochronous stream of finger taps and a suitable template for the alignment between expected auditory feedback from the taps and phase. We use mPIPPET to simulate an expectation for isochronous taps (green notes and trace on the left). For simplicity, taps are placed every 0.5 sec; however, even noisy taps generated based on estimated phase could serve to reduce phase uncertainty and avoid a total phase tracking failure.

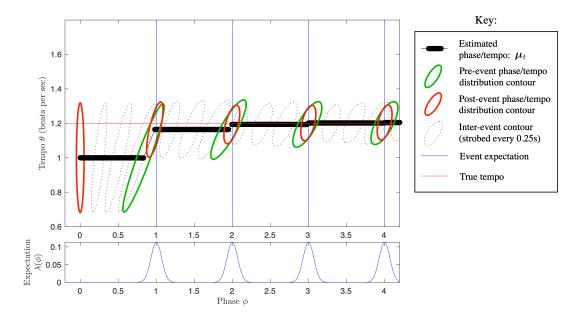


Figure 5: The PATIPPET filter estimates phase and tempo. PATIPPET is initialized with high tempo uncertainty. The first event occurs relatively early, causing the estimated tempo to increase. Each subsequent event occurs close to the time expected based on the estimated phase and tempo, causing the posterior to contract in both the phase and tempo direction as its prediction of event time is fulfilled and its phase and tempo estimates are corroborated. Ultimately, PATIPPET settles on a narrow distribution around the appropriate tempo as it continues to accurately estimate phase.

subjects, α has repeatedly been observed to increase linearly with metronome period ("inter-onset interval,"

²⁹⁹ or IOI), exceeding 1 (i.e., over-correction) for sufficiently long IOIs [33, 34].

The phase inference framework offers a principled explanation for α increasing with IOI. During an eventfree interval, phase uncertainty increases over time. When an event does occur, the precision of the prior distribution on phase and tempo is weighed against the precision of the likelihood function associated with the expectation of that event. If the prior is less precise due to accumulated uncertainty, the precision of the likelihood weighs more heavily against it and the adjustment in phase is more thorough. Thus, all else being equal, events spaced more widely apart in time induce more extensive phase corrections.

Since the strongest phase correction PIPPET can make at an event is to fully update the phase estimate to the expected event time, it cannot account for α values above 1. However, it has been previously suggested that α may exceed 1 for long metronome periods due to some period correction occurring in addition to phase correction [33]. We were therefore curious to see whether PATIPPET could reproduce the linear increase of α with increasing IOI up to and beyond $\alpha = 1$.

In Figure 6, we show that with appropriate parameters, PATIPPET can indeed reproduce the experimental observation of a near-linear increase in α from below to above 1 as IOI increases. In PATIPPET, this

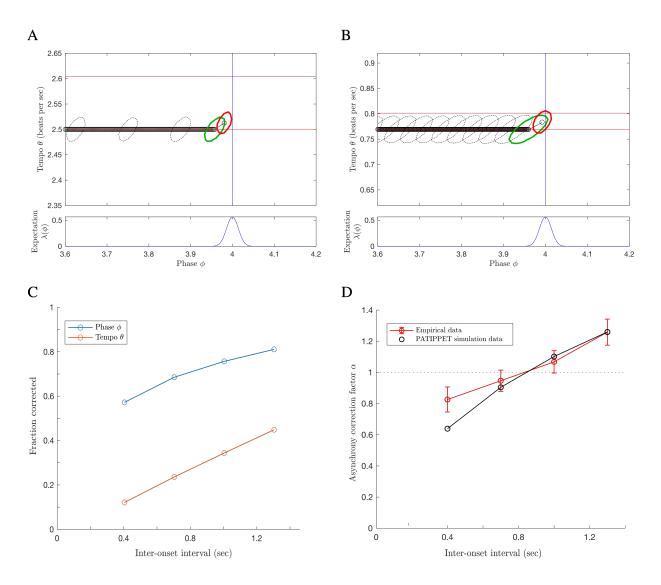


Figure 6: **PATIPPET reproduces human tapping data showing stronger error correction for longer inter-onset intervals.** A and B) The distribution on phase and tempo leading up to and following a phase shift at the fourth event in an isochronous sequence for two different metronome tempi, i.e., two different inter-onset intervals. (Same color key as Figure 5, but with phase/tempo distribution contours strobed every .05 sec.) Note that when the IOI is short, PATIPPET arrives at the phase-shifted event with a high degree of phase and tempo certainty. C) PATIPPET makes a proportionally larger correction to phase and tempo for long IOIs than for short IOIs due to the greater degree of uncertainty preceding each event. D) Alpha (α) is the proportion of a phase shift that is corrected at the next tap time. With this set of parameters, PATIPPET reproduces the empirical observation from [34] that the phase shift is undercorrected when IOIs are short and overcorrected $\alpha > 1$ when IOIs are long.

- ³¹³ phenomenon is a natural consequence of optimal inference in the context of phase and tempo uncertainty
- 314 that accumulates between observed events.

315 3.6 Time warping in the absence of expected events

When an event in a rhythmic stimulus is strongly expected but no event occurs, an optimal Bayesian observer should initially be biased to believe that in spite of their current phase estimate, the stimulus may not have reached the expected event phase yet. The result should be that a perfectly timed event later in the stimulus will seem to be arriving earlier than expected: in other words, the tempo of the stimulus will seem to accelerate. The degree of this effect will depend on the observer's degree of phase and tempo uncertainty.

There is evidence of such an effect in human rhythm perception. The "filled duration" illusion is the 321 impression that an isochronous sequence has changed tempo when it is initially subdivided by additional 322 predictable events and then subdivisions are eliminated. According to multiple reports, the magnitude of this 323 effect is reduced or eliminated if the empty intervals precede the filled intervals [35, 36, 37, 38] (though there 324 is some disagreement about this [39]), suggesting that it is indeed the established expectation of continuing 325 subdivision that interferes with the perceived passage of time when the subdivisions cease. A second result 326 that could be similarly accounted for is the surprising finding in [40] that a participant tapping along with 327 a subdivided beat delays their tap following the omission of an expected subdivision. If taps are planned to 328 coincide with the arrival of a specific mean estimated phase, then the slowing of estimated phase induced by 329 an omission of a strongly expected event should indeed delay the subsequent tap. 330

We stimulated PATIPPET with a strong isochronous expectation template by scaling up λ and presented it with a "filled duration" in which all expected events occurred and an "empty duration" in which events occurred only at the beginning and end of the interval (Figure 7). PATIPPET loyally tracked phase through the filled duration; however, when strongly expected events were omitted, the mean phase estimate slowed down at each expected event phase, leading to an overall slowing in estimated phase advance and an unexpectedly early onset of the event marking the end of the empty duration (Figure 7A).

Specifically timed event expectations are not necessary to produce a filled duration illusion: random 337 raindrop sounds were sufficient to lengthen produced intervals during audiomotor synchronization task [41]. 338 In PATIPPET, a filled duration effect was also produced when the expectation template consisted only of 339 a high expected background rate of events λ_0 . In this case, estimated phase advance slowed during the 340 empty interval because estimated tempo dropped. The PATIPPET filter effectively noted that not as many 341 events were occurring as expected, and in response it lowered estimated tempo because a lower event rate is 342 expected at a lower tempo. This type of explanation could be invoked to offer a normative account for other 343 non-rhythmic filled interval illusions, though doing so is beyond the scope of this work. 344

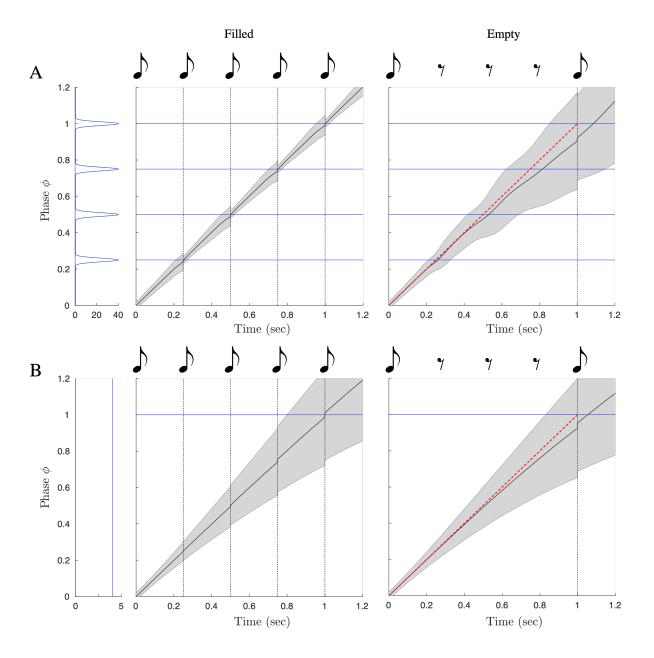


Figure 7: The filled duration illusion: time warping by the omission of strongly expected events. (Same image key as 4, with shading displaying PATIPPET phase variance.) A) PATIPPET is simulated with strong expectations for isochronous events. Left: When a set of strongly expected events occur as expected (a filled duration), estimated phase stays on track, advancing (on average) at a rate of 1. Right: When the duration is empty, estimated phase deviates from steady progression (red diagonal) by dragging as each expected event point approaches and passes, leading to the illusion that the event marking the end of the interval has arrived earlier than expected. B) PATIPPET is simulated with a high expected background rate of events λ_0 , but no phase-specific event expectations ϕ_i . In this case, too, an empty duration leads to dragging estimated phase and an unexpectedly early final event.

345 4 Discussion

Here were have presented PIPPET, a framework representing entrainment to a time series of discrete events 346 based on a template of temporal expectations. PIPPET treats the event stream as the output of a point 347 process modulated by the state of a hidden phase variable. The PIPPET filter uses variational Bayes 348 to continuously estimate phase and track phase uncertainty based on this generative model. PATIPPET 349 extends PIPPET to include a generative model of tempo change, and the PATIPPET filter simultaneously 350 estimates phase, tempo, and the covariance matrix representing their uncertainty and their codependence. 351 This framework is intended to serve as a hypothesis for how the human brain integrates auditory event 352 timing to inform and update an estimate of the state and rate of an underlying temporal process. 353

PIPPET and PATIPPET reproduce several qualitative features of human entrainment, including realistic failures to track overly perfectly-timed but over-syncopated rhythms, perceived acceleration of a metronomic pulse when strongly expected events are omitted, and error correction after metronome timing perturbations that increases with increasing inter-onset interval. We show that these three phenomena all follow naturally from our framing of entrainment as a process of Bayesian inference based on specific phase-based temporal expectations.

³⁶⁰ 4.1 Relationship to other models of timing

The dynamics of PIPPET and PATIPPET in response to sensory events are similar to dynamics of other 361 entrainment models that correct phase and period based on event timing, e.g., [42, 43]. Models based on 362 Dynamic Attending Theory, e.g., [11, 12], are also similar in explicitly modeling timing expectations and 363 their effect on phase and period adjustment. The phase inference framework differ from these existing models 364 in four key ways. First, they are derived as optimal solutions to specific inference problems, and therefore all 365 modeling decisions can be justified within a normative framework. Second, they are formulated in sufficient 366 generality to describe entrainment based on non-isochronous and even aperiodic temporal expectations, an 367 area that has lately received increasing experimental attention [6, 44, 45] but has been largely neglected in 368 entrainment modeling. Third, they allow expectations to influence the inferred phase even in the absence 369 of sensory events, creating the time-warping effect of disappointed expectations evidenced in humans by the 370 "filled duration" illusion. Finally and most critically, they explicitly track uncertainty in phase and tempo, 371 providing a system for moderating between assimilation of new timing data and loyalty to an internal sense 372 of time. 373

Bayesian methods have been used elsewhere to analyze rhythmic structure as time series of point events. Some of these are application-focused methods that require offline analyses [46, 47] and therefore do not

³⁷⁶ serve as satisfying models of real-time behavior. Cemgil et al (2000) [27] use a Kalman filter that tracks a ³⁷⁷ distribution on phase and tempo similarly to PATIPPET. However, this model is structured to infer phase ³⁷⁸ and tempo event-by-event rather than in continuous time, and is not equipped to handle complex rhythms ³⁷⁹ or temporal structures more complex than approximate isochrony.

Bayesian inference has also been used to model timing estimation in the brain (e.g., [20, 21]), but it is 380 generally used to describe inferences about discrete variables like interval durations and event times, whereas 381 PIPPET describes a continuous inference process underlying predictions about event times. One such model 382 leading to particularly PIPPET-like results was presented in Elliot et al 2014 [22]. The authors created a 383 Bayesian model to explain the results of an experiment that had participants tap along to a stimulus consist-384 ing of two jittered metronomes. The model behaves similarly to PIPPET in that it estimates the next event 385 time using a weighted average of previous event times and prior beliefs, with weights informed by expected 386 timing precision. However, like [27], their model infers the anticipated timing of discrete, metronomic events, 387 whereas PIPPET predicts and updates an underlying phase in continuous time and can therefore generalize 388 to non-isochronous and complex rhythms and account for the effects of event omissions. Additionally, in or-389 der to account for participants ignoring events far from predicted time points, they introduce the assumption 390 that participants repeatedly test the hypotheses that events come from one or two separate streams, whereas 391 PIPPET naturally accounts for this phenomenon by attributing stray events to a uniform background event 392 rate λ_0 . 303

³⁹⁴ 4.2 Interpreting the generative model

The PIPPET generative model is formulated as though it implements perfect variational Bayesian inference 305 on inherently stochastic stimuli. However, Bayesian computations in the brain are often invoked to com-396 pensate for internal as well as external sources of stochasticity [48], and in the case of PIPPET the most 397 reasonable interpretation may be a combination of the two possibilities. In reality, we do not often listen 398 to musical rhythms with random timing and phase jitter; however, neural noise and interaction with other 399 ongoing processes may introduce timing variability into the processing of sensory events and give rise to 400 variability in the process of tracking estimated phase. This interpretation also allows for changes in gener-401 ative model parameters based on internal states that might affect internal noise levels, e.g., attentiveness 402 (which has been shown to affect tempo correction but not phase correction [49], and which therefore might 403 be modeled through its effect on σ_{θ}). Ideally, the phase inference framework could be reconstructed based 404 on assumptions of a combination of internal and external noise; however, that is beyond the scope of the 405 current work. 406

Given this ambiguity, the generative model parameters may ultimately reflect some combination of the 407 empirical statistics of rhythmic stimuli and internal factors. We briefly discuss the precision parameters v_i 408 as an example. First, an upper bound on the precision of expected event timing is the precision of sensory 409 timing perception, which is, for example, high for human audition and significantly lower for human vision¹. 410 Second, expected event timing precision may further reflect the observed relative timing distributions of 411 event streams. These observations may inform expectations on time scales ranging from a single sitting to 412 a lifetime of listening. Expected timing may be learned separately for different sensory modalities, different 413 musical genres (e.g., techno vs. funk), or even different instruments (e.g., kick drum, snare, hi-hat, as 414 discussed below). The precision of a beat-based temporal expectation is closely related to the width of a 415 "beat bin," the window of time (rather than a single time point) that is proposed to constitute the "beat" 416 in [50], and to the width of the temporal "expectancy region" described in dynamic attending theory [11]; 417 in both cases, this width is increased by imprecision in the immediately preceding stimulus. 418

419 4.3 Testable behavioral predictions

Given the ambiguous interpretation of the generative model discussed above, the question of whether human expectation-based entrainment is truly described by a normative framework may be ill-posed. However, two key qualitative elements of this framework can be tested directly: the tracking of phase uncertainty and the influence of expectations in the absence of events. Seeking further experimental evidence of these two phenomena would help determine the value of phase-inference-based models in describing human entrainment behavior.

The phase inference framework predicts that the accumulation of uncertainty over the course of empty time has a critical effect on the perceptual interpretation of subsequent events. In Figure 4, we show a rhythm that is perceptually misinterpreted due in part to empty time preceding syncopation. An experiment could be designed along the lines of [32] to test this aspect of the phase inference framework by measuring the effect of empty time on the interpretation of rhythmic stimuli that follow.

A second prediction along these lines is that various measurable perceptual phenomena, including perioddependent error correction in motor entrainment, perceptual parsing of ambiguous rhythms, and susceptibility to temporal illusions such as the filled duration illusion, should depend critically on levels of phase and tempo uncertainty. Assuming that the parameters of uncertainty tracking vary across individuals, the PIPPET/PATIPPET framework would predict correlations in measurements across these domains: certain

¹An event can only be experienced after it occurs, so (as pointed out in [21]) the likelihood function on underlying phase associated with this type of uncertainty should be asymmetrical. The analytically tractable incarnation of our framework presented here uses Gaussian likelihood peaks, so cannot account for the effect of asymmetrical likelihoods; however, we could posit a λ function with asymmetrical peaks and use numerical methods rather than the explicit solution derived here to estimate underlying phase at each time step.

individuals should show increased sensitivity to temporal illusion, misleading rhythms, and the effect of period on error correction. Further, stimulus manipulations that affect phase and tempo uncertainty, including
the temporal precision of the auditory events and the length of the click train establishing an initial tempo
estimate, should have direct and predictable effects on these perceptual and behavioral measures.

Third, the phase inference framework predicts that omissions of strongly expected events should sys-440 tematically distort estimates of phase and tempo, or, perhaps indistinguishably, of elapsed time. These 441 effects could be explored by parametrically manipulating event expectations through priming stimuli and 442 then measuring distortions induced by event omissions through perceptual report or timed motor response. 443 If we find situations in which human behavior qualitatively differs from solutions to the inference prob-444 lems posed by PIPPET and PATIPPET, these can be interpreted in two perfectly valid ways: either human 445 behavior has not been optimally tuned for the task at hand, or we have not correctly identified and encap-446 sulated the task and its survival-relevant objective. If we follow the latter interpretation, we might attempt 447 to refine the generative model, e.g., by introducing the belief that tempo changes occur in jumps or ramps 448 rather than as random drift, or to modify the objective of the task, e.g., by including additional cost functions 449 or priors associated with perceptual report or motor output as discussed above. 450

451 4.4 Application to analysis of behavioral data

The phase inference framework offers a predictive processing lens for understanding the results of rhythm perception and production experiments. Given a perceptual or behavioral task, we can suppose that motor or perceptual human entrainment behavior is optimally solving an inference problem, and determine the parameters of that problem by fitting them with appropriate methods. These parameters come with natural interpretations in the language of prediction and precision. We can then study the changes in these parameters over the course of an experiment, over different variations on the same experiment, over the human lifespan, across cultures, etc.

For some experimental data, the many parameters available in PIPPET may prove redundant. For 459 example, the observation of weak error correction in entrained tapping could be explained by imprecise 460 auditory timing expectations (high v_i), an overly precise internal model of phase (low V_t , caused perhaps by 461 low σ), or overly precise tap feedback timing expectations (as discussed below). However, we believe these to 462 be meaningful distinctions that call for disambiguation through carefully designed experiments – for example, 463 skipping taps to separate out the precision effects of tapping feedback or varying silent durations within the 464 stimulus to separate the accumulating effects of phase uncertainty V_t from the history-independent effects of 465 timing expectation uncertainty v_i . For experiments that do not take such measures, redundant parameter 466

⁴⁶⁷ sets that fit the data may be interpreted as meaningfully different possible interpretations of the results.

468 4.4.1 Multiple event characteristics

mPIPPET generalizes the PIPPET/PATIPPET framework to cases of multiple distinguishable event types, 469 each with its own set of expectations as a function of phase. One example could be listening, tapping, or 470 dancing to a kit drum track with bass drum, snare, and hi-hat cymbal. Timing perturbations of different 471 instruments in drum rhythms have been shown to differently affect human entrainment [51]. By letting j472 take values from $\{bass, snare, hihat\}$ and choosing appropriate values for ϕ_i^j, v_i^j , and λ_i^j for each event i on 473 the metrical grid, one could create a set of timing expectations with strength and precision dependent on 474 the specific drum and metrical position that could then be used to optimally track underlying phase and 475 tempo through a complex kit drum rhythm. A similar setup could be used to implement the assumption that 476 pitches in a melody match the harmonic context more often in strong metrical positions, allowing rhythm 477 parsing during melody listening to be influenced by scale degree. 478

Alternatively, the j index may be used to treat events over multiple sensory modalities. Visual event timing is judged with less precision than auditory event timing in perceptual report [21] and in timingsensitive sensory pathways [52], and might therefore be modeled with a less precise expectation template. (Note, however, that visual information may not have the same access to motor-related brain regions used for auditory entrainment [53], so the same modeling framework may not be appropriate.)

⁴⁸⁴ mPIPPET with $j \to \infty$ can be used to account for a continuum of event types. Thus, we could create a ⁴⁸⁵ forward model in which it is more likely for notes played with stronger accents to fall on strong beats, or in ⁴⁸⁶ which lower pitches are expected with higher timing precision [54] and therefore exert greater influence on ⁴⁸⁷ neural entrainment [55].

The phase inference framework could be further generalized to take into consideration additional stream of continuous input. This could be visual input from watching a pendulum, auditory input from a continuously modulated sound, or proprioceptive feedback from continuous entrained motion (as opposed to discrete, timed proprioceptive feedback like tapping). This goes beyond the scope of the mathematics presented here, but is a straightforward application of results proven in [23].

493 **4.4.2** Tapping

As illustrated in Figure 4, mPIPPET can be used to describe entrained tapping data. Experiments have shown that the presence of entrained tapping prior to temporal perturbations in a metronomic stimulus reduces the phase correction response [56], indicating that the estimate of moment-by-moment phase is influenced by the proprioceptive, tactile, and auditory feedback from tapping. The phase inference framework

is well-suited to modeling this influence as its own separate stream of informative input, though a thorough 498 tapping model would require introducing noise into tap execution and into the phase tracking process itself. 499 Importantly, using tap times to inform an estimate of underlying phase challenges our interpretation of 500 this phase representing a purely external source of temporally patterned events. Instead, the inferred phase 501 would be a hybrid of an external phase and the phase of one's own motor cycle. Functionally, this is similar 502 to the perceptual oscillator forced by both an external stimulus and one's own periodic action proposed by 503 [57]. This may be an especially useful way to think about synchronization with another agent, where one 504 can adopt strategies ranging from following (assigning high precision to input from the other) to leading 505 (assigning low precision to input from the other, and possibly higher precision to self-generated events). See 506 [58] for a discussion of such a coding strategy as a means of minimizing representational neural resources. 507

⁵⁰⁸ 4.4.3 Aperiodic rhythm, speech, and musical grammar

One specific question that the phase inference framework might help resolve is how periodic and nonperiodic entrainment differ. PIPPET does not intrinsically differentiate between these two processes; however, since it is sufficiently general to model both, it could guide an exploration of parameter differences between the performance of similar tasks in periodic and aperiodic contexts. (For neural and behavioral evidence of differences between memory-based and periodicity based entraiment, see, e.g., [45, 6].)

By accommodating aperiodic expectations with any degree of precision or imprecision, the phase inference 514 framework may be especially well-suited to modeling the loose temporal regularities of speech [59]. However, 515 as currently formulated, it is limited in that expectations are not history-dependent: the occurrence or 516 absence of an event does nothing to the expectancy of an event at a later timepoint. This is appropriate 517 for modeling the metrical aspect of rhythmic expectancy, but does not address the grammar-like structure 518 of music rhythm [60], i.e., the expectation of certain temporal patterns of events over others regardless of 519 their metrical positions. Speech, of course, is even more thoroughly grammatical, with certain sound events 520 strongly shaping the temporal and spectral patterns expected in the immediate future. 521

Such effects could be readily incorporated into the phase inference framework by adding history dependence to the expectation template λ , though that is beyond the scope of this work. The precise details of this history dependence in rhythm parsing could be based on any suitable formal model for rhythmic grammar (e.g., [61, 62, 60]), and for speech applications could include whatever aspects of the co-dependence of timing and content expectations were appropriate for the task at hand.

⁵²⁷ 4.5 Limitations and possible extensions of the phase inference framework

528 4.5.1 Perceptual vs. motor entrainment

PIPPET is formulated as a perceptual process, without specific reference to how entrained movement is produced by this process. In presenting the PIPPET framework and using it to explain tapping results, we have posited that perceptual and motor entrainment are rooted in the same internal tracking of the phase of an external process. However, perceptual and motor measures of entrainment sometimes give conflicting results: for example, exposure to musical performance with expressively irregular timing affects perceptual reports of timing in subsequent stimuli [63], but does not affect phase correction in tapping to subsequent stimuli [64].

We expect that both physical entrainment and perceptual report are informed by a neural process of 536 estimating underlying phase. Principles of economy suggest that they should share in such an estimate rather 537 than drawing on separately instantiated processes of neural inference, and experimental correlations between 538 motor and perceptual results tentatively support this conclusion (e.g., [65]). However, it is possible that rapid, 539 automatic audiomotor adjustment mechanisms have been selected to prioritize speed over precision (e.g., the 540 spinocerebellar vermis [66]), especially in the case of entrainment to simple isochronous stimuli, and thus 541 may not take uncertainty into account. If this is the case, then motor entrainment experiments not be clean 542 indicators of perceptual management of uncertainty until the effects of these mechanisms are separated out. 543

544 4.5.2 Learning expectation templates

If the brain does treat entrainment as a process of inference based on a generative model, this raises the question of how the properties of the generative model are established in the first place. The PIPPET framework does not address this question directly, but by examining the parameters necessary to formulate PIPPET, we can clearly see what components need to be in place before a process of continuous phase and tempo updating can begin.

First, the brain must learn the temporal structures of the expectation template for rhythmic expectation. Learning these underlying structures from an experiential corpus of noisy, complex rhythms is not trivial. It seems likely to involve some type of bootstrapping in which a recognition of some degree of temporal structure allows for attribution of events to positions in that structure, allowing for deeper structure learning. Earlier exposure to simpler, less complex rhythms would likely help with such a bootstrapping process. (For a discussion of the challenges of this type of simultaneous learning and filtering and a proposed solution for non-point-process data, see [67].)

⁵⁵⁷ The brain must also learn noise and precision parameters for the model. Note that neither the temporal

expectation variance parameters v_i nor the noise parameter σ necessarily correspond to the actual precision of the neural or external timing mechanisms in play. The brain may underestimate the noisiness of the timing process it uses to track underlying phase, leading to under-adjustment to auditory event timing and minimal time-warping between events, or do the opposite. Presumably, these parameters must be learned through experience and prediction error.

563 4.5.3 Selecting and updating expectation templates

When the brain is exposed to a rhythmic stimulus, it must first recognize that a predictable pattern exists and 564 select an appropriate expectation template from its learned repertoire. This is its own process of inference, 565 and may be amenable to a Bayesian description. Since the PIPPET filter maintains a unimodal posterior, 566 it is not well-suited to model this initial inference process, which may require maintaining a distribution 567 over multiple distinct possible starting phases and expectation templates. This problem might be partially 568 addressed by incorporating a model that evaluates multiple distinct hypotheses for beat or meter (e.g. [68, 569 69], or [70] with appropriate probabilistic interpretation) as an additional level of inference in parallel with 570 ongoing phase and tempo inference. 571

572 4.6 PIPPET in the brain

Though PIPPET and PATIPPET are abstract models not committed to a particular brain-based implementation, advances in the brain basis of timing and beat-keeping combined with the hypothesized neural bases of predictive processing suggest the beginnings of a plausible approximation of PIPPET in the brain, described below.

The essential aspect of the PIPPET framework that qualitatively differentiates its behavior from previous 577 models is the explicit tracking of uncertainty over time for the purpose of informing the relative weights of 578 sensory event timing and internal state estimates. There have been various proposals of how uncertainty 579 is represented and utilized in the brain, and the system likely differs by task and type of uncertainty [71. 580 48]. One proposal is of particular interest in relation to timing: uncertainty about a hidden state may be 581 computed in medial frontal cortex and signalled via dopaminergic neurons in the ventral tegmental area [72]. 582 In this case, the hidden state would be the phase and tempo of the stimulus. This proposal is consistent 583 with the observations that dopaminergic neurons encode of certainty in the temporal expectation of sensory 584 cues [73] and that dopamine receptor antagonism in humans causes increased timing uncertainty [74]. 585 In the predictive processing literature, dopamine is often given the role of signaling certainty ("expected 586

⁵⁸⁷ precision") across levels of hierarchical processing [75]. In this framework, it participates in probabilistic

computations by weighting the input to error-calculating neural populations, causing these errors to be weighted more heavily in the ongoing process of error-minimization that implements variational Bayesian estimation of hidden states. Different dopaminergic populations may signal precision at different levels of processing; in particular, dopamine may signal precision of both higher-level state estimates and lower-level sensory expectations. Thus, phase certainty V_t^{-1} and expected timing precision v_i^{-1} may both influence computation through dopaminergic signalling.

Experiments with non-human primates have shown neural trajectories in medial premotor cortex (MPC, 594 encompassing the supplementary and pre-supplementary motor areas) that represent progress through self-595 generated behavioral processes. The author hypothesizes in [76] that similar trajectories represent rhythmic 596 phase in human MPC. A representation of a linear phase ϕ , used in the phase inference framework for 597 flexibility and mathematical tractability, would seem to be a limiting factor for implementation in the 598 brain. For shorter, aperiodic learned patterns of temporal expectation, phase could be represented by short. 599 aperiodic trajectories [77], as observed in primates in timed response tasks; for simple periodic patterns, phase 600 could be represented circularly [78], as observed in isochronous tapping tasks; and for longer, hierarchical 601 patterns, phase could be represented by hierarchically structured trajectories that loop but also evolve in 602 other dimensions, as observed in cyclic behaviors whose sensory components change from one cycle to the 603 next [79]. 604

Guided by the "Action Simulation for Auditory Prediction" (ASAP) hypothesis presented in [80] and 605 further developed in [76], the theory of hierarchical predictive processing [81], and the predictive functions 606 proposed for the dorsal auditory pathway [82, 83], we propose a neural implementation of PIPPET's phase 607 estimation in Figure 8. An essential aspect of this account is that it does not insist on the mathematical 608 convenience of instantaneous phase updates, which are obviously implausible in the brain. Instead, precise 609 timing predictions are issued with appropriate timing to intercept rising sensory signals, and the resulting 610 timing errors are then be used to update phase through an error minimization process over the next few 611 hundred milliseconds. 612

⁶¹³ Briefly, phase is represented by stereotyped trajectories of population firing rates in MPC, and phase ⁶¹⁴ uncertainty is also represented locally in medial frontal cortex [72]. Basal ganglia selects and activates ⁶¹⁵ an expectation template appropriate to the context. This template is combined with phase and phase ⁶¹⁶ uncertainty estimates in MPC to compute a momentary subjective hazard rate Λ . The hazard rate is sent ⁶¹⁷ to parietal cortex as a prediction of event-based input, where it meets ascending pulses from the auditory ⁶¹⁸ system associated with auditory events (which may be relayed rapidly from the dorsal cochlear nucleus via ⁶¹⁹ cerebellum [19]). "Event prediction error" from parietal cortex returns to MPC, where it pushes μ in the ⁶²⁰ direction that reduces error: toward expected event phases at events and away from them between events.

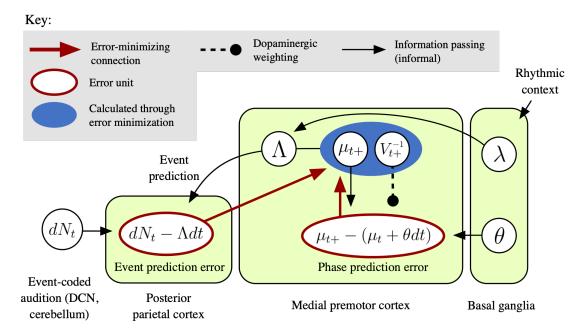


Figure 8: A possible implementation of PIPPET in the brain. This diagram embeds a formal predictive coding error minimization scheme is embedded within an informal information-passing schematic to outline how estimated phase μ_t might be calculated and updated on each dt time step by a network of interacting brain regions. Estimated phase μ_t and phase uncertainty V_t are represented in medial premotor cortex (MPC). These estimates are used to calculate instantaneous subjective hazard rate Λ with the help of basal ganglia, which has selected an expectation template λ based on recent rhythmic context. The hazard rate is sent to parietal cortex, where it acts as a prediction of pulses rising from the event-based auditory pathway. An "event prediction error" signal comparing pulses to their prediction is sent back up to MPC, where it pushes μ_{t+} in the direction that reduces prediction error – strongly toward local expectancy peaks when events occur, and weakly away from them when there are no events. (Note that phase updating at events is assumed to be rapid but not instantaneous as represented in the PIPPET filter.) The event prediction error is counterbalanced by a local "phase prediction error" signal generated through local interactions within MPC that pushes μ_{t+} to continue its steady forward progress. Phase prediction error is weighted by dopaminergic signaling of state precision V_t^{-1} through VTA.

 $_{621}$ This influence is opposed by a "phase prediction error" signal within MPC that pulls μ to progress steadily

at tempo θ . This error signal is weighted by phase precision V^{-1} .

Note that this is not a full, formal error-minimization scheme for implementing PIPPET, which is beyond the scope of this manuscript. In particular, it leaves out an updating scheme for V; see [84] for a discussion of the neurophysiology of precision updating. Further, it does not yet include an appropriate scheme for weighting event prediction error.

Although it would be difficult to directly test this neurophysiological setting of PIPPET in humans, it may be possible to indirectly observe a PIPPET-like process in neural data. At the scalp level and in intracortical electrodes, slow electrical oscillations do seem to anticipatorily track the structure of periodic auditory stimuli [85, 86], and this tracking is associated with the subjective passage of time [87]; these oscillations could be explored as possible estimates of mean underlying phase, with particular focus on those

⁶³² in motor areas. Ideally, timing prediction errors could be observed in the evoked EEG response to events ⁶³³ (the ERP), allowing a direct measurement of event expectancy at each event time, and there are indeed ⁶³⁴ indicators that the ERP is sensitive to temporal predictability (e.g., [88, 89]); however, the sensitivity of the ⁶³⁵ ERP to recent stimulus history makes this approach unpromising. However, timing prediction errors may be ⁶³⁶ observable in EEG/MEG through their effect on gamma oscillations [90, 91]. Further, the subjective hazard ⁶³⁷ rate Λ itself may be observable by using techniques recently applied to decode the temporal hazard function ⁶³⁸ from EEG data [92], or through its correlation with beta oscillations [93].

Although human-like beat-based perceptual and audio-motor entrainment seems to be unique to humans, other primates do show rudimentary rhythmic timing abilities, especially in the visual modality [94], and represent phase of self-generated cyclic behavioral processes in MPC [79, 78]. Experimental paradigms appropriately modified to engage mechanisms of self-action tracking might activate in non-human primates the same mechanisms of uncertainty-informed event-timing-based phase tracking that we hypothesize for auditory rhythm tracking in humans. Thus, primate neurophysiology in MPC and the dopaminergic system may be a promising avenue for indirectly testing the phase inference framework as a description of the human faculty of rhythmic entrainment.

647 5 Acknowledgments

Thanks to Tom Kaplan for extensive discussions and key insights motivating this manuscript, and to
 Aniruddh Patel, Darren Rhodes, and Nori Jacoby for helpful feedback.

650 6 Appendix

651 6.1 The PATIPPET filter

We let $\boldsymbol{\mu} = \begin{pmatrix} \bar{\phi} \\ \bar{\theta} \end{pmatrix}$ denote the posterior mean and $\mathbf{V} = \begin{pmatrix} V^{11} & V^{12} \\ V^{21} & V^{22} \end{pmatrix}$ denote the posterior covariance. The expressions for the evolution of the PATIPPET filter, which we derive in the following section, are:

$$\begin{cases} d\boldsymbol{\mu} = \begin{pmatrix} \bar{\theta} \\ 0 \end{pmatrix} dt + (\hat{\boldsymbol{\mu}} - \boldsymbol{\mu}_t) \cdot (dN_t - \Lambda dt) \\ d\mathbf{V} = \begin{pmatrix} 2V^{12} + \sigma^2 & V^{22} \\ V^{22} & \sigma_{\theta}^2 \end{pmatrix} dt + (\hat{\mathbf{V}} - \mathbf{V}_t) \cdot (dN_t - \Lambda dt) \end{cases}$$
(9)

₆₅₄ where we define

$$\begin{cases} \Lambda := \sum_{i=0,1,\dots} \Lambda_{i} \hat{\theta}_{i} \\ \hat{\mu} = \frac{1}{\Lambda} \sum_{i=0,1,\dots} \Lambda_{i} \begin{pmatrix} K_{i}^{12} + \hat{\phi}_{i} \hat{\theta}_{i} \\ K_{i}^{22} + \hat{\theta}_{i}^{2} \end{pmatrix} \\ \hat{\mathbf{V}} := \frac{1}{\Lambda} \sum_{i=0,1,\dots} \Lambda_{i} \begin{pmatrix} \hat{\theta}_{i} \mathbf{K}_{i} + \hat{\theta}_{i} (\hat{\mu}_{i} - \mu_{t+}) (\hat{\mu}_{i} - \mu_{t+})^{T} \\ + (\hat{\mu}_{i} - \mu_{t+}) \begin{pmatrix} K_{i}^{21} & K_{i}^{22} \end{pmatrix} + \begin{pmatrix} K_{i}^{12} \\ K_{i}^{22} \end{pmatrix} (\hat{\mu}_{i} - \mu_{t+})^{T} \end{pmatrix}$$
(10)

655 and where

656 $\mathbf{K}_0 := \mathbf{V}, \, \mathbf{K}_i := \left(\mathbf{P}_i + \mathbf{V}^{-1}\right)^{-1} \text{ for } i > 0.$

1

$$K_i^{kl}$$
 denotes the entries in \mathbf{K}_i .

$$\begin{aligned} & \overset{658}{\mathbf{h}_{0}} & & \Lambda_{0} := \lambda_{0}, \Lambda_{i} := \lambda_{i} \varphi(\phi_{i} | \bar{\phi}, v_{i}^{-1} + (V^{11})^{-1}) \text{ for } i > 0. \\ & \overset{659}{\mathbf{h}_{i}} & = \begin{pmatrix} \hat{\phi}_{i} \\ \hat{\theta}_{i} \end{pmatrix} := \mathbf{K}_{i} \left(\begin{pmatrix} v_{i}^{-1} \phi_{i} \\ 0 \end{pmatrix} + \mathbf{V}^{-1} \boldsymbol{\mu} \right) \text{ for } i > 0, \text{ and } \hat{\boldsymbol{\mu}}_{0} := \boldsymbol{\mu} \\ & \overset{660}{\mathbf{h}_{i}} & = \begin{pmatrix} v_{i}^{-1} & 0 \\ 0 & 0 \end{pmatrix} \end{aligned}$$

661 6.2 Derivation of differential equations and update equations.

Here we derive the PATIPPET filter; the PIPPET filter can be derived similarly or as a special case of
 PATIPPET.

Snyder [23] provides a partial differential equation describing the evolution of a probability distribution on
 a continuously stochastically evolving state that drives the emission of point process events. If the evolution
 of the underlying state is described by a Gauss-Markov diffusion process:

$$d\mathbf{x} = \mathbf{A}\mathbf{x}dt + \mathbf{B}d\mathbf{W}_t \tag{11}$$

and events are generated at rate $\lambda(\mathbf{x})$, then the evolution of the probability distribution $p_t(\mathbf{x})$ is described by

$$dp_t(\mathbf{x}) = \mathcal{L}[p_t(\mathbf{x})]dt + p_t(\mathbf{x})\left(\frac{\lambda(\mathbf{x})}{\Lambda} - 1\right) \cdot (dN_t - \Lambda dt)$$
(12)

where $\Lambda := \mathbb{E}[\lambda(\mathbf{x})]$ (with \mathbb{E} denoting expectation under distribution $p_t(\mathbf{x})$), dN_t is the increment in the event count over each dt time step (assumed to be either 1 or 0 with probability 1), and \mathcal{L} is the Kolmogorov

⁶⁷¹ forward operator associated with (11):

$$\mathcal{L}[p(\mathbf{x})] = -\sum_{i} \frac{\partial}{\partial x_{i}} [\mathbf{A}\mathbf{x}]_{i} p(\mathbf{x}) + \frac{1}{2} \sum_{i,j} \frac{\partial^{2}}{\partial x_{i} \partial x_{j}} [\mathbf{B}\mathbf{B}^{T}]_{ij} p(\mathbf{x})$$
(13)

Here we project p onto a Gaussian distribution at each time step by matching mean μ and covariance **V**, which is also the projection with minimal KL divergence. We do this by finding the differentials of these moments of p_t and using them to drive the evolution of these two variables:

$$d\boldsymbol{\mu}_{t} = \boldsymbol{\mu}_{t+} - \boldsymbol{\mu}_{t} = \int_{\mathbf{x}} \mathbf{x} p_{t+}(\mathbf{x}) d\mathbf{x} - \int_{\mathbf{x}} \mathbf{x} p_{t}(\mathbf{x}) d\mathbf{x}$$
$$= \int_{\mathbf{x}} \mathbf{x} \left(p_{t+}(\mathbf{x}) - p_{t}(\mathbf{x}) \right) (\mathbf{x}) d\mathbf{x} = \int_{\mathbf{x}} \mathbf{x} dp_{t}(\mathbf{x}) d\mathbf{x}$$
$$= \int_{\mathbf{x}} \mathbf{x} \mathcal{L}[p_{t}(\mathbf{x})] dt d\mathbf{x} + (\hat{\boldsymbol{\mu}} - \boldsymbol{\mu}_{t}) \cdot (dN_{t} - \Lambda dt)$$
(14)

where we define $\hat{\boldsymbol{\mu}} := \mathbb{E} [\mathbf{x} \lambda(\mathbf{x})]$, and

$$d\mathbf{V}_t = \mathbf{V}_{t+} - \mathbf{V}_t = \int_{\mathbf{x}} [[\mathbf{x} - \boldsymbol{\mu}_{t+}]]^2 p_{t+}(\mathbf{x}) d\mathbf{x} - \int_{\mathbf{x}} [[\mathbf{x} - \boldsymbol{\mu}_t]]^2 p_t(\mathbf{x}) d\mathbf{x}$$

where $[[\mathbf{x}]]^2$ denotes $\mathbf{x}\mathbf{x}^T$.

$$d\mathbf{V}_{t} = \int_{\mathbf{x}} [[\mathbf{x} - \boldsymbol{\mu}_{t+1}]]^{2} \left(p_{t+}(\mathbf{x}) - p_{t}(\mathbf{x}) \right) d\mathbf{x}$$

+
$$\int_{\mathbf{x}} \left([[\mathbf{x} - \boldsymbol{\mu}_{t+1}]]^{2} - [[\mathbf{x} - \boldsymbol{\mu}_{t}]]^{2} \right) p_{t}(\mathbf{x}) d\mathbf{x}$$

=
$$\int_{\mathbf{x}} [[\mathbf{x} - \boldsymbol{\mu}_{t+1}]]^{2} dp_{t}(\mathbf{x}) - [[\boldsymbol{\mu}_{t+1} - \boldsymbol{\mu}_{t+1}]]^{2}$$

=
$$\int_{\mathbf{x}} [[\mathbf{x} - \boldsymbol{\mu}_{t+1}]]^{2} \mathcal{L}[p_{t}(\mathbf{x}|N_{t})] dt d\mathbf{x} + \left(\hat{\mathbf{V}} - \mathbf{V}_{t}\right) \cdot (dN_{t} - \Lambda dt)$$
(15)

where we define $\hat{\mathbf{V}} := \mathbb{E}\left[[[\mathbf{x} - \boldsymbol{\mu}_{t+1}]]^2 \lambda(\mathbf{x}) \right].$

Integrating by parts (or following [26]), we can calculate the appropriate integrals of $\mathcal{L}[p_t(\mathbf{x}|N_t)]$, arriving at a general expression for the variational Bayesian filter for point process data:

$$\begin{cases} d\boldsymbol{\mu}_{t} = \mathbf{A}\boldsymbol{\mu}_{t}dt + (\hat{\boldsymbol{\mu}} - \boldsymbol{\mu}_{t}) \cdot (dN_{t} - \Lambda dt) \\ d\mathbf{V}_{t} = (\mathbf{A}\mathbf{V}_{t} + \mathbf{V}_{t}\mathbf{A}^{T} + \mathbf{B}\mathbf{B}^{T})dt + (\hat{\mathbf{V}} - \mathbf{V}_{t}) \cdot (dN_{t} - \Lambda dt) \end{cases}$$
(16)

From (4), the PATIPPET generative model is described by the Gauss-Markov diffusion process (11) with

$$\mathbf{x} = \begin{pmatrix} \phi \\ \theta \end{pmatrix} \text{ and } \boldsymbol{\mu} = \begin{pmatrix} \bar{\phi} \\ \bar{\theta} \end{pmatrix}$$
$$\mathbf{V} = \begin{pmatrix} V^{11} & V^{12} \\ V^{21} & V^{22} \end{pmatrix}$$
$$:= \begin{pmatrix} 0 & 1 \\ 0 & 0 \end{pmatrix} \text{ and } \mathbf{B} := \begin{pmatrix} \sigma & 0 \\ 0 & \sigma_{\theta} \end{pmatrix}$$

677

676

$$\begin{cases} d\boldsymbol{\mu}_{t} = \begin{pmatrix} \bar{\theta} \\ 0 \end{pmatrix} dt + (\hat{\boldsymbol{\mu}} - \boldsymbol{\mu}_{t}) \cdot (dN_{t} - \Lambda dt) \\ d\mathbf{V} = \begin{pmatrix} 2V^{12} + \sigma^{2} & V^{22} \\ V^{22} & \sigma_{\theta}^{2} \end{pmatrix} dt + (\hat{\mathbf{V}} - \mathbf{V}_{t}) \cdot (dN_{t} - \Lambda dt) \end{cases}$$
(17)

We complete the derivation by calculating Λ , $\hat{\mu}$, and $\hat{\mathbf{V}}$. This proceeds by first deriving a simple expression for $p(\mathbf{x})\lambda(\mathbf{x})$ as a sum of scaled normal distributions.

Let $||x||_A^2$ denote $x^T A x$. We will make use of the following result, a generalized form of a well-known result about quadratic forms that allows us to write products of multivariate normal distributions as normal distributions (see [95] for proof and similar application):

$$\|x - a\|_{A}^{2} + \|x - b\|_{B}^{2} = \|a - b\|_{A(A+B)^{-1}B}^{2} + \|x - (A+B)^{-1}(Aa + Bb)\|_{A+B}^{2}$$
(18)

⁶⁸⁴ In the PATIPPET generative model, events are generated at rate

Α

$$\lambda(\mathbf{x}) = \theta \left(\lambda_0 + \sum_{i=1,2,\dots} \frac{\lambda_i}{\sqrt{2\pi v_i}} e^{-\frac{1}{2} \|\mathbf{x} - \mathbf{x}_i\|_{\mathbf{P}_i}^2} \right)$$
$$\mathbf{P}_i = \begin{pmatrix} v_i^{-1} & 0\\ 0 & 0 \end{pmatrix}, \ \mathbf{x}_i = \begin{pmatrix} \phi_i\\ 0 \end{pmatrix}.$$

685

 $p(\mathbf{x})$ is assumed (forced) to be Gaussian, so we can write:

$$p(\mathbf{x}) = \frac{1}{\sqrt{2\pi |\mathbf{V}|}} e^{-\frac{1}{2} \|\mathbf{x} - \boldsymbol{\mu}\|_{\mathbf{V}^{-1}}^2}$$

We calculate:

$$p(\mathbf{x})\lambda(\mathbf{x}) = \frac{\theta}{\sqrt{2\pi|\mathbf{V}|}} e^{-\frac{1}{2}\|\mathbf{x}-\boldsymbol{\mu}\|_{\mathbf{V}^{-1}}^{2}} \left(\lambda_{0} + \sum_{i=1,2,\cdots} \frac{\lambda_{i}}{\sqrt{2\pi v_{i}}} e^{-\frac{1}{2}\|\mathbf{x}-\mathbf{x}_{i}\|_{\mathbf{P}_{i}}^{2}}\right)$$
$$= \frac{\lambda_{0}\theta}{\sqrt{2\pi|\mathbf{V}|}} e^{-\frac{1}{2}\|\mathbf{x}-\boldsymbol{\mu}\|_{\mathbf{V}^{-1}}^{2}} + \theta \sum_{i=1,2,\cdots} \frac{\lambda_{i}}{2\pi\sqrt{v_{i}|\mathbf{V}|}} e^{-\frac{1}{2}\|\mathbf{x}-\mathbf{x}_{i}\|_{\mathbf{P}_{i}}^{2} - \frac{1}{2}\|\mathbf{x}-\boldsymbol{\mu}\|_{\mathbf{V}^{-1}}^{2}}$$

Applying (18),

$$p(\mathbf{x})\lambda(\mathbf{x}) = \frac{\lambda_{0}\theta}{\sqrt{2\pi|\mathbf{V}|}} e^{-\frac{1}{2}\|\mathbf{x}-\boldsymbol{\mu}\|_{\mathbf{V}^{-1}}^{2}} \\ + \theta \sum_{i=1,2,\cdots} \lambda_{i} \left(\frac{1}{\sqrt{2\pi(v_{i}^{-1}+V^{-1})}} e^{-\frac{1}{2}\|\mathbf{x}_{i}-\boldsymbol{\mu}\|_{\mathbf{P}_{i}\mathbf{K}_{i}}^{2}(\mathbf{V}^{1}_{1})^{-1}}\right) \left(\frac{1}{\sqrt{2\pi\frac{v_{i}|\mathbf{V}|}{v_{i}^{-1}+(\mathbf{V}^{1}_{1})^{-1}}}} e^{-\frac{1}{2}\|\mathbf{x}-\mathbf{K}_{i}(\mathbf{P}_{i}\mathbf{x}_{i}+\mathbf{V}^{-1}\boldsymbol{\mu})\|_{\mathbf{K}_{i}^{-1}}^{2}}\right)$$
(19)

where we define $\mathbf{K}_i := (\mathbf{P}_i + \mathbf{V}^{-1})^{-1}$. These two final terms are both expressions for normal distributions, so we can rewrite (19) as

$$p(\mathbf{x})\lambda(\mathbf{x}) = \lambda_0 \theta \varphi(\mathbf{x}|\boldsymbol{\mu}, \mathbf{V}) + \theta \sum_{i=1,2,\cdots} \lambda_i \varphi(\phi_i | \bar{\phi}, v_i^{-1} + (V^{11})^{-1}) \varphi(\mathbf{x} | \mathbf{K}_i(\mathbf{P}_i \mathbf{x}_i + \mathbf{V}^{-1} \boldsymbol{\mu}), \mathbf{K}_i)$$
(20)

We simplify this expression by defining $\Lambda_i := \lambda_i \varphi(\phi_i | \bar{\phi}, v_i^{-1} + (V^{11})^{-1})$ for i > 0, and setting $\Lambda_0 := \lambda_0$ and $\mathbf{K}_0 = \mathbf{V}$. We define $\hat{\boldsymbol{\mu}}_i := \begin{pmatrix} \hat{\phi}_i \\ \hat{\theta}_i \end{pmatrix} := \mathbf{K}_i (\mathbf{P}_i \mathbf{x}_i + \mathbf{V}^{-1} \boldsymbol{\mu})$ for i > 0 and set $\hat{\boldsymbol{\mu}}_0 := \boldsymbol{\mu}$. This lets us write

$$p(\mathbf{x})\lambda(\mathbf{x}) = \sum_{i=0,1,\cdots} \Lambda_i \theta \varphi(\mathbf{x}|\hat{\boldsymbol{\mu}}_i, \mathbf{K}_i)$$
(21)

We use this expression and the moments of normal distributions to calculate Λ , $\hat{\mu}$, and $\hat{\mathbf{V}}$:

$$\Lambda := \mathbb{E}_{p} \left[\lambda(\mathbf{x}) \right] = \sum_{i=0,1,\cdots} \Lambda_{i} \int \theta \varphi(\mathbf{x} | \hat{\boldsymbol{\mu}}_{i}, \mathbf{K}_{i}) d\mathbf{x} = \sum_{i=0,1,\cdots} \Lambda_{i} \hat{\theta}_{i}$$
(22)
$$\hat{\boldsymbol{\mu}} := \frac{1}{\Lambda} \mathbb{E} \left[\mathbf{x} \lambda(\mathbf{x}) \right] = \frac{1}{\Lambda} \sum_{i=0,1,\cdots} \Lambda_{i} \int \begin{pmatrix} \phi \theta \\ \theta^{2} \end{pmatrix} \varphi(\mathbf{x} | \hat{\boldsymbol{\mu}}_{i}, \mathbf{K}_{i}) d\mathbf{x}$$

This expression picks out non-central second moment terms of each normal distributions in (21), each of which can be written in terms of the covariance matrix and mean of the distribution. Using K_i^{kl} to denote the entries in \mathbf{K}_i , we can write

$$\hat{\boldsymbol{\mu}} = \frac{1}{\Lambda} \sum_{i=0,1,\cdots} \Lambda_i \begin{pmatrix} K_i^{12} + \hat{\phi}_i \hat{\theta}_i \\ K_i^{22} + \hat{\theta}_i^2 \end{pmatrix}$$
(23)

The third-order expression for $\hat{\mathbf{V}}$ can also be written in terms of covariance matrices and means since the central third moments of normal distributions are zero.

$$\hat{\mathbf{V}} := \frac{1}{\Lambda} \mathbb{E}_{p} \left[\left[[\mathbf{x} - \boldsymbol{\mu}_{t+1}] \right]^{2} \lambda(\mathbf{x}) \right] \\
= \frac{1}{\Lambda} \sum_{i=0,1,\cdots} \Lambda_{i} \int \left[[\mathbf{x} - \boldsymbol{\mu}_{t+1}] \right]^{2} \theta \varphi(\mathbf{x} | \hat{\boldsymbol{\mu}}_{i}, \mathbf{K}_{i}) d\mathbf{x} \\
= \sum_{i=0,1,\cdots} \Lambda_{i} \left[\hat{\theta}_{i} \int \left[[\mathbf{x} - \hat{\boldsymbol{\mu}}_{i}] \right]^{2} \varphi(\mathbf{x} | \hat{\boldsymbol{\mu}}_{i}, \mathbf{K}_{i}) d\mathbf{x} \cdots \\
+ \hat{\theta}_{i} \left[[\hat{\boldsymbol{\mu}}_{i} - \boldsymbol{\mu}_{t+1}] \right]^{2} \cdots \\
+ \left(\hat{\boldsymbol{\mu}}_{i} - \boldsymbol{\mu}_{t+1} \right) \int (\mathbf{x} - \hat{\boldsymbol{\mu}}_{i})^{T} (\theta - \hat{\theta}_{i}) \varphi(\mathbf{x} | \hat{\boldsymbol{\mu}}_{i}, \mathbf{K}_{i}) d\mathbf{x} \cdots \\
+ \left(\int (\mathbf{x} - \hat{\boldsymbol{\mu}}_{i}) (\theta - \hat{\theta}_{i}) \varphi(\mathbf{x} | \hat{\boldsymbol{\mu}}_{i}, \mathbf{K}_{i}) d\mathbf{x} \right) \left(\hat{\boldsymbol{\mu}}_{i} - \boldsymbol{\mu}_{t+1} \right)^{T} \right] \\
= \frac{1}{\Lambda} \sum_{i=0,1,\cdots} \Lambda_{i} \left[\hat{\theta}_{i} \mathbf{K}_{i} + \hat{\theta}_{i} \left[[\hat{\boldsymbol{\mu}}_{i} - \boldsymbol{\mu}_{t+1}] \right]^{2} \cdots \\
+ \left(\hat{\boldsymbol{\mu}}_{i} - \boldsymbol{\mu}_{t+1} \right) \left(K_{i}^{21} \quad K_{i}^{22} \right) + \left(K_{i}^{12} \\
K_{i}^{22} \right) \left(\hat{\boldsymbol{\mu}}_{i} - \boldsymbol{\mu}_{t+1} \right)^{T} \right] \tag{24}$$

Expressions (22), (23), and (24) coupled with (17) constitute the PATIPPET filter.

The PIPPET filter can be derived as a special case of the PATIPPET filter by setting $\sigma_{\theta} = 0, \theta_0 = 1$, and all terms in **V** to zero except *V*. However, this requires finessing various degeneracies, e.g. wherever **V** is inverted. More straightforward is to follow the same process as above, starting from the PIPPET generative model (1) and (2). Either way ultimately yields the PIPPET filter (3).

⁶⁹² For multiple event streams j,:

$$dp_t(\mathbf{x}) = \mathcal{L}[p_t(\mathbf{x})]dt + p_t(\mathbf{x})\sum_j \left(\lambda_j(\mathbf{x}) - \mathbb{E}_p[\lambda_j(\mathbf{x})]\right) \cdot \left(\mathbb{E}_p[\lambda_j(\mathbf{x})]^{-1}dN_j - dt\right)$$
(25)

This follows directly from application of the derivation above to equation (5) in [96] with a discrete spatial dimension. By the methods above, it yields the mPIPPET filter (8) and the mPATIPPET filter:

$$\begin{cases} d\boldsymbol{\mu}_{t} = \begin{pmatrix} \bar{\theta} \\ 0 \end{pmatrix} dt + \sum_{j} \left(\hat{\boldsymbol{\mu}}^{j} - \boldsymbol{\mu}_{t} \right) \cdot \left(dN_{t}^{j} - \Lambda^{j} dt \right) \\ d\mathbf{V} = \begin{pmatrix} 2V^{12} + \sigma^{2} & V^{22} \\ V^{22} & \sigma_{\theta}^{2} \end{pmatrix} dt + \sum_{j} \left(\hat{\mathbf{V}}^{j} - \mathbf{V}_{t} \right) \cdot \left(dN_{t}^{j} - \Lambda^{j} dt \right) \end{cases}$$
(26)

695 6.3 Simulation parameters.

All code used to create figures in this manuscript is available at https://github.com/joncannon/PIPPET. PIPPET simulations were conducted by numerical simulation of (3) with dt = .001 and initialized with $\phi_0 = 0$ and $V_0 = .0002$. Parameters for the simulations shown in each figure are listed below, with t_n used to denote simulated event times (in units of seconds).

Figure 2:
$$\phi_1 = .5, v_1 = .0005, \lambda_1 = 1, \mu_t = .43, V_t = .001, \lambda_0 = 0$$
 or .5, except as otherwise specified.
Figure 3A:

$$\begin{split} \{t_n\} =& \{0, .150, .5, .75, .9, 1.25\} \\ \{\phi_i\} =& \{0, .15, .25, .4, .5, .65, .75, .9, 1, 1.15, 1.25, 1.4\} \\ \{v_i\} =& \{.0001, .0005, .0001, .0005, .0001, .0005, .0001, .0005, .0001, .0005\} \\ \{\lambda_i\} =& \{.05, .01, .05, .01, .05, .01, .05, .01, .05, .01\} \\ \lambda_0 =& .01 \\ \sigma =& .05 \end{split}$$

- Figure 3B: Same as Figure 3A, but with $t_3 = .45$ (50 ms negative event time shift).
- Figure 3C: Same as Figure 3A, but with $\{t_n\} = \{0, .15, .5, .7, .85, 1.2\}$ (50 ms negative phase shift).
- Figure 4A: Same as Figure 3, but with $\{t_n\} = \{0, .150, .65, .9, 1.15, 1.25\}.$
- Figure 4B: Same as Figure 4A, but with $\sigma = .3$.

Figure 4A: Same as Figure 4A, but with additional tap times and tap feedback expectations:

$$\begin{split} \{t_n^{tap}\} = & \{\phi_i^{tap}\} = \{0, .5, 1\} \\ v_i^{tap} = & .0005 \\ \lambda_i^{tap} = & .05 \\ \lambda_0^{tap} = & .01 \end{split}$$

PATIPPET simulations were conducted by numerical simulation of (4) with dt = .001. Parameters for the simulations shown in each figure are listed below.

Figure 5:

$$t_n = \frac{n}{1.2Hz}$$

$$\phi_i = i$$

$$v_i = .005$$

$$\lambda_i = .02$$

$$\lambda_0 = .0001$$

$$\sigma = .05$$

$$\sigma_\theta = .05$$

$$\mu_0 = \begin{pmatrix} 0\\ 1 \end{pmatrix}$$

$$V_0 = \begin{pmatrix} .001 & 0\\ 0 & .04 \end{pmatrix}$$

Figure 6: In four simulations, we set the inter-onset interval Δ to .4s, 0,7s, 1.0s, and 1.3s. In each

simulation, we set the perturbation δ to $\frac{\Delta}{25}.$

$$\{t_n\} = \{\Delta, 2\Delta, 3\Delta, 4\Delta + \delta\}$$

$$\phi_i = i$$

$$v_i = .0002$$

$$\lambda_i = .02$$

$$\lambda_0 = 10^{-5}$$

$$\sigma = .01$$

$$\sigma_{\theta} = .01$$

$$\mu_0 = \begin{pmatrix} 0 \\ 1 \end{pmatrix}$$

$$\mathbf{V}_0 = \begin{pmatrix} 10^{-4} & 0 \\ 0 & 10^{-4} \end{pmatrix}$$

$$\phi_i = .25i$$

$$v_i = .0001$$

$$\lambda_i = 1$$

$$\lambda_0 = .0001$$

$$\sigma = .015$$

$$\sigma_\theta = .2$$

$$\mu_0 = \begin{pmatrix} 0\\ 1 \end{pmatrix}$$

$$\mathbf{V}_0 = \begin{pmatrix} .0001 & 0\\ 0 & .005 \end{pmatrix}$$

⁷⁰⁷ Left: $\{t_n\} = \{.25, .5, .75, 1\}$. Right: $\{t_n\} = \{1\}$.

Figure 7B: Same as Figure 7A, but with $\lambda_i = 0$ and $\lambda_0 = 4$.

709 References

- Repp BH and Su YH. Sensorimotor synchronization: A review of recent research (2006-2012). Psychonomic Bulletin and Review 2013; 20:403-52. DOI: 10.3758/s13423-012-0371-2. arXiv: NIHMS150003
- 2. Merchant H, Grahn J, Trainor L, Rohrmeier M, and Fitch WT. Finding the beat: a neural perspective
- across humans and non-human primates. Philosophical transactions of the Royal Society of London.
- Series B, Biological sciences 2015; 370. DOI: 10.1098/rstb.2014.0093. Available from: http:
- //www.ncbi.nlm.nih.gov/pubmed/25646516
- Obleser J and Kayser C. Neural Entrainment and Attentional Selection in the Listening Brain. Trends
 in Cognitive Sciences 2019; 23:1–14. DOI: 10.1016/j.tics.2019.08.004. Available from: https:
 //doi.org/10.1016/j.tics.2019.08.004
- Lawrance ELA, Harper NS, Cooke JE, and Schnupp JWH. Temporal predictability enhances auditory
 detection. The Journal of the Acoustical Society of America 2014; 135:EL357–EL363. DOI: 10.1121/
 1.4879667. Available from: http://dx.doi.org/10.1121/1.4879667
- Nobre AC and Van Ede F. Anticipated moments: Temporal structure in attention. Nature Reviews
 Neuroscience 2018; 19:34-48. DOI: 10.1038/nrn.2017.141. Available from: http://dx.doi.org/10.
 1038/nrn.2017.141
- Morillon B, Schroeder CE, Wyart V, and Arnal LH. Temporal prediction in lieu of periodic stimulation.
 Journal of Neuroscience 2016; 36:2342–7. DOI: 10.1523/JNEUROSCI.0836-15.2016
- 727 7. Lange K. Brain correlates of early auditory processing are attenuated by expectations for time and
 728 pitch. Brain and Cognition 2009; 69:127-37. DOI: 10.1016/j.bandc.2008.06.004. Available from:
 729 http://dx.doi.org/10.1016/j.bandc.2008.06.004
- Jazayeri M and Shadlen MN. Temporal context calibrates interval timing. Nature Neuroscience 2010;
 13:1020-6. DOI: 10.1038/nn.2590
- 9. Herrmann B, Henry MJ, Haegens S, and Obleser J. Temporal expectations and neural amplitude
 fluctuations in auditory cortex interactively influence perception. NeuroImage 2016; 124:487–97. DOI:
 10.1016/j.neuroimage.2015.09.019
- Rajendran VG, Teki S, and Schnupp JW. Temporal Processing in Audition: Insights from Music.
 Neuroscience 2018; 389:4–18. DOI: 10.1016/j.neuroscience.2017.10.041. Available from: https:
 //doi.org/10.1016/j.neuroscience.2017.10.041
- Large EW and Jones MR. The dynamics of attending: How people track time-varying events. Psychological Review 1999; 106:119–59. DOI: 10.1037//0033-295x.106.1.119

- Large EW and Palmer C. Perceiving temporal regularity in music. Cognitive Science 2002; 26:1–37.
 DOI: 10.1016/S0364-0213(01)00057-X
- Friston K. A theory of cortical responses. Philosophical Transactions of the Royal Society B: Biological
 Sciences 2005; 360:815–36. DOI: 10.1098/rstb.2005.1622
- Vuust P and Witek MA. Rhythmic complexity and predictive coding: A novel approach to modeling
 rhythm and meter perception in music. Frontiers in Psychology 2014; 5:1–14. DOI: 10.3389/fpsyg.
 2014.01111
- ⁷⁴⁷ 15. Vuust P, Dietz MJ, Witek M, and Kringelbach ML. Now you hear it: A predictive coding model for
 ⁷⁴⁸ understanding rhythmic incongruity. Annals of the New York Academy of Sciences 2018; 1423:19–29.
 ⁷⁴⁹ DOI: 10.1111/nyas.13622
- Proksch S, Comstock DC, Médé B, Pabst A, and Balasubramaniam R. Motor and Predictive Processes
 in Auditory Beat and Rhythm Perception. 2020; 14. DOI: 10.3389/fnhum.2020.578546
- ⁷⁵² 17. Friston K, Stephan K, Li B, and Daunizeau J. Generalised filtering. Mathematical Problems in Engi ⁷⁵³ neering 2010; 2010. DOI: 10.1155/2010/621670
- Buckley CL, Kim CS, McGregor S, and Seth AK. The free energy principle for action and perception:
 A mathematical review. Journal of Mathematical Psychology 2017; 81:55–79. DOI: 10.1016/j.jmp.
 2017.09.004. arXiv: 1705.09156. Available from: http://dx.doi.org/10.1016/j.jmp.2017.09.004
- ⁷⁵⁷ 19. Schwartze M and Kotz SA. A dual-pathway neural architecture for specific temporal prediction. Neu-
- roscience and Biobehavioral Reviews 2013; 37:2587–96. DOI: 10.1016/j.neubiorev.2013.08.005.
- Available from: http://dx.doi.org/10.1016/j.neubiorev.2013.08.005
- Egger SW and Jazayeri M. A nonlinear updating algorithm captures suboptimal inference in the pres ence of signal-dependent noise. Scientific Reports 2018 :18–20. DOI: 10.1038/s41598-018-30722-0
- ⁷⁶² 21. DI Luca M and Rhodes D. Optimal Perceived Timing: Integrating Sensory Information with Dynami ⁷⁶³ cally Updated Expectations. Scientific Reports 2016; 6:1–15. DOI: 10.1038/srep28563
- ⁷⁶⁴ 22. Elliott MT, Wing AM, and Welchman AE. Moving in time: Bayesian causal inference explains movement
 ⁷⁶⁵ coordination to auditory beats. Proceedings of the Royal Society B: Biological Sciences 2014; 281. DOI:
 ⁷⁶⁶ 10.1098/rspb.2014.0751
- ⁷⁶⁷ 23. Snyder DL. Filtering and Detection for Doubly Stochastic Poisson Processes. IEEE Transactions on
 ⁷⁶⁸ Information Theory 1972; 18:91–102. DOI: 10.1109/TIT.1972.1054756
- ⁷⁶⁹ 24. Opper M. A Bayesian Approach to On-line Learning. On-Line Learning in Neural Networks 2010
 ⁷⁷⁰ :363-78. DOI: 10.1017/cbo9780511569920.017

- 771 25. Friston K. The free-energy principle: A unified brain theory? Nature Reviews Neuroscience 2010;
 772 11:127–38. DOI: 10.1038/nrn2787
- ⁷⁷³ 26. Eden UT and Brown EN. Continuous-time filters for state estimation from point-process models of
 ⁷⁷⁴ neural data. Statistica Sinica 2008; 18:1293–310
- 775 27. Cemgil AT, Kappen B, Desain P, and Honing H. On tempo tracking: Tempogram representation and
- 776
 Kalman filtering. Journal of New Music Research 2000; 29:259–73. DOI: 10.1080/09298210008565462
- London J, Polak R, and Jacoby N. Rhythm histograms and musical meter: A corpus study of Malian
 percussion music. Psychonomic Bulletin and Review 2017; 24:474–80. DOI: 10.3758/s13423-0161093-7
- Polak R, London J, and Jacoby N. Both isochronous and non-isochronous metrical subdivision afford
 precise and stable ensemble entrainment: A corpus study of malian jembe drumming. Frontiers in
 Neuroscience 2016; 10:1–11. DOI: 10.3389/fnins.2016.00285
- ⁷⁸³ 30. Friberg A and Sundström A. Swing Ratios and Ensemble Timing in Jazz Performance: Evidence for a
 ⁷⁸⁴ Common Rhythmic Pattern. Music Perception 2002; 19:333–49. DOI: 10.1525/mp.2002.19.3.333
- Warren RM and Gregory RL. An Auditory Analogue of the Visual Reversible Figure. The American
 Journal of Psychology 1958; 71:612–3
- 787 32. Fitch WT and Rosenfeld AJ. Perception and Production of Syncopated Rhythms. Music Perception
 2007; 25:43–58
- Repp BH. Tapping in synchrony with a perturbed metronome: The phase correction response to small
 and large phase shifts as a function of tempo. Journal of Motor Behavior 2011; 43:213–27. DOI: 10.
 1080/00222895.2011.561377
- Repp BH, Keller PE, and Jacoby N. Quantifying phase correction in sensorimotor synchronization:
 Empirical comparison of three paradigms. Acta Psychologica 2012; 139:281–90. DOI: 10.1016/j.
 actpsy.2011.11.002. Available from: http://dx.doi.org/10.1016/j.actpsy.2011.11.002
- 795 35. Hall GS and Jastrow J. Studies of Rhythm. Mind 1886 Jan; os-XI:55-62. DOI: 10.1093/mind/os-
- 796 XI.41.55. eprint: https://academic.oup.com/mind/article-pdf/os-XI/41/55/9358438/os-
- 797 XI_41_55.pdf. Available from: https://doi.org/10.1093/mind/os-XI.41.55
- ⁷⁹⁸ 36. Nakajima Y. A psychophysical investigation of divided time intervals shown by sound bursts. Journal
 ⁷⁹⁹ of the Acoustical Society of Japan 1979; 35:145–51
- 37. Meumann E. Beiträge zur Psychologie des Zeitbewußtseins [contributions to the psychology of time
 consciousness]. Philosophische Studien 1896; 12:128–254

802	38.	Grimm K. der einfluß der Zeitform auf die Wahrnehmung der Zeitdauer [the influence of time-form on
803		the perception of duration]. Zeitschrift für Psychologie 1934; 132:104–32
804	39.	Repp BH and Bruttomesso M. A filled duration illusion in music: Effects of metrical subdivision on

- the perception and production of beat tempo. Advances in Cognitive Psychology 2009; 5:114–34. DOI:
 10.2478/V10053-008-0071-7
- ⁸⁰⁷ 40. Repp B and Jendoubi H. Flexibility of temporal expectations for triple subdivision of a beat. Advances
 ⁸⁰⁸ in Cognitive Psychology 2009; 5:27–41. DOI: 10.2478/v10053-008-0063-7
- 41. Wohlschläger A and Koch R. Synchronization error: An error in time perception. *Rhythm perception and production.* Ed. by Desain P and Winsdor L. Swets:115–27

42. Wing AM and Kristofferson AB. Response delays and the timing of discrete motor responses. Perception
& Psychophysics 1973; 14:5–12. DOI: 10.3758/BF03198607

43. Mates J. A model of synchronization of motor acts to a stimulus sequence - II. Stability analysis, error
estimation and simulations. Biological Cybernetics 1994; 70:475–84. DOI: 10.1007/BF00203240

- ⁸¹⁵ 44. Breska A and Deouell LY. Neural mechanisms of rhythm-based temporal prediction: Delta phase⁸¹⁶ locking reflects temporal predictability but not rhythmic entrainment. PLoS Biology 2017; 15:1–30.
 ⁸¹⁷ DOI: 10.1371/journal.pbio.2001665
- ⁸¹⁸ 45. Bouwer FL, Honing H, and Slagter HA. Beat-based and memory-based temporal expectations in
 ⁸¹⁹ rhythm: similar perceptual effects, different underlying mechanisms. 2019; 8:55
- 46. Fox C, Rezek I, and Roberts S. Drum 'N 'Bayes : on-Line Variational Inference for Beat Tracking
 and Rhythm Recognition. International Computer Music Conference 2007. DOI: 10.1016/j.chieco.
 2016.10.003
- 47. Pesek M, Leonardis A, and Marolt M. An Analysis of Rhythmic Patterns with Unsupervised Learning.
 Applied Sciences 2019. DOI: 10.3390/app10010178
- 48. Ma WJ and Jazayeri M. Neural coding of uncertainty and probability. Annual Review of Neuroscience
 2014; 37:205–20. DOI: 10.1146/annurev-neuro-071013-014017
- ⁸²⁷ 49. Repp BH and Keller PE. Adaptation to tempo changes in sensorimotor synchronization: Effects of
 ⁸²⁸ intention, attention, and awareness. Quarterly Journal of Experimental Psychology Section A: Human
 ⁸²⁹ Experimental Psychology 2004; 57:499–521. DOI: 10.1080/02724980343000369
- 50. Danielsen A. Here, There, and Everywhere: three accounts of pulse in D'Angelo's 'Left and Right'.
 2010 Jan :19-36. DOI: 10.4324/9781315596983-2

- ⁸³² 51. Witek MA, Clarke EF, Kringelbach ML, and Vuust P. Effects of Polyphonic Context, Instrumentation,
 ⁸³³ and Metrical Location on Syncopation in Music. Music Perception 2014; 32:201–17
- ⁸³⁴ 52. Rauschecker JP. Where, When, and How: Are they all sensorimotor? Towards a unified view of the
 ⁸³⁵ dorsal pathway in vision and audition. Cortex 2018; 98:262-8. DOI: 10.1016/j.cortex.2017.10.020.
 ⁸³⁶ Available from: https://doi.org/10.1016/j.cortex.2017.10.020
- 53. Comstock DC, Hove MJ, and Balasubramaniam R. Sensorimotor synchronization with auditory and
 visual modalities: Behavioral and neural differences. Frontiers in Computational Neuroscience 2018;
 12:1–8. DOI: 10.3389/fncom.2018.00053
- ⁸⁴⁰ 54. Hove MJ, Marie C, Bruce IC, and Trainor LJ. Superior time perception for lower musical pitch explains
 ⁸⁴¹ why bass-ranged instruments lay down musical rhythms. Proceedings of the National Academy of
 ⁸⁴² Sciences of the United States of America 2014; 111:10383–8. DOI: 10.1073/pnas.1402039111
- ⁸⁴³ 55. Lenc T, Keller PE, Varlet M, and Nozaradan S. Neural tracking of the musical beat is enhanced by
 ⁸⁴⁴ low-frequency sounds. Proceedings of the National Academy of Sciences of the United States of America
 ⁸⁴⁵ 2018; 115:8221–6. DOI: 10.1073/pnas.1801421115
- 56. Repp BH. Phase Correction, Phase Resetting, and Phase Shifts After Subliminal Timing Perturbations in Sensorimotor Synchronization. Journal of Experimental Psychology: Human Perception and
 Performance 2001; 27:600-21. DOI: 10.1037//0096-1523.27.3.600
- ⁸⁴⁹ 57. Heggli OA, Cabral J, Konvalinka I, Vuust P, and Kringelbach ML. A Kuramoto model of self-other
 ⁸⁵⁰ integration across interpersonal synchronization strategies. PLoS Computational Biology 2019; 15:1–
 ⁸⁵¹ 17. DOI: 10.1371/journal.pcbi.1007422
- ⁸⁵² 58. Koban L, Ramamoorthy A, and Konvalinka I. Why do we fall into sync with others? Interpersonal
 ⁸⁵³ synchronization and the brain's optimization principle. Social Neuroscience 2019; 14:1–9
- Rimmele JM, Morillon B, Poeppel D, and Arnal LH. Proactive Sensing of Periodic and Aperiodic
 Auditory Patterns. Trends in Cognitive Sciences 2018; 22:870-82. DOI: 10.1016/j.tics.2018.08.003.
 Available from: https://doi.org/10.1016/j.tics.2018.08.003
- ⁸⁵⁷ 60. Rohrmeier M. Towards a formalization of musical rhythm. Proc. of the 21st Int. Society for Music
 ⁸⁵⁸ Information Retrieval Conf. 2020
- ⁸⁵⁹ 61. Pearce MT. The construction and evaluation of statistical models of melodic structure in music per⁸⁶⁰ ception and composition. PhD thesis. City University, London, 2005

- ⁸⁶¹ 62. Sioros G, Davies ME, and Guedes C. A generative model for the characterization of musical rhythms.
- ⁸⁶² Journal of New Music Research 2018; 47:114–28. DOI: 10.1080/09298215.2017.1409769. Available ⁸⁶³ from: http://doi.org/10.1080/09298215.2017.1409769
- Repp BH. Obligatory "expectations" of expressive timing induced by perception of musical structure.
 Psychological Research 1998; 61:33–43. DOI: 10.1007/s004260050011
- ⁸⁶⁶ 64. Repp BH. Compensation for subliminal timing perturbations in perceptual-motor synchronization.
 ⁸⁶⁷ Psychological Research 2000; 63:106–28. DOI: 10.1007/PL00008170
- 65. Schwartze M and Kotz SA. The Timing of Regular Sequences: Production, Perception, and Covariation.
- Journal of Cognitive Neuroscience 2015; 27:139. DOI: 10.1162/jocn. Available from: https://www.
- apa.org/ptsd-guideline/ptsd.pdf%7B%5C%%7D0Ahttps://www.apa.org/about/offices/ directorates/guidelines/ptsd.pdf
- 66. Chauvigné LaS, Gitau KM, and Brown S. The neural basis of audiomotor entrainment: an ALE meta analysis. Frontiers in human neuroscience 2014 Jan; 8:776. DOI: 10.3389/fnhum.2014.00776.
 Available from: http://www.pubmedcentral.nih.gov/articlerender.fcgi?artid=4179708%7B%
 5C&%7Dtool=pmcentrez%7B%5C&%7Drendertype=abstract
- Kneissler J, Drugowitsch J, Friston K, and Butz MV. Simultaneous learning and filtering without delusions: A bayes-optimal combination of predictive inference and adaptive filtering. Frontiers in Computational Neuroscience 2015; 9:1–12. DOI: 10.3389/fncom.2015.00047
- ⁸⁷⁹ 68. Weij B van der, Pearce MT, and Honing H. A probabilistic model of meter perception: Simulating
 enculturation. Frontiers in Psychology 2017; 8:1–18. DOI: 10.3389/fpsyg.2017.00824
- Alejandro M, Id M, Sigman M, and Slezak DF. From beat tracking to beat expectation : Cognitive based beat tracking for capturing pulse clarity through time. PLoS ONE 2020; 15:e0242207. DOI:
 10.17605/0SF.IO/P3QTV
- ⁸⁸⁴ 70. Large EW, Almonte FV, and Velasco MJ. A canonical model for gradient frequency neural networks.
 ⁸⁸⁵ Physica D: Nonlinear Phenomena 2010; 239:905–11. DOI: 10.1016/j.physd.2009.11.015. Available
 ⁸⁸⁶ from: http://dx.doi.org/10.1016/j.physd.2009.11.015
- Pouget A, Beck JM, Ma WJ, and Latham PE. Probabilistic brains: Knowns and unknowns. Nature
 Neuroscience 2013; 16:1170–8. DOI: 10.1038/nn.3495
- 72. Gershman SJ and Uchida N. Believing in dopamine. Nature Reviews Neuroscience 2019; 20:703–14. DOI:
- ⁸⁹⁰ 10.1038/s41583-019-0220-7. Available from: http://dx.doi.org/10.1038/s41583-019-0220-7

891	73.	Sarno S, De Lafuente V, Romo R, and Parga N. Dopamine reward prediction error signal codes the
892		temporal evaluation of a perceptual decision report. Proceedings of the National Academy of Sciences
893		of the United States of America 2017; 114:E10494–E10503. DOI: 10.1073/pnas.1712479114

- ⁸⁹⁴ 74. Tomassini A, Ruge D, Galea JM, Penny W, and Bestmann S. The Role of Dopamine in Temporal Un-
- certainty. Journal of Cognitive Neuroscience 2016. DOI: 10.1162/jocn. arXiv: 1511.04103. Available
- from: http://dx.doi.org/10.1162/jocn%7B%5C_%7Da%7B%5C_%7D00409%7B%5C%%7D5Cnhttp:
- ⁸⁹⁷ //www.mitpressjournals.org/doi/abs/10.1162/jocn%7B%5C_%7Da%7B%5C_%7D00409
- 75. Friston KJ, Shiner T, FitzGerald T, Galea JM, Adams R, Brown H, Dolan RJ, Moran R, Stephan KE,
- and Bestmann S. Dopamine, affordance and active inference. PLoS Computational Biology 2012; 8.
 DOI: 10.1371/journal.pcbi.1002327
- 76. Cannon J and Patel AD. How beat perception coopts motor neurophysiology: a proposal. bioRxiv 2020.
 DOI: https://doi.org/10.1101/805838
- ⁹⁰³ 77. Wang J, Narain D, Hosseini EA, and Jazayeri M. Flexible timing by temporal scaling of cortical
 responses. Nature Neuroscience 2018; 21:102–12. DOI: 10.1038/s41593-017-0028-6. Available from:
 http://dx.doi.org/10.1038/s41593-017-0028-6
- ⁹⁰⁶ 78. Gámez J, Mendoza G, Prado L, Betancourt A, and Merchant H. The amplitude in periodic neural state
 ⁹⁰⁷ trajectories underlies the tempo of rhythmic tapping. PLoS biology 2019; 17:e3000054
- ⁹⁰⁸ 79. Russo AA, Khajeh R, Bittner SR, Perkins SM, Cunningham JP, Abbott LF, and Churchland MM.
 ⁹⁰⁹ Neural trajectories in the supplementary motor area and primary motor cortex exhibit distinct ge⁹¹⁰ ometries, compatible with different classes of computation. Neuron 2020; 107. DOI: 10.1101/650002.
 ⁹¹¹ Available from: https://www.biorxiv.org/content/10.1101/650002v1.abstract
- 80. Patel AD and Iversen JR. The evolutionary neuroscience of musical beat perception: the Action Simulation for Auditory Prediction (ASAP) hypothesis. Frontiers in Systems Neuroscience 2014; 8:1–14.
 ⁹¹⁴ DOI: 10.3389/fnsys.2014.00057. Available from: http://journal.frontiersin.org/article/10.
- 915 3389/fnsys.2014.00057/abstract
- ⁹¹⁶ 81. Friston K. Hierarchical models in the brain. PLoS Computational Biology 2008; 4. DOI: 10.1371/
 ⁹¹⁷ journal.pcbi.1000211
- ⁹¹⁸ 82. Schubotz RI. Prediction of external events with our motor system: towards a new framework. Trends
 ⁹¹⁹ in Cognitive Sciences 2007; 11:211-8. DOI: 10.1016/j.tics.2007.02.006

- 84. Kanai R, Komura Y, Shipp S, and Friston K. Cerebral hierarchies: Predictive processing, precision and
 the pulvinar. Philosophical Transactions of the Royal Society B: Biological Sciences 2015; 370. DOI:
 10.1098/rstb.2014.0169
- 85. Schroeder CE and Lakatos P. Low-frequency neuronal oscillations as instruments of sensory selection.
 Trends in neurosciences 2009; 32. DOI: 10.1016/j.tins.2008.09.012.Low-frequency
- 86. Arnal LH and Giraud AL. Cortical oscillations and sensory predictions. Trends in Cognitive Sciences
 2012; 16:390-8. DOI: 10.1016/j.tics.2012.05.003. Available from: http://dx.doi.org/10.1016/
 j.tics.2012.05.003
- 87. Arnal LH and Kleinschmidt AK. Entrained delta oscillations reflect the subjective tracking of time.
 Cerebral Cortex 2017 :e1349583. DOI: 10.1093/cercor/bhu103
- 88. Schwartze M, Farrugia N, and Kotz SA. Dissociation of formal and temporal predictability in early
 auditory evoked potentials. Neuropsychologia 2013; 51:320–5. DOI: 10.1016/j.neuropsychologia.

⁹³⁵ 2012.09.037. Available from: http://dx.doi.org/10.1016/j.neuropsychologia.2012.09.037

⁹³⁶ 89. Ungan P, Karsilar H, and Yagcioglu S. Pre-attentive Mismatch Response and Involuntary Attention
⁹³⁷ Switching to a Deviance in an Earlier-Than-Usual Auditory Stimulus: An ERP Study. Frontiers in
⁹³⁸ Human Neuroscience 2019; 13:1–16. DOI: 10.3389/fnhum.2019.00058

- 939 90. Todorovic A, Ede F van, Maris E, and Lange FP de. Prior expectation mediates neural adaptation to
 repeated sounds in the auditory cortex: An MEG study. Journal of Neuroscience 2011; 31:9118–23.
 941 DOI: 10.1523/JNEUROSCI.1425-11.2011
- 942 91. Bastos AM, Usrey WM, Adams Ra, Mangun GR, Fries P, and Friston KJ. Canonical microcircuits for
 predictive coding. Neuron 2012 Nov; 76:695-711. DOI: 10.1016/j.neuron.2012.10.038. Available
 from: http://www.pubmedcentral.nih.gov/articlerender.fcgi?artid=3777738%7B%5C&%7Dtool=
 pmcentrez%7B%5C&%7Drendertype=abstract
- ⁹⁴⁶ 92. Herbst SK, Fiedler L, and Obleser J. Tracking temporal hazard in the human electroencephalogram
 ⁹⁴⁷ using a forward encoding model. eNeuro 2018; 5:1–17. DOI: 10.1523/ENEURO.0017-18.2018
- 93. Tavano A, Schröger E, and Kotz SA. Beta power encodes contextual estimates of temporal event
 probability in the human brain. PLoS ONE 2019; 14. DOI: 10.1371/journal.pone.0222420

46

Rauschecker JP. An expanded role for the dorsal auditory pathway in sensorimotor control and integration. Hearing Research 2011; 271:16-25. DOI: 10.1016/j.heares.2010.09.001. Available from:
 http://dx.doi.org/10.1016/j.heares.2010.09.001

950	94.	Merchant H and Honing H. Are non-human primates capable of rhythmic entrainment? Evidence for
951		the gradual audiomotor evolution hypothesis. Frontiers in neuroscience 2014 Jan; 7:274. DOI: 10.
952		3389/fnins.2013.00274. Available from: http://www.pubmedcentral.nih.gov/articlerender.
953		fcgi?artid=3894452%7B%5C&%7Dtool=pmcentrez%7B%5C&%7Drendertype=abstract
954	95.	Harel Y, Meir R, and Opper M. A tractable approximation to optimal point process filtering: Applica-
955		tion to neural encoding. Advances in Neural Information Processing Systems 2015; 2015-Janua:1603–
956		11
957	96.	Snyder DL and Fishman P. How to track a swarm of fireflies by observing their flashes. IEEE Trans-

actions on Information Theory 1975; 21