SLOW ADAPTATION IN FAST SPIKING NEURONS OF VISUAL CORTEX

Running title: Adaptation in fast spiking neurons

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Abstract

Fast spiking (FS) neurons are a class of inhibitory interneurons, classically characterized as having short duration action potentials (<0.5 ms at half height) and displaying little to no spike frequency adaptation during short (< 500 ms) depolarizing current pulses. As a consequence, the resulting injected current intensity versus firing frequency relationship is typically steep and they can achieve firing frequencies of up to 1 KHz. Here we have investigated the properties of FS neurons discharges on a longer time scale. Twenty second discharges were induced in electrophysiologically identified FS neurons by means of current injection, either with sinusoidal current or with square pulses. We found that virtually all FS neurons recorded in cortical slices do show spike frequency adaptation, but with a slow time course ($\tau = 2\text{–}19 \text{ s}$). This slow time course has precluded the observation of this property in previous studies that used shorter pulses. Contrary to the classical view of FS neurons functional properties, long duration discharges were followed by a slow after-hyperpolarization (AHP) lasting up to 23 seconds. During this post-adaptation period the excitability of the neurons was decreased on average for $16.7 \pm 6.8 \text{ sec}$, therefore rendering the cell less responsive to subsequent afferent inputs. Slow adaptation is also reported here for FS neurons recorded in vivo. This longer time scale of adaptation in FS neurons may be critical for balancing excitation and inhibition as well as for the understanding of cortical network computations.
Introduction

Fast spiking (FS) neurons are an electrophysiologically defined subtype of cortical neuron that corresponds to several morphological subtypes of non-pyramidal cells in the hippocampus and cortex (Kawaguchi and Kubota 1993, 1997; McCormick et al. 1985; Tasker et al. 1996; Thomson et al. 1996). Classically, cortical neurons have been classified as FS based upon their short duration action potentials (< 0.5 ms spike width at half amplitude), rapid rate of spike repolarization, lack of spike frequency adaptation, and a steep frequency versus injected current relationship (McCormick et al. 1985; Nowak et al. 2003). The lack of repetitive burst firing in FS neurons differentiates this cell class from another type neuron, chattering cells, that display equally fast action potentials but generate them in bursts (Brumberg et al. 2000; Gray and McCormick 1996, Nowak et al. 2003). FS neurons can discharge at high rates, sometimes reaching 1 KHz, and they have a steep frequency versus injected current relationship. These properties have been characterized in the cortex in vitro (Connors and Gutnick 1990; Kawaguchi 1993; Llinas et al. 1991; McCormick et al. 1985; Galarreta and Hestrin 2001; Thomson et al. 1996) and in vivo (Mountcastle et al. 1969; Simons 1978, Azouz et al. 1997; Contreras and Palmer 2003; Nowak et al. 2003; Nunez et al. 1993). A significant point of agreement between the in vivo and in vitro studies is that FS neurons can discharge action potentials at high rates with little or no spike frequency adaptation or attenuation in spike height. The basis
for some of these functional properties have been traced to the molecular level, and they are attributable to the expression of specific Na⁺ and K⁺ channels that are characterized by fast activation and recovery from inactivation kinetics (Martina and Jonas 1997; McBain and Rudy 2001). The rising phase of Na⁺ channel activation in FS and regular spiking neurons are governed by similar kinetics, but the rate in which the Na⁺ channels recover from inactivation is faster in FS neurons, thus allowing them to generate subsequent action potentials sooner (Martina and Jonas 1997). In a synergistic arrangement FS neurons express K⁺ channels of the Kv3 family that mediate a very fast repolarization, therefore shortening the action potential duration and facilitating recovery from Na⁺ channel inactivation (Erisir et al. 1999; McBain and Fisahn 2001; Rudy et al. 1999; Rudy and McBain 2001).

In the present report we show that the presence or absence of spike frequency adaptation in FS neurons is dependent upon the length of time they are stimulated. For short duration stimuli (e.g. <500 ms depolarizing current pulses), FS neurons show little or no spike frequency adaptation. However, we find that, although they do not show adaptation with short pulses, FS neurons do display adaptation during long (~20 sec) periods of activation. Additionally, this prolonged activation evokes a slow afterhyperpolarization (AHP) that decreases excitability for tens of seconds. Slow adaptation and AHP’s of this time scale have been previously described for cortical pyramidal neurons, the
underlying current being a Na\(^+\) dependent K\(^+\) current (Constanti and Sim 1987; Schwindt et al. 1989; Kubota and Saito 1991; Safronov and Vogel 1996; Sanchez-Vives et al. 2000b). This current has been implicated in vivo as one of the cellular mechanisms that underlie the phenomenon of contrast adaptation (Sanchez-Vives et al. 2000a, b). The finding that slow frequency adaptation also occurs in FS neurons may have important implications for understanding the computations performed by the cortical network.
Methods

In vitro experiments were performed on cortical slices from 2- to 6-month-old ferrets of either sex that were deeply anesthetized with sodium pentobarbital (40 mg/kg) and decapitated. Their brains were quickly removed and placed in ice cold cutting solution (see below). Ferrets were cared for and used in accordance with all appropriate regulatory guidelines. Four hundred-micrometer-thick coronal slices of the primary visual cortex were cut on a vibratome. A modification of the technique developed by Aghajanian and Rasmussen (1989) was used to increase tissue viability. After preparation, slices were placed in an interface-style recording chamber (Fine Sciences Tools, Foster City, CA) and bathed in ACSF containing (in mM): NaCl, 124; KCl, 2.5; MgSO₄, 2; NaHPO₄, 1.25; CaCl₂, 2; NaHCO₃, 26; and dextrose, 10, and was aerated with 95% O₂, 5% CO₂ to a final pH of 7.4. Bath temperature was maintained at 34-35°C. Intracellular recordings were initiated after 2 hr of recovery. Sharp intracellular recording electrodes were formed on a Sutter Instruments (Novato, CA) P-80 micropipette puller from medium-walled glass and beveled on a Sutter Instruments beveller to final resistances of 50-100 MΩ. Micropipettes were filled with 2 M KAc. Recordings were digitized, acquired and analyzed using a data acquisition interface and software from Cambridge Electronic Design (Cambridge, UK).

Intracellular recordings in vivo were obtained in the primary visual cortex of cats following the methodology that we have previously described (Sanchez-Vives et al., 2000a). In short, adult cats were anesthetized with ketamine (12-15 mg/kg,
i.m.) and xylazine (1 mg/kg, i.m.). The cat was then mounted in a stereotaxic frame and ventilated with either a mixture of nitrous oxide and oxygen 2:1 with halothane (1.5%), or with oxygen and isoflurane (2.5%). A craniotomy (3-4 mm wide) was made overlying the representation of the area centralis of area 17. During recording, anesthesia was maintained with 0.4-1% halothane or with 0.5-2% isoflurane. The heart rate, expiratory CO₂ concentration, rectal temperature, and blood O₂ concentration were monitored throughout the experiment and maintained at 150-180 bpm, 3-4%, 37-38°C, and >95%, respectively. The EEG and the absence of reaction to noxious stimuli were regularly checked to insure an adequate depth of anesthesia. To minimize pulsation arising from the heartbeat and respiration, a cisternal drainage and a bilateral pneumothorax were performed, and the animal was suspended by the rib cage to the stereotaxic frame. After the recording session, the animal was given a lethal injection of sodium pentobarbital. This protocol was approved by the Yale University Institutional Animal Care and Use Committees and conforms to the guidelines recommended in Preparation and Maintenance of Higher Mammals During Neuroscience Experiments, National Institutes of Health publication No. 91-3207.

Intracellular recordings were obtained with identical micropipettes to the ones used to record from the cortical slices. Data recording and acquisition were made using the same methods as for the in vitro data.
In order to identify neurons as FS we used 300-500 ms depolarizing current pulses. Longer periods of firing were induced by sinusoidal current injection at a frequency of 2 Hz for 20 sec or by depolarizing square pulses of 20 sec. The instantaneous frequency was computed as the inverse of the interspike interval. Responses to several repetitions of the current injection were averaged and they were quantified as spikes per cycle for sinusoidal current injections or in spikes per second for the square pulse current injections (500 ms bins). The slow spike frequency adaptation was measured as the firing rate decay between the first and the last 500 ms of the 20 sec of stimulation. The decay time constant was determined by fitting an exponential to the 40 values corresponding to the average number of spikes per cycle (sinusoids) or to the mean frequency for 500 ms bins (square pulse) (Fig. 1E, F). In order to minimize the effects of the synaptic noise *in vivo*, the adaptation strength *in vivo* was determined from the amplitude at time 0 and 20 sec given by the exponential function. Post-adaptation excitability change was assayed by injecting depolarizing current pulses (120 – 300 ms). Data are reported as mean ± SD.
Results

Fifteen fast spiking (FS) neurons recorded from slices of ferret visual cortex were included in this study. In order to be included, the neurons fulfilled the following criteria (McCormick et al. 1985; Nowak et al. 2003): short duration action potentials (0.35 ± 0.06 ms), absence of burstiness, little or no spike frequency adaptation during < 500 ms depolarizing pulses, and a steep relationship of the intensity of current injection versus the frequency of the discharge. Figure 1 (A, B, C and D) illustrates all these properties. The lack of spike frequency adaptation during 300 ms depolarizing pulses regardless of current intensity is manifest in Fig. 1D. However, when in the same neuron longer spike trains were induced by means of a prolonged depolarizing pulse (20 sec, Fig. 1E, F) or sinusoidal current injection (Fig. 1G, H), a slow spike frequency adaptation was revealed. In both cases the adaptation time constant was 2 sec and resulted in a 76% decrease in the firing frequency for the square pulse and of 48% for the sinusoidal current injection.

In our in vitro population data, all the cells (n=15) showed some degree of slow adaptation after 20 secs of high frequency firing. Slow frequency adaptation was studied in detail in 12 out of 15 cells with sinusoidal current injection at 2 Hz. The average decay in the number of spikes per cycle was of 44.0 ± 15.8 % and the average time constant of this decay was 8.8 ± 5.8 sec (n=12). Slow frequency adaptation was also explored by injecting 20 sec duration pulses in 5 out of 15 neurons (Fig. 1E, F). The average firing rate decay during 20 sec depolarizing
pulses was $53.5 \pm 30.7\%$ (last 0.5 sec with respect to the first 0.5 sec) and the time constant of the decay was $6.7 \pm 3.3$ secs.

A slow after-hyperpolarization (AHP) of the membrane potential was observed following the high frequency discharge induced using either protocol (Fig. 1E, G and Fig. 2A). This slow AHP had an amplitude that varied between 0.7 and 10 mV ($5.5 \pm 2.7$ mV) and was long lasting, between 3 and 23 secs ($12.4 \pm 5.6$ sec; n=11 cells in which the characteristics of the protocol allowed AHP quantification). A significant linear relationship ($p<0.01$) was found between the duration of the AHP and the time constant of the preceding spike frequency adaptation (not shown). The slow AHP decreased the excitability of the neurons during the post-discharge period (Fig. 2 C and E). This decrease in excitability was assessed in 7 FS cells by injecting depolarizing current pulses (120 - 300 ms) before and following the cessation of the 20 second square pulse or 20 seconds of sinusoidal current injection. The decrease in excitability lasted for up to 23 sec ($16.7 \pm 6.8$ sec). Figure 2D illustrates the linear relationship between the decrease of the AHP and the recovery of neuronal excitability. The distribution of the values of slow AHP duration and the duration of decreased excitability following a 20 sec discharge are represented in Fig. 2F.

**Intracellular recordings in vivo**

The effect of long lasting current injection has been studied in 9 FS neurons recorded intracellularly in cat area 17 in vivo. These cells were classified as FS according to the
criteria established by Nowak et al. (2003) (Fig. 3 A, B and C). One of these neurons was previously reported in (Sanchez-Vives et al. 2000a). Action potentials were induced by intracellular injection of 2 Hz sinusoidal current during 20 sec in 8 of these 9 neurons (Fig. 3F, G). As a result, 62.5% of the neurons (n=5/8) showed slow spike frequency adaptation. Adaptation strength determined as the firing rate for the last 0.5 sec of current injection relative to the firing rate for the first 0.5 sec of current injection (see Methods) was 29.7 ± 14.0 % (range: 10.2 – 46.4 %). In these 5 neurons the average time constant of the firing rate decay was 5.6 ± 4.7 sec. No adaptation was observed in the other 3 FS neurons.

Long duration (20 sec) square depolarizing pulses were injected in 3 FS neurons (including one that did not receive sinusoidal current injection) (Fig. 3E). This type of stimulus induced a larger amount of adaptation (40.5 ± 18.8 %). In two of the recordings a single exponential could be fitted to the decay of the firing, the time constants being 2.1 and 5.0 sec respectively.

In 2 out of the 6 neurons in vivo that showed slow spike frequency adaptation, a significant reduction in firing rate following the high frequency firing was observed (note PSTH in Fig. 3E). The membrane potential in the remainder cells (n=4/6) was subthreshold in between trials, and changes in the firing following the high frequency discharge could not be assessed. A slow AHP that followed the high frequency discharge and may have mediated the reduction in discharge during adaptation and after adaptation could be measured in 4 of the neurons. The AHP amplitude was between 3 and 8 mV (mean = 5.6 ± 2.7 mV) and the
duration between 5 and 10 sec (Fig. 3E, lower panel). The number of FS neurons showing slow AHPs may have been underestimated since the AHP may have been precluded by the large amount of synaptic noise typical of in vivo recordings. In none of the FS neurons that did not exhibit spike frequency adaptation was there an observable AHP (n=3).
Discussion

FS neurons have been characterized electrophysiologically classically as neurons with fast action potentials, the capability to reach high (up to 1 KHz) firing frequencies, steep slopes in discharge frequency versus current intensity relationship, and little or no spike frequency adaptation in response to depolarizing current injection. These characteristics have been established both with in vitro (Connors and Gutnick 1990; McCormick et al. 1985) and in vivo studies (Nowak et al. 2003; Nunez et al. 1993). Here we describe, to our knowledge for the first time, the existence of spike frequency adaptation in these classically defined FS neurons. In the present study we demonstrate that neurons with virtually no adaptation in response to short (< 0.5 sec) depolarizing current pulses, do actually adapt following long (tens of seconds) periods of activation. Both 20 sec long square depolarizing pulses and sinusoidal currents (2 Hz) have been used, and both evoked spike frequency adaptation. It is important to note that repetitive firing or prolonged activation as utilized in this study may be more relevant to the workings of the brain in vivo, where neurons discharge repetitively in response to external inputs or as part of internal rhythms.

One of the interesting aspects of FS neurons is that they constitute the only electrophysiological cell class unambiguously linked to inhibitory neurons. Neurons with identical or similar electrophysiological properties to those described here, and which have previously been termed FS neurons, have been associated with several subtypes of GABAergic non-pyramidal
neurons such as basket cells and chandelier cells (McCormick et al. 1985; Kawaguchi and Kubota 1993, 1997; Thomson and Deuchars 1997; Azouz et al. 1997), however in this study we did not attempt to discern between these various morphological types. It is likely that electrophysiological subtypes exist within the FS category (e.g. see Nowak et al. 2003). For example, one criteria used in some studies is the appearance of an abrupt transition between no action potentials to full trains of action potentials with small increases in the amplitude of injected current pulses (e.g. Kawaguchi and Kubota 1993, 1997). This criteria, however, is not useful for the classifications of in vivo recordings, since the presence of background synaptic activity prevents the measurement of threshold behaviors. In this study we have used our quantitative criteria to classify neurons as FS (McCormick et al. 1985; Nowak et al. 2003), with the understanding that this represents a heterogeneous population of cortical GABAergic neurons. In our hands (McCormick et al. 1985; Azouz et al. 1997; Nowak et al. 2003), all of the cells that we have classified as FS neurons and labelled with the same recording methods used here, have been morphologically consistent with what is expected for parvalbumin (+) cells (for a review see Kawaguchi and Kubota, 1997). Parvalbumin (+) cells have been reported to be up to 50% (Gonchar and Burkhalter, 1997 -rat-) and even 74% (van Brederode et al. 1990 -monkey-) of the total number of inhibitory cells. Therefore the slow adaptation that occurs in at least part of the
inhibitory cell population may have a large impact in its associated networks’ global discharges.

Fast spiking neurons recorded in vitro showed slow frequency adaptation with a time constant in the range of seconds for both sinewave current injections (8.8 ± 5.8 sec) and long (20 sec) depolarizing pulses (6.7 ± 3.3 sec). The reduction of the firing was similar in response to both protocols, with an average reduction of about 50% after 20 sec. In our previous study on putative excitatory neurons (non-FS neurons; Sanchez-Vives et al. 2000b), we found a very similar reduction of the firing rate (54.7%) at the end of 20 sec of sinusoidal current injection. Both adaptation strength and time constant (3.3 sec) showed a large heterogeneity between cells (see Figure 2C in Sanchez-Vives et al. 2000b) that could correspond to the inclusion of different cell types such as pyramidal and spiny stellate cells. When compared with the present data, we found that the average decay time constant is significantly faster for non-FS neurons than for FS neurons (p < 0.0006). This difference may be due to a relevant role in some of the non-FS neurons of adaptation currents with faster time course such as calcium-dependent potassium currents in non-FS cells (Madison and Nicoll, 1984; Pennefather et al. 1985; Schwindt et al. 1992; Vogalis et al. 2003).

Following long (20 sec) discharges, FS neurons recorded in vitro displayed a slow AHP that lasted up to 23 sec. This AHP was associated with a concurrent decrease in excitability of analogous duration. Therefore, the membrane event underlying the slow adaptation in FS neurons appears to be a slow AHP that lasts up to tens of seconds. Furthermore, we find a significant correlation between the time constant of the spike frequency
adaptation and the duration of the subsequent AHP, suggesting that the same mechanism underlies both phenomena. In our study on cortical non-FS neurons (Sanchez-Vives et al. 2000b), we found a slow AHP of 12-75 sec following long (20-60 sec) discharges, that also induced a long lasting decrease of excitability. In non-FS neurons we demonstrated this hyperpolarization was due, at least in large part, to the activation of a $K^+$ current activated by $Na^+$ (Sanchez-Vives et al. 2000b). This current has been described in many other neurons (Bader et al. 1985; Foehring et al. 1989; Dryer et al. 1989; Schwindt et al. 1989; Kubota and Saito 1991; Egan et al. 1992; Safronov and Vogel 1996; Kim and McCormick 1998; Bischoff et al. 1998) as well as in heart cells (Kameyama et al. 1984; Luk and Carmeliet 1990). The channels responsible for this current have recently been cloned (Yuan et al. 2003; Bhattacharjee et al. 2003). In the view of the similarity of adaptation and postadaptation time course, we suggest a $Na^+$-dependent $K^+$ current as a plausible candidate to underlie slow adaptation in FS neurons as well. Other possibilities cannot be, however, ruled out. A $Na^+/K^+$ ATPase electrogenic pump (Thomas 1972; Gustafson and Wigstrom 1983; Thompson and Prince 1986) would yield a similar time course and could contribute to the slow hyperpolarization. This possibility was tested in non-FS neurons (Sanchez-Vives et al. 2000b), where an increase in conductance following activation of the neurons and a lack of blockade by strophanthidin argued against an important contribution of a $Na^+/K^+$ ATPase electrogenic pump.

While all neurons studied in vitro (15 out of 15) displayed slow adaptation, even if small, only 67% of the FS neurons recorded in vivo did. In our in vivo studies we gave long
duration (20 sec) stimulus to 9 neurons, 6 out of which showed a significant adaptation. Also, the strength of adaptation was less in vivo (30% for the sinusoids, 40% for the pulse) than in vitro (44% for the sinusoids, 53% for the pulse). One explanation for this difference is that we more often used sinusoidal current injections in the in vivo studies, and these are less efficient than long square depolarizing pulses to induce adaptation. Another possibility is that ongoing activity in vivo may have an effect on the steady state of adaptation (Castro-Alamancos 2004): a fraction of Na⁺-dependent K⁺ current may be activated due to the presence of spontaneous activity in vivo, such that neurons are already partially adapted in the baseline condition. On the other hand, the lack of spontaneous activity in slices would result in neurons that have not been “pre-adapted”. This implies that FS neurons in vivo possess the same slow conductance as they do in vitro, but the basal state of their activation may be different in the two instances. Thus a neuron’s ability to be in different states of adaptation suggests a mechanism that may allow the neuron to integrate new inputs while still tracking its firing history. Finally, it is reasonable to suggest that adaptation in vitro may be more prominent than in vivo owing to a decrease in efficiency for meeting the metabolic demands of the neurons.

The slow adaptation of a whole population of inhibitory neurons may have important consequences for the computational properties of the cortical network. Neurons in vivo slowly adapt when they are stimulated for extended periods of time, i.e. in response to sustained high contrast stimulation (Maffei et al. 1973; Movshon and Lennie 1979; Albrecht et al. 1984; Ohzawa et al. 1985; Carandini and Ferster 1997; Sanchez-Vives et al. 2000a)
or as a result of long lasting current injection (Sanchez-Vives et al. 2000a). Intrinsic mechanisms such as slow adaptation currents have an effect somewhat similar to recurrent inhibition, in the sense that the spike discharge is regulated by the neuron’s own previous activity. The fact that both excitatory and inhibitory neurons share similar mechanisms for slow adaptation probably contributes to the excitatory/inhibitory balance within the cortical network (Shu et al. 2003).
Disclosures

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Figure legends

Figure 1. Slow frequency adaptation in a FS neuron recorded in vitro. A. Spike triggered average of the action potential (top trace). Below, the first derivative shows the fast rate of depolarization and repolarization characteristic of FS neurons. B. The interspike interval histogram reveals the lack of burstiness and short ISIs. C. Current intensity – firing frequency relationship. Each current intensity was applied twice and the responses have been averaged. The standard error of the mean (s.e.m.) is displayed in error bars, although in some cases the s.e.m. is too small to be seen. D. Instantaneous frequency (top) and spike discharge (middle) in response to the injection of four square current pulses of 300 ms duration and increasing intensity (bottom). Note that there is no spike frequency adaptation, but a slight acceleration of the discharge during the duration of the current pulse. E. Response of the same cell to the injection of a long duration (20 sec) square pulse of 0.75 nA. The rate of discharge (top) is represented in bins of 500 ms and the voltage trace (middle) shows the action potential discharge. Note that spike frequency adaptation develops slowly. F. Expanded traces of the voltage recording showed in E, corresponding to the beginning (a), middle (b) and end (c) of the long duration (20 s) pulse. G. Response of the same neuron to sinusoidal current (2 Hz, 1.1 nA peak-to-peak amplitude) injection for 20 sec. The slow adaptation of the number of spikes / cycle (top) had a time constant of 2.1 sec. AHP amplitude 8 mV; AHP duration 10–14 sec. H. Expanded traces of the voltage recording depicted in G, corresponding to the beginning (a) and end (b) of the 20 sec sinusoidal injection.
Figure 2. AHP and decreased excitability following prolonged firing in a representative FS neuron recorded in vitro. A. Slow AHP (4.6 mV amplitude) following a spike discharge induced by 20 sec of sinusoidal current injection (average of two trials). Hyperpolarizing pulses (-0.25 nA, 200 ms) were given to test changes in input resistance. Note the decrease in input resistance following the current injection (compare traces a, b). The PSTH plots the mean frequency of the firing calculated with a bin width (0.5 sec) equal to the sinewave period. The mean and standard deviation for two trials are plotted. B. The response of the neuron illustrated in part A to the first (top trace, Ba) and last (top trace Bb) cycles of the injected sinewave. Note the decrease in number of spikes/cycle. Below the traces are the instantaneous spike discharge frequency and the injected current trace (Bc). C. The excitability before and after high frequency firing (induced as in A) was explored by injecting depolarizing pulses of 0.2 nA and 200 ms duration at 0.5 Hz. The first two traces were obtained prior to the 20 sec adaptation protocol, the subsequent pulses correspond to 1.5, 3.5, 7.5, 11.5 and 15.5 sec after the end of the high frequency firing. Note that the excitability took longer than 10 seconds to recover. On average this neuron recovered to its pre-adapted stage in 22.7 ± 8.2 sec. D. Scatter plot representing the amplitude of the AHP preceding the pulses used to test the excitability of the cell (as in C) versus the number of evoked action potentials. AHP was induced by high frequency firing as in A. The linear regression is significant with p < 0.0001. Prior high frequency firing, the number of spikes evoked by the same 0.2 nA pulses was 9.5 ± 1.3.
E. Neuronal discharge for a sinusoidal current injection of low intensity (0.6 nA peak-to-peak amplitude), high intensity (0.9 nA) and low intensity. Mean frequency per cycle (2 Hz) is represented in the PSTH. Note the slow adaptation during the high frequency firing and long period of post-adaptation (firing rate reduction). The lower trace represents the current injection protocol, the frequency has been decreased for clarity. A, B, C, D and E correspond to the same neuron. F. Population histogram of the AHP durations and time of decreased excitability in bins of 5 sec.

**Figure 3.** Slow frequency adaptation in a FS neuron recorded in vivo. A. Spike triggered average of the action potential (top trace). Below, the first derivative shows the profile of fast depolarization and repolarization rate characteristic of FS neurons. B. Interspike interval distribution. Note lack of burstiness and short ISIs. C. Intensity – frequency relationship (slope = 292 Hz / nA). D. Voltage response (top) to the injection of four square current pulses of 300 ms duration and increasing intensity. The instantaneous frequency of the spike discharge is displayed (middle trace). E. Peristimulus histogram of the same cell to the injection of a long duration (20 sec) square pulse of 0.4 nA. The discharge rate is represented in bins of 500 ms and the error bars correspond to the s.e.m. for two trials. Note that there is adaptation during the 20 sec of firing and a period of decreased spontaneous discharge following the 20 sec pulse. The average membrane potential for the same two trials is displayed below the PSTH. The voltage trace has been low pass filtered to eliminate the action potentials and better show the slow AHP that follows the high frequency firing. F. Response of the same neuron
to sinusoidal current (2 Hz, 0.8 nA peak-to-peak amplitude) injection for 15 sec. The histogram displays the average number of spikes/cycle for 3 trials. Error bars represent s.e.m. The time course of the decay did not allow fitting by an exponential curve. G. Response of the neuron illustrated in F to two cycles of the injected sinewave. Top, membrane voltage, middle, instantaneous spike discharge frequency and bottom, the injected current.
Figure 1. Descalzo et al. 2004
Figure 2. Descalzo et al. 2004
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