Gain modulation and balanced synaptic input in a conductance-based neural model

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Abstract

Gain modulation of neural responses by the balanced component of the synaptic input is analyzed in the gaussian approximation using a single compartment conductance-based neural model. The model is analyzed in the "normal operating regime", in which the output spiking-rate of the neuron is equal to the spontaneous spiking-rate in the absence of any stimulus. The gain in response to both additional excitatory synaptic input and injected current is found to be modulated in a non-linear way by the level of balanced synaptic input.

Key words: Gain modulation, integrate-and-fire neurons, gaussian approximation, synaptic input, reversal potential

1 Introduction

Gain modulation is a change in the amplitude of the response that a neuron generates due to one input (the modulatory one), but which does not affect the receptive field characteristics (or selectivity) of the neuron. It is a non-linear mechanism by which information is combined (or integrated) from within a single pathway or between different pathways of neural processing, which may be of sensory, motor or cognitive origin. Gain modulation provides a computational means by which neural systems may transform, combine or compare the representations that they carry of the physical world and by which they more accurately control (i.e. modulate) their output. Gain modulation plays an important role in sensory-motor integration, such as eye and reaching movements, and in spatial perception, as well as in auditory masking, attentional processing, object recognition and navigation. Gain modulation is one of the

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few neural computational principles that pervades neural processing, since it plays an important role in a number of different modalities (sensory, motor and cognitive), different brain areas (cortical and sub-cortical), and different neural functions [9].

Although experimental studies have shown gain modulation to play an important role in neural processing, our understanding of the underlying biophysical mechanisms by which neural systems implement gain modulation is lacking. The central question is: "How do neurons achieve the non-linear, multiplicative behavior characteristic of gain modulation, when their inputoutput relationship is basically integrative?". A number of possible different mechanisms have been proposed [9], although they are not necessarily exclusive. One suggested mechanism involves the effect of non-linear interactions in the dendritic processing of neurons [7]. Another approach involves the effect of recurrent connections between neurons, which may give rise to non-linear interactions [8,5]. A third approach is through correlations in the synaptic input [10]. A fourth approach is that addressed in this paper, namely the effect of the balanced component of the synaptic input in providing the non-linear gain modulation of the neural response [4]. The various mechanisms are, of course, by no means mutually exclusive, e.g., the balanced component of the input could partly arise via feedback from recurrent connections.

This paper examines how balanced synaptic input can modulate neuronal gain. This modulation is a result of the effect of the balanced synaptic input upon the variance of the synaptic current, which can be varied independently of the mean synaptic current (which is determined by the difference of the excitatory and inhibitory inputs). The possible role of balanced input in neuronal gain modulation was highlighted by a recent in vitro study in which a variable current (with zero mean) was injected into a rat cortical pyramidal neuron and the gain associated with the injection of an additional constant current was measured [4]. The results indicated that the variability of the injected current affected the neuronal gain multiplicatively. In this paper we examine gain modulation as a result of balanced synaptic inputs in a conductance-based model. The analysis will be carried out in the gaussian approximation [2,3,1], which is most accurate for neurons with a large number of small-amplitude synaptic inputs. The effect of synaptic input over a range of input conditions (input rates, numbers of synapses and EPSP/IPSP amplitudes) is analyzed. The gain is measured as the rate of increase of the output spiking-rate of the neuron in response to increased input. Two different forms of input are examined: excitatory synaptic input (i.e. in excess of the balanced component of the synaptic input) and injected current.

2 Methods

A conductance-based leaky integrate-and-fire neuron (with reversal potentials) is used in which the membrane potential V(t) receives from its presynaptic inputs both excitatory and inhibitory contributions and decays in time with a characteristic time constant (the membrane time constant τ):

$$dV(t) = -\frac{(V(t) - v_0)}{\tau} dt + a_E(V_E - V(t)) dP_E(t) + a_I(V_I - V(t)) dP_I, (1)$$

where v_0 is the reset potential, V_E and V_I are the (constant) reversal potentials ($V_I \leq v_0 \leq V(t) \leq V_{\rm th} < V_E$). $dP_E(t)$ and $dP_I(t)$ are independent temporally homogeneous Poisson processes with constant intensities $N_E \lambda_E$ and $N_I \lambda_I$ respectively, i.e., each of the N_E excitatory input fibers (and N_I inhibitory input fibers) has a rate λ_E (resp. λ_I). In the balanced neuron considered here we choose $N = N_E = N_I$. The postsynaptic potential amplitudes a_E and a_I are chosen to be nonnegative (and are identical for all excitatory and inhibitory inputs respectively). The values of a_E and a_I are chosen so that the average excitatory and inhibitory inputs are balanced at the potential V_B ($V_I < V_B < V_E$), denoted the balance-potential,

$$a_E \frac{(V_E - V_B)}{\theta} = a_I \frac{(V_B - V_I)}{\theta} \equiv b = b_0 / \sqrt{N}$$
 , $\theta = V_{\text{th}} - v_0$. (2)

When the membrane potential reaches a threshold, an output spike is generated and the membrane potential is reset to its resting value v_0 . In the absence of spike generation, the membrane potential reaches an equilibrium value, V_Q , about which it fluctuates with variance σ_Q . The membrane potential approaches V_Q with a time constant given by τ_Q (see [3] for the values of V_Q , σ_Q , τ_Q). The analysis is carried out in the Gaussian approximation [2], in which the probability density of the membrane potential p(v, t | v', 0) is parameterized as

$$p(v, t \mid v', 0) = \frac{1}{\sqrt{2\pi\Gamma(t; v')}} \exp\left\{-\frac{(v - \Upsilon(t; v'))^2}{2\Gamma(t; v')}\right\},\tag{3}$$

where $\Upsilon(t;v)$ and $\Gamma(t;v)$ are the (time-dependent) mean and variance of the membrane potential. The Gaussian approximation is accurate in the limit of a large number of input synapses, N, which allows the probability density of the membrane potential to be evaluated using a self-consistent analysis [1]. The output spike distribution $f_{\theta}(t)$ obeys the renewal equation

$$p(V_{\rm th}, t \mid v_0, 0) = \int_0^t dt' \ f_\theta(t') \ p(V_{\rm th}, t \mid V_{\rm th}, t'). \tag{4}$$

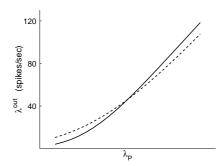
The effect of reversal potentials is to impose a lower bound to the membrane potential, and consequently it is necessary to consider their effect when inhibition plays an important role. In the model where reversal potentials are neglected, the amplitudes of the excitatory and inhibitory post-synaptic potentials (EPSPs and IPSPs resp.) are constant (i.e. independent of the existing membrane potential). In this case the gaussian approximation gives results that are identical with the diffusion method, in which the membrane potential is modelled as a stochastic process that is characterized by its first two moments. However, the gaussian approximation differs from the diffusion method when reversal potentials are included, since it accounts for non-linear summation and allows quite different time-courses for the mean, $\Upsilon(t; v)$, and variance, $\Gamma(t; v)$, of the membrane potential (these are given explicitly in [1]).

3 Results

An essential part of the analysis is defining the "normal operating regime" of a neuron, to ensure that the chosen parameter values correspond to biologically relevant neural behavior. This is achieved in our model by the condition that the output spiking-rate in the absence of a stimulus corresponds to the spontaneously active spiking-rate. This requires that there be a slight excess of excitation in the spontaneously active neuron (for the values used here, this is satisfied by $\lambda_E = \lambda_I + \lambda_P = \lambda_{\rm spon}$ with $\lambda_P = 1$ spike/sec).

The parameter values chosen are: $V_I = -75$ mV, $v_0 = -65$ mV, $V_{\rm th} = -55$ mV, $V_E = 0$ mV, so that the ratios of the potential differences are $(v_0 - V_I)$: $(V_{\rm th} - v_0)$: $(V_E - V_{\rm th}) = 1$: 1: 5.5. The membrane time constant is $\tau = 10$ ms, the amplitude of the post-synaptic potentials is given by $b_0 = 1$, the spontaneous spiking-rate is $\lambda_{\rm spon} = 5$ spikes/sec, the balance potential is $V_B = V_Q$, and the number of excitatory and inhibitory synapses is $N = N_E = N_I = 4000$. The results are not very sensitive to the choice of $V_B = V_Q$ and $b_0 = 1$ so long as λ_P is chosen appropriately to fulfill the above condition on the spontaneous spiking-rate.

Figure 1 shows the output spiking-rate (left plot) for the parameter values given above: The solid line is for the "1X" case in which the balanced input is just the spontaneous activity, and the dashed line is for the "2X" case in which the balanced input has twice the spiking-rate of the spontaneous activity. The plot on the right shows the ratio of the gains for the "2X" and "1X" conditions. Figure 2 shows the equivalent results for injected current.



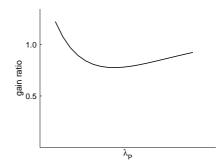
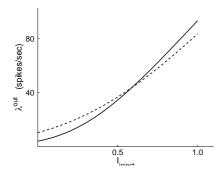


Fig. 1. Plot of results for neuronal gain in the case with excitatory synaptic input (a) Output spiking-rate, λ^{out} , for the "1X" (solid line) and "2X" (dashed line) cases, and (b) the ratio of the gain of the "2X" and "1X" conditions.



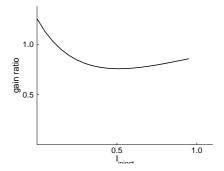


Fig. 2. Plot of results for neuronal gain in the case with injected current input, as for Fig. 1.

4 Discussion and Conclusions

There has been considerable interest in the role of balanced synaptic inputs recently, since neurons receiving balanced input produce a variability of spike times (usually measured by the coefficient of variation of the interspike interval distribution) that agrees with values observed in cortical neurons [11]. Neurons that receive predominately excitatory input, on the other hand, would be expected to have a membrane potential that is relatively smooth, resulting in regular firing and a small value of the coefficient of variation [12].

The results of this study indicate that, in the normal operating regime for the neuron, the effect upon the neuronal gain of balanced synaptic input is not simply multiplicative. However, the balanced inputs do give rise to a nonlinear gain modulation effect, when the gain is measured both with respect to synaptic input and to injected current. Although the results presented here represent only one set of parameters, they are representative of the results over most of the normal operating regime of a typical cortical neuron.

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References

- [1] A.N. Burkitt, Balanced neurons: Analysis of leaky integrate-and-fire neurons with reversal potentials, *Biol. Cybern.* 85 (2001) 247–255.
- [2] A.N. Burkitt, G.M. Clark, Calculation of interspike intervals for integrate-and-fire neurons with Poisson distribution of synaptic inputs, *Neural Comput.* 12 (2000) 1789–1820.
- [3] A.N. Burkitt, G.M. Clark, Synchronization of the Neural Response to Noisy Periodic Synaptic Input, *Neural Comput.* 13 (2001) 2639–2672.
- [4] F.S. Chance, L.F. Abbott, A.D. Reyes, Synaptic input variance controls the gain not the variability of neuronal responses, *Brandeis Uni preprint* (2002).
- [5] R. Hahnloser, R.J. Douglas, M. Mahowald, K. Hepp, Feedback interactions between neuronal pointers and maps for attentional processing, *Nature Neurosci.* 2 (1999) 746-752.
- [6] N. Hohn, A.N. Burkitt, Shot noise in the leaky integrate-and-fire neuron, Phys. Rev. E 63 (2001) 031902.
- [7] B.W. Mel, Synaptic integration in an excitable dendritic tree, *J. Neurophysiol.* 70 (1993) 1086–1101.
- [8] E. Salinas, L.F. Abbott, A model of multiplicative neural response is parietal cortex, *Proc. Natl. Acad. Sci. USA* 93 (1996) 6461–6474.
- [9] E. Salinas, P. Thier, Gain modulation: A major computational principle of the central nervous system, *Neuron* 27 (2000) 15–21.
- [10] E. Salinas, T.J. Sejnowski, Impact of correlated synaptic input on output firing rate and variability in simple neuronal models, J. Neurosci. 20 (2000) 6193– 6209.
- [11] M.N. Shadlen, W.T. Newsome, Noise, neural codes and cortical organization, Curr. Opin. Neurobiol. 4 (1994) 569–579.
- [12] W.R. Softky, C. Koch, The highly irregular firing of cortical cells is inconsistent with temporal integration of random EPSPs, J. Neurosci. 13 (1993) 334–350.