Sluggish and brisk ganglion cells detect contrast with similar sensitivity

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Abstract

Roughly half of all ganglion cells in mammalian retina belong to the broad class, termed “sluggish”. Many of these cells have small receptive fields and project via lateral geniculate nuclei to visual cortex. However, their possible contributions to perception have been largely ignored because sluggish cells seem to respond weakly compared to the more easily studied “brisk” cells. By selecting small somas under infrared DIC optics and recording with a loose seal, we could routinely isolate sluggish cells. When a spot was matched spatially and temporally to the receptive field center, most sluggish cells could detect the same low contrasts as brisk cells. Detection thresholds for the two groups determined by an “ideal observer” were similar: threshold contrast for sluggish cells was 4.7 ± 0.5% (mean ± SE), and for brisk cells was 3.4 ± 0.3% (Mann-Whitney test: p>0.05). Signal-to-noise ratios for the two classes were also similar at low contrast. However, sluggish cells saturated at somewhat lower contrasts (contrast for half-maximum response was 14 ± 1% vs. 19 ± 2% for brisk cells) and were less sensitive to higher temporal frequencies (when the stimulus frequency was increased from 2 Hz to 4 Hz, the response rate fell by 1.6-fold). Thus the sluggish cells covered a narrower dynamic range and a narrower temporal bandwidth, consistent with their reported lower information rates. Because information per spike is greater at lower firing rates, sluggish cells may represent “cheaper” channels that convey less urgent visual information at a lower energy cost.

Key words: retinal ganglion cells, brisk, sluggish, contrast threshold
Introduction

The best studied ganglion cells in mammalian retina are classified as “brisk” because they respond rapidly at stimulus onset with relatively high firing rates (Cleland and Levick 1974; Stone and Fukuda 1974). Brisk cells are broadly tuned to spatio-temporal features of the stimulus, and they are highly sensitive to stimulus contrast; for example, the most sensitive ones can detect a spot over the receptive field center when contrast is as low as 0.5% (Linsenmeier et al. 1982; Dhingra et al, 2003; Dhingra and Smith, 2004). But brisk cells constitute only half of the population (Wässle and Boycott 1991; Rowe and Palmer, 1994; Troy and Shou 2002).

The rest can be classified as “sluggish” because they respond slowly at stimulus onset with low peak firing rates (Cleland and Levick 1974a, 1974b; Caldwell and Daw, 1978; Amthor et al., 1989). Sluggish cells can be quite selective for particular spatial and temporal features of the stimulus, such as local edge, slow motion, or motion in a particular direction (Levick, 1967; Stone and Fukuda 1974; Rowe and Cox 1993; Troy and Shou, 2002). These properties, which arise from strong inhibitory mechanisms (Caldwell et al., 1978), combine to make sluggish cells hard to “drive”, and this led us to wonder if they are fundamentally less sensitive than brisk cells.

This question seemed worth tackling for two reasons. First, although sluggish cells innervate brainstem structures, such as accessory optic and pretectal nuclei, and superior colliculus, they also innervate thalamic nuclei that project to striate and extrastriate cortex (Wilson and Stone, 1975; Troy and Shou, 2002; Dacey et al. 2003; Van Hooser et al., 2003). Thus, sluggish cells probably affect high level processes related to perception. Second, sluggish cells apparently receive far fewer excitatory synapses than brisk cells and relatively more inhibitory inputs (Kolb, 1979; Watanabe et al, 1985).
Therefore, to establish the relative sensitivities of brisk and sluggish cells should help clarify how sensitivity emerges from the balance of excitation and inhibition.

Sluggish cells are hard to record in vivo because their somas are small and their axons are fine. However, in vitro they are easier to record because a soma can be visualized with DIC optics and then attached stably to a loose-patch electrode (Koch et al, 2004). This approach, also used here, begins to restore the balance of knowledge between the brisk and sluggish components of the optic nerve.

**Methods**

**Tissue preparation and recording**

An adult guinea pig was anesthetized with ketamine (100mg/kg), xylazine (20mg/kg) and pentobarbital (50mg/kg). Then an eye was removed, and the animal was killed by overdose of pentobarbital (100mg/kg). The retina, attached to the pigment epithelium, choroid and sclera, was incised radially and flattened with ganglion cells up on a membrane filter in oxygenated (95% O₂, 5%CO₂) Ames medium (Sigma, St. Louis, MO) containing sodium bicarbonate (1.9g/l) and glucose (0.8g/l). After resting in darkness for about half an hour, the retina was mounted in a chamber on a microscope stage and superfused with oxygenated Ames medium (4-8 ml/min, 34-36°C).

A ganglion cell soma from the visual streak was selected under infrared DIC optics and cleared of Muller cell end-feet by squirting Ames medium from a pipette under mild pressure. A glass pipette (tip resistance 3-4 MΩ) was then attached loosely by mild suction. Spikes were amplified (Neurodata IR-283, Cygnus Technologies Inc, Delaware Water Gap, PA), high-pass filtered at 100Hz, and sampled at 5kHz by Axoscope software (Axon Instruments, Foster City, CA). Sometimes after recordings, we
replaced the patch pipette with a sharp electrode, injected DiI (2%, Molecular Probes, Eugene, OR), and subsequently imaged the cell with a confocal microscope.

**Visual stimulation**

Visual stimuli were generated with Matlab (MathWorks, Natick, MA) and displayed on a miniature monochrome computer monitor (640 x 480 pixels, 60 Hz) (Lucivid MR1-103, Microbrightfield, Colchester, VT) projected through a 4x objective and focused on the photoreceptors. Mean background intensity was 7900 photons/µm²/s, (equivalent to ~2x10⁴ R*/cone/s or ~0.6x10⁴ R*/rod/s at 535 nm), well into the photopic range. The relation between gun voltage and monitor intensity was linearized in software with a lookup table. Contrast was defined as (I_{max}-I_{mean})/I_{mean}, where I_{mean}=[d*I_{max}+(1-d)*I_{min}], where I_{max} and I_{min} are the maximum and minimum light intensities and d is the duty cycle. The stimulus was a spot of variable size, duration, temporal frequency, and contrast. Typically the spot was a 100 or 250 ms square-wave repeated at 0.5-4 Hz, and for the remainder of the cycle the mean background intensity was presented. At 2 Hz, the 100ms square wave constituted a duty cycle of 20%. For contrast threshold measurements, an ideal observer analyzed the responses within the whole cycle. When testing a cell’s temporal frequency tuning, we used a moderate contrast (10 -20%) sine-wave-modulated spot with 50% duty cycle, and the response was measured as the first harmonic component (F1), because this component was the largest one among other components that originated in nonlinearities and most nonlinear cells gave half-wave rectified responses to sine-wave stimuli.

To measure the time course of the mean effective stimulus, a random flickering checkerboard was presented to the retina. The checkerboard consisted of at least 5x5
squares with a size of 10x10 to 50x50 µm each, where the central one matched the receptive field center of the cell. Each of the squares changed its intensity independently according to a pseudorandom sequence at a frequency of 30 Hz. Then the sequence of patterns was cross-correlated with the cell’s spike response to obtain the mean effective stimulus for this cell (Fig. 1D, Fig. 6A). Temporal bandwidths of this cell were calculated from the Fourier-transform of the time course to produce the power spectra (Fig. 6B).

**Ideal Observer Analysis**

To determine a cell’s threshold for contrast detection, we constructed an ideal observer using a single-interval two-alternative forced-choice paradigm (Geisler et al, 1991; Dhingra et al, 2003). Responses were recorded to 100-200 repeated presentations of several contrasts between 0% and 30%. A given contrast was presented in blocks of 10-40 trials, which were interleaved with similar blocks at other contrasts in pseudorandom order. To reduce the effect of adaptation, blocks were interleaved with 5 s of mean background, and the first trial in each block was discarded.

Half of the responses (randomly chosen) at a given contrast were used to construct a histogram that was normalized to create a probability density function (PDF). Separate functions were constructed for each stimulus. Then the ideal observer was presented the remaining responses individually and decided for each response which of the two stimuli (a non-zero contrast stimulus vs. 0% or mean background) had most likely been presented. Because it was impractical to collect a sufficient amount of data to construct the ideal observer using a multidimensional histogram, we approximated the ideal observer with unidimensional histograms based on the temporal pattern of the spike trains. This method is more sensitive than one based on total spike count because it
emphasizes the time bins that contain the most information (Dhingra et al, 2003). The responses at each contrast were divided into temporal bins (40 ms), and the probability of 0 to n spikes was calculated for each bin. The spike pattern was obtained by multiplying the PDFs of all bins to give their joint probability. Then ideal observer made a choice by computing the likelihood ratio (Geisler et al., 1991) given by:

\[ L = \prod_{i=1}^{n} P_i(N_i / B) / \prod_{i=1}^{n} P_i(N_i / A) \]  

(1)

where \( N_i \) is the number of spikes in \( i^{th} \) bin, and \( n \) is the number of temporal bins, and \( A \) and \( B \) are the two stimuli (Dhingra et al., 2003). When \( L>1 \), the ideal observer chose stimulus \( B \); for \( L<1 \), it chose stimulus \( A \). For some analyses, we did not compute the joint probability but instead compared PDFs from single bins (Fig. 4D). The likelihood ratio computed in this way, since it was not an approximation, was optimal (Geisler et al., 1991; Dhingra & Smith, 2004). When the choice corresponded to the stimulus that had actually been presented, it was “correct”. A neurometric function was defined as the fraction of correct responses vs. contrast and fitted with a cumulative Weibull function (Weibull, 1951; Quick, 1974) (Fig. 4B) as follows:

\[ P=1-0.5\exp(-C/\alpha)^{\beta}, \]  

(2)

where \( P \) is the probability of correct choices, \( C \) is the contrast, \( \alpha \) is the threshold and \( \beta \) is the slope of the Weibull function. Threshold was defined as the contrast that gave 68% correct choices (Geisler et al., 1991; Dhingra et al, 2003).

**Signal and noise from contrast response function**

To understand the contrast detection thresholds in terms of signal-to-noise ratio, signal and noise were measured for each cell at different contrasts ranging from 0 to
100%. The signal was defined as the average evoked spike frequency summed over all trials (20 – 40 trials) within the single time bin (40ms) containing the peak firing rate. Noise was defined as the standard deviation of the response frequency in that bin across trials.

Although the noise computed by this method could be reduced by summing over more time bins, this would also decrease the signal. An attempt to generate an ideal filter to weight each time bin did not provide a better estimate of signal and noise because the response wave shape varied with contrast. Furthermore, because the number of trials we tested (20-40) in this series of experiments was limited, it was not possible to create a reliable estimate of the ideal temporal pattern to weight each time bin before summing them. Thus as a compromise, we choose a single bin.

The signal and noise responses vs. contrast were fitted with a modified Naka-Rushton equation (Naka and Rushton, 1966; Boynton et al., 1999; Dhingra and Smith, 2004) as:

\[ R(C) = R_{\text{max}} \times \left( \frac{C^p}{C^q + \sigma^q} - T \right) \]  

(4)

where \( R \) is the response, \( C \) is the stimulus contrast, and the other symbols determine the shape of contrast-response function. Increment thresholds were estimated from signal and noise measurements by assuming that at any contrast the minimum discriminable contrast increment was proportional to the noise in the response measured at that contrast (Dhingra and Smith, 2004).

Unless stated otherwise, we applied the non-parametric, Mann-Whitney test for statistical comparison.

**Results**
We studied 45 ganglion cells from the visual streak of 19 animals. Twenty-four cells responded briskly, firing at short latency and relatively high rates. Among them, brisk-transient cells (n=14) resembled the Y/alpha cells in cat and rabbit, with large somas, broad dendritic fields, and broad receptive field centers with modest surround antagonism (Fig. 1). They responded well to stimuli above 4 Hz. Brisk-sustained cells (n=10) resembled X/beta cells in cat and rabbit, with medium somas and medium dendritic fields. They resembled brisk-transient cells in latency, peak rates, and temporal sensitivity.

Twenty-one cells responded sluggishly, firing with long latency (Fig. 1D, Fig. 6A) and relatively low rates (Fig. 1B). Many of these cells had small somas, narrow dendritic fields, and narrow receptive field centers with strong surround antagonism, which resembled “W” cells in cat and rabbit (Amthor et al., 1989; Row and Cox, 1993; Troy and Shou, 2002). They responded best to stimuli around 1-2 Hz and hardly responded to stimuli above 4 Hz (Fig. 1C). Thus, considering three parameters: peak firing rate, optimal temporal frequency, and receptive field size, sluggish and brisk cells formed distinct clusters in multi-parametric space (Fig. 1E). The plot illustrates that no single parameter could separate brisk from sluggish cells, but the three parameters together clearly separated the tight sluggish cluster from the brisk cluster.

Sluggish cells comprise more than seven types (Troy and Shou, 2002), and we could identify several of them. For example, the local-edge cell had a narrow receptive field center, strong suppressive surround, and narrow dendritic field that stratified narrowly at mid-level of the inner plexiform layer (Fig 1). Similarly, the ON-OFF DS cell was directionally sensitive, with a medium dendritic field that stratified in both ON and OFF levels of the inner plexiform layer (Fig 1). Our sample of sluggish cells was too
small to evaluate the sensitivity of all the types, but within the group, we found no obvious differences in contrast sensitivity.

**Optimizing the stimulus and the ideal observer**

Before measuring a cell’s contrast threshold, we first determined the optimal spot size and temporal frequency. Brisk and sluggish cells preferred different temporal frequencies (Fig. 2A) which affected their spike responses and contrast sensitivities (Fig. 2B). The optimal temporal frequency for sluggish cells was less than 2 Hz, while for brisk cells it was 4 Hz or in some cases slightly higher. Using this estimation procedure, we selected the repetition rate used in each square wave spot experiment, typically 2 or 4 Hz for brisk cells and 1 or 2 Hz for sluggish cells (Fig. 2). Above 4 Hz, the brisk cells’ response to the stimulus flash did not recover to baseline completely between stimuli. Below 1 Hz, sluggish cells responded well, but the longer inter-trial intervals restricted the number of responses that could be collected. Stimulus duration was 100 ms for brisk cells (Dhingra et al., 2003) and usually 100 ms or 250 ms for sluggish cells; prolonging the stimulus to 400 ms gave no improvement.

With the stimulus optimized, we presented a cell with 100-200 trials of this optimal square-wave spot at 6 -10 contrasts (Fig. 3), in a session extending over 60 minutes. Stability of the cell was checked by measuring the contrast response function before and after these measurements.

We directed the ideal observer to evaluate the temporal pattern of spikes because this gives the lowest threshold (Dhingra et al, 2003). Since bin width would affect the performance of the ideal observer, we tested bin widths between 10-200 ms. For brisk cells, thresholds were generally lowest with a 40 ms bin (confirming Dhingra et al, 2003);
for sluggish cells, thresholds were insensitive to the bin widths tested, so for all cells we report their responses in 40 ms bins.

**Contrast thresholds**

Contrasts greater than ~10% evoked much stronger responses in brisk than in sluggish cells, but contrasts below ~10% evoked similar responses in both classes. Since detection thresholds were mainly determined by the weakest responses, both cell classes had similar thresholds. This feature, shown for a brisk and sluggish pair in Figure 3, was also characteristic for the two populations: different mean response at high contrast, but similar responses at low contrast, thus similar contrast thresholds (Fig. 4A,B).

The mean detection threshold for brisk cells was 3.4 ± 0.3% (mean ± SE, n=24), and for sluggish cells it was 4.7± 0.5% (n=19) (Fig. 4). The difference was slight and insignificant (p>0.05, Mann-Whitney -test). Furthermore, more than one-third of the cells in each class had thresholds between 3% and 4%, and ~85% of sluggish cells shared the same range of thresholds as brisk cells (1-6%, Fig. 4C). Thus, we conclude that contrast detection thresholds for the two broad classes are, if not identical, certainly close.

Within the brisk category, brisk-transient (BT) had similar contrast thresholds as brisk-sustained (BS) cells (BT: 3.2 ± 0.4%, n=14; BS: 3.6 ± 0.3 %, n=10; p>0.4, but see Troy, 1983; Kaplan and Shapley, 1982,1986). Within the sluggish category, two local-edge cells and one ON-sustained sluggish cell showed unusually low sensitivity (thresholds > 6%), but other cells of the same types were found to be more sensitive. For the five local-edge cells recorded, their thresholds ranged between 3.8- 8.4 %, the mean value (6.4 ± 0.9%) was not significantly different from five ON-OFF DS cells (4.0 ± 0.5
Other types of sluggish cell were too sparsely represented in our sample to allow meaningful comparison.

To understand how a neuron's ideal performance varied in time, we measured performance for individual time bins and compared sluggish with brisk cell (Fig. 4D). Performance showed two peaks, a robust one during the initial excitatory phase of the response and a weaker one during the recovery phase. At a near-threshold (4%) contrast, performance rose slowly then decreased, peaking for a brisk cell at 100 ms and for a sluggish cell at 160 ms (Fig. 4D), which corresponded well to the longer response latency of sluggish cell. Furthermore, performance for the sluggish cell in a single time bin was as high as (or even higher than) that for the brisk cell. However the overall thresholds determined by the joint probability from all the bins were less affected by these differences in the single bin performance. Thus, despite their different latencies and peak performance, the thresholds were nearly identical for these two cells (brisk, 3.9%; sluggish, 4.0%).

On the other hand, brisk and sluggish cells differed in the dynamic range and slope of their contrast response function. Brisk responses rose faster and saturated at higher contrast (Fig. 4A). For brisk cells the contrast for half-maximum response was 19 ±2% (mean ± SE, n=22), and for sluggish cells it was 13 ± 1% (n=19, Fig. 4E). This difference was significant (p<0.05). For individual cells, contrasts at threshold and at half–maximum response were uncorrelated (r=0.017, p>0.9, n=41).

**Contrast threshold vs. other response properties**

Contrast thresholds were uncorrelated with either maintained rate (r=0.01, p=0.95, Fig. 4F) or peak firing rate (r=-0.28, p>0.05) at high contrast when brisk cells fired more
strongly (at 50% contrast, 209±20 spikes s\(^{-1}\) vs. 102±8 spikes s\(^{-1}\) for sluggish, p<0.001).

Brisk cells had larger receptive field centers than sluggish cells (367 ± 28 \(\mu\)m vs. 180±21 \(\mu\)m, p<0.001), but their thresholds were uncorrelated with center size\((r=0.02, p>0.8)\).

Finally, although the optimal temporal frequency was higher for brisk cells than for sluggish (5.2±0.4 Hz vs.1.5±0.2 Hz, p<0.001), this was uncorrelated with threshold \((n=35, r = -0.32, p>0.05)\).

**Signal and noise**

Signal and noise were calculated separately at each contrast (see methods, Fig. 5A). Signal increased slowly at very low contrast and then steeply for both brisk and sluggish cells, but this nonlinear behavior was more pronounced for sluggish cells. Both signal and noise amplitudes were consistently lower for sluggish cells, so at low contrast the signal-to-noise ratios were the same for both classes (Fig. 5B), which contributed to their similar detection thresholds. At higher contrasts (>20%), signal-to-noise ratios for brisk cells improved relative to sluggish cells, and the difference was greatest for contrasts above 60% \((p<0.05)\), as shown in Fig. 5C.

We calculated contrast sensitivity (Fig. 5C) by inverting the increment thresholds estimated from the signal and noise measurements (see Methods; also Dhingra and Smith, 2004). Contrast sensitivity of the brisk and sluggish groups was quite close at low contrast (\(<10\%\)); sensitivity peaked near 4\% contrast for both groups; and then declined at higher contrasts faster for sluggish cells.

To estimate the total gray levels, we integrated contrast sensitivity over contrast. With a greater sensitivity above 20\% contrast, brisk cells had more gray levels than
sluggish (5.5 vs. 3.8). However, within the 0 - 10% contrast range, the gray levels were similar for brisk and sluggish (3.4 vs. 3.1).

**Temporal bandwidth**

Temporal bandwidths were estimated from the Fourier transform of the mean effective stimulus, testing with flickering checkerboards (Fig. 6A,B). Sluggish cells had a longer time to peak and had a lower temporal bandwidth than brisk cells.

The power spectra showed smaller differences in cut-off frequency between the two cell groups than shown in the temporal tuning curves (Fig. 2A), probably because of the different stimuli and contrast used. For temporal tuning curves, a sine-wave spot at 20% contrast was applied with a sequence of frequencies up to 10Hz; whereas the checkerboards comprised a greater range of frequencies, with contrasts up to 100%.

Previous work has shown that optimal frequency increased with contrast (Dhingra et al., 2003, Fig. 2B), so one would expect the optimal frequency to be closer for the two groups at higher contrast.

**Discussion**

**Contrast detection thresholds and sensitivities near threshold are similar for brisk and sluggish cells**

Our basic finding is that the threshold for detecting an optimal spot is similar for brisk and sluggish cells (3-4% contrast; Fig. 4). This sensitivity resembles previous studies of brisk-transient cells in the superfused, flat-mounted guinea pig retina (Dhingra et al, 2003) and *in vivo* measurements in other species (Derrington and Lennie, 1982,
Sensitivity found here for sluggish cells is slightly higher than reported in rabbit by Merwine et al. (1995). Since mammalian species seem to have rather similar sensitivities, the differences in this case may reflect differences in defining threshold and in sensitivity of the preparation.

Contrast gain is reportedly lower for sluggish cells than for brisk (Fig. 4A; Rowe and Cox, 1993; Merwine et al., 1995; Van Hooser et al., 2003). However, this measurement is based on a cell’s “responsivity” (spikes s\(^{-1}\) contrast\(^{-1}\)) and excludes the effect of noise. Although the brisk signal is greater than the sluggish signal, the noise is also larger (Fig. 5A). Therefore, despite the many clear differences between brisk and sluggish cells, their detection thresholds are similar (~4%), and at low contrasts the signal-to-noise ratios are similar, as are the numbers of gray levels that they encode (Fig. 5B, C).

Initially this seemed puzzling because the sluggish cells probably have far fewer excitatory synapses than brisk cells: the dendritic fields tend to be narrower (Fig. 1A), and the fraction of bipolar (excitatory) inputs seems much lower. For example, in cat, bipolar synapses comprise 40-80% of input to brisk-sustained cells and 15-50% for brisk-transient cells, but no more than 20% for sluggish cells (Cohen and Sterling, 1992; Freed and Sterling, 1988; Weber and Stanford, 1994; Watanabe et al., 1985; Kolb, 1979). Our working hypothesis was that a sluggish cell with far fewer excitatory synapses would collect fewer miniature postsynaptic excitatory currents (mEPSCs) with relatively greater Poisson fluctuations than a brisk cell and thus be both absolutely and relatively less sensitive (Kier et al., 1995, Croner et al., 1993).

On the other hand, sluggish cells tend to have higher membrane resistance. For example, for a brisk-transient (alpha) cell input resistance \((R_m)\) was \(\sim 31\text{M}\Omega\); whereas
for an ON-OFF DS cell (iota) $R_m \sim 396 \text{ M}\Omega$, and for a local-edge cell (zeta) $R_m \sim 1048 \text{ M}\Omega$. (O’Brien et al, 2002). Consequently, in smaller cells the same mEPSC evokes a larger depolarization. The higher membrane resistance also gives the smaller cell a longer time constant, for example, for the brisk-transient, $\tau \sim 4.5$ ms for the ON-OFF DS cell $\tau \sim 36.3$ ms, and for the local-edge cell $\tau \sim 81.6$ ms (O’Brien et al, 2002). The larger depolarization per quantal conductance plus the longer integration time might combine with the biophysical properties that comprise the spike generator (Dhingra and Smith, 2004) to render the small cells as sensitive as the large ones. There may be additional factors, such as differences in postsynaptic glutamate receptors (as found for cone input to different bipolar types -- see Devries, 2000) and differences in vesicle release rate (as found for output to different ganglion cell types --see Freed, 2000a,b). Finally, the sluggish cells, like the brisk cells, may use synchronous release of synaptic vesicles to minimize the effects of synaptic noise in the spike output (Demb et al 2004).

One could argue that a simple spot would underestimate contrast sensitivity for cell types selective for specific stimuli, such as direction of motion or local edges, because to such stimuli, the cell type may be more sensitive. However, our purpose in this study was to compare the basic contrast sensitivities of brisk and sluggish cells using a stimulus that synchronously activates all bipolar cells presynaptic to the ganglion cell. Thus, our measurements of contrast sensitivity to a spot optimized to fit the receptive field represent a lower bound.

Our measurements also reveal differences between brisk and sluggish cells. With a broader dynamic range and more gray levels, brisk cells can respond more effectively at higher contrasts. Although sluggish cells respond reliably at higher contrasts, their
responses saturate above 20%, sacrificing any ability to discriminate between different contrasts.

**Why should the retina need multiple pathways of nearly identical sensitivity?**

Despite resembling brisk cells in contrast sensitivity, contrast gain, and number of gray levels at low contrast, sluggish cells express markedly lower peak firing rates. This limits their information rate (bits s\(^{-1}\)), but increases their spike efficiency (bits spike\(^{-1}\); Koch et al., 2004). Sluggish cells also differ structurally from brisk cells, notably by having much finer axons (reviewed by Troy and Shou, 2002). Since the brisk cells cover a broader temporal bandwidth (Fig. 6) and broader dynamic range (Fig. 4), why should the retina expend additional resources on a multiplicity of ganglion cell pathways with low information rates?

In this respect the ganglion cells continue a trend that begins with the photoreceptors and bipolar cells. There, too, are many parallel lines where those with the lowest information rates use the finest axons with the fewest microtubules and the fewest synaptic outputs (Hsu et al., 1996; Sterling, 2004). These trends suggest a hypothesis, namely that each channel uses energy in proportion to its information rate (Laughlin et al., 1998). As in daily life, if some messages are important but not urgent, it might be energetically cheaper to send them at a lower rate. Consistent with this, sluggish somas stain weakly for cytochrome oxidase compared to brisk somas (Kageyama and Wong-Riley, 1984), although quantitative comparisons for the complete neurons are not available. If this is roughly correct, then the biophysical properties of sluggish cells, which limit the spike rate (O’Brien et al., 2002), and thus the information rate (Koch et al., 2004), offer clear advantages.
Acknowledgements

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Figure Legends

Figure 1. Identifying brisk and sluggish ganglion cells.

A. Each cell type expressed a distinctive morphology. Brisk-transient cell (BT) had a large soma and broad dendritic field. Sluggish cells, here a local-edge cell (LE) and an ON-OFF directionally selective cell (DS), had smaller somas and much narrower dendritic fields. All cells were located in the visual streak.

B. Sluggish cells fired at lower rates. Stimulus was a high contrast (50%) square-wave spot filling the receptive field center.

C. Sluggish cells responded at lower temporal frequencies. Stimulus was a sinusoidally modulated spot at 10% contrast. Ordinate: amplitude of the first harmonic responses (F1).

D. Time courses of mean effective stimulus for the three cells showed a longer time-to-peak for the two sluggish cells. The mean effective stimulus was obtained by cross-correlating a quasi-random stimulus sequence (“flickering checkerboard”) with each spike in a cell’s response (see methods).

E. Sluggish cells and brisk cells formed distinct clusters in a multiparametric space constructed from peak firing rate, size of receptive field and optimal frequency.

Figure 2. Optimize temporal frequencies for sluggish and brisk cells

A: Mean temporal tuning curves for sluggish and brisk cells. The optimal frequency for the sluggish cells was 1 or 2Hz, while that for brisk cells was around 4Hz or even higher.

B: Stimulus at 4Hz elicited stronger response thus higher contrast sensitivity for a brisk cell than at 2 Hz, while a sluggish cell was more sensitive to a stimulus at 1 Hz than either 2 Hz or 0.5 Hz.
Figure 3. Sluggish cell showed lower gain than brisk cell but similar threshold for contrast detection.

A. Spike responses of a brisk and a sluggish cell to 100 trials at each of several contrasts near threshold. Stimulus was presented to the brisk cell at twice the frequency and for a shorter duration than for the sluggish cell.

B. Below ~10% contrast, evoked responses were similar for the sluggish and brisk cells, rising sharply with high gain. At higher contrasts the brisk cell’s response continued to rise and saturated around 45%, whereas the sluggish cell’s response rose more slowly and saturated earlier, at ~ 30%.

C. Contrast detection threshold was determined by an ideal observer using all the information available in the temporal pattern of spiking (see Methods). Thresholds for these two cells were nearly identical.

Figure 4. Contrast sensitivity functions and detection thresholds for sluggish and brisk populations.

A. Sluggish cell responses to contrast rose more slowly and saturated earlier. Vertical bars, standard error (SE).

B. Sluggish and brisk detection thresholds were similar. The fraction of correct responses vs. contrast was fitted with cumulative Weibull function (equation 2) for the averaged values across cells.

C. Most sluggish cells (>85 %) showed detection thresholds between 1.8 - 5.8%, and this range was bracketed by the brisk cells (0.9% - 5.9%). The population means did not differ significantly.
D. Sluggish cell detection sensitivity during the stimulus cycle (4% square-wave spot, near threshold for this pair of cells) peaked about 60 ms later than for brisk cells. Each bar represents the cell's ideal performance for one time bin (20ms), see Methods.

E. Distribution of contrasts that gave half-maximal response overlapped for sluggish and brisk cells, but the population means differed (p <0.05) because brisk cells tended to saturate at higher contrasts.

F. Contrast detection thresholds were uncorrelated with maintained rate.

**Figure 5. Similar signal-to-noise ratios and gray levels for sluggish and brisk cells at low contrasts (<10%).**

A. Signal and noise were both slightly lower for sluggish cells. Nonlinearities were observed at very low contrast (inset) and high contrast. Signal curves were fitted with equation (4).

B. Signal-to-noise ratios. At contrasts lower than 10%, no difference was observed between two cell groups, though at higher contrast, brisk cells achieved a greater signal-to-noise ratio. Curves were fitted with equation (4).

C. Contrast sensitivity, the inverse of the increment threshold estimated from the measured signal and noise curves, showed similar integrated area (gray levels) for brisk and sluggish cells up to 10% contrast (3.4 vs. 3.1). At higher contrasts brisk cells continued to show substantial sensitivity.

**Figure 6. Sluggish cells transmit at lower bandwidths**
A. Time course of the mean effective stimulus showed that sluggish cells took longer times to peak and to recovery. The functions for ON cells (4 sluggish cells and 7 brisk cells) were inverted here to combine with OFF cells.

B. Power spectra (Fourier-transform of the mean effective stimulus) showed lower bandwidth for sluggish cells.
Fig. 1

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Fig. 2

Panel A: Graph showing F1 (spikes/s) vs. temporal frequency (Hz). The graph compares 'Brisk (n=18)' and 'Slug (n=21)' conditions.

Panel B: Graph showing firing rate (spikes/s) vs. contrast (%). The graph includes lines for 'Brisk 4Hz', 'Brisk 2Hz', 'Slug 1Hz', 'Slug 2Hz', and 'Slug 0.5Hz'.

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Fig. 4

A. Peak rate (spikes s⁻¹) vs. contrast (%)

B. Correct response vs. contrast (%)

C. Number of cells vs. contrast threshold (%)

D. Correct response vs. stimulus duration (ms)

E. Number of cells vs. half-max contrast (%)

F. Contrast threshold vs. maintained rate (spikes s⁻¹)

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Fig. 6

A

B

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