A theory of direction selectivity for Macaque primary visual cortex

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Abstract

This paper offers a new theory for the origin of direction selectivity in the Macaque primary visual cortex, V1. Direction selectivity (DS) is essential for the perception of motion and control of pursuit eye movements. In the Macaque visual pathway, DS neurons first appear in V1, in the Simple cell population of the Magnocellular input layer 4Cα. The LGN cells that project to these cortical neurons, however, are not direction-selective. We hypothesize that DS is initiated in feedforward LGN input, in the summed responses of LGN cells afferent to a cortical cell, and it is achieved through the interplay of (a) different visual response dynamics of ON and OFF LGN cells, and (b) the wiring of ON and OFF LGN neurons to cortex. We identify specific temporal differences in the ON/OFF pathways that together with (b) produce distinct response time-courses in separated subregions; analysis and simulations confirm the efficacy of the mechanisms proposed. To constrain the theory, we present data on Simple cells in layer 4Cα in response to drifting gratings. About half of the cells were found to have high DS, and the DS was broad-band in spatial and temporal frequency (SF and TF). The proposed theory includes a complete analysis of how stimulus features such as SF and TF interact with ON/OFF dynamics and LGN-to-cortex wiring to determine the preferred direction and magnitude of DS.

Significance Statement

Motion perception is important for primates, and direction selectivity (DS), the ability to perceive the direction a target is moving, is an essential part of motion perception. Yet no satisfactory mechanistic explanation has been proposed for the origin of DS in primate visual cortex up until now. In this paper, we hypothesize that DS is initiated in feedforward LGN input as a result of the dynamic differences between the ON/OFF pathways. The mechanisms we propose are biology-based, and our theory explains experimental data for all spatial and temporal frequencies in visual stimuli. Exploiting temporal biases in parallel pathways is relevant beyond visual neuroscience; similar ideas likely apply to other types of neural signal processing.

Introduction

This paper proposes a solution to a longstanding question in visual neuroscience, namely the origin of direction selectivity in the visual cortex of Macaque monkeys. Motion perception is a vital visual capability well developed in primates. As perceiving motion requires perceiving the direction in which a target moves, direction selectivity (DS), the ability of visual neurons to sense the direction of movement, is essential for motion perception (1) and for the control of pursuit eye movements (2). For these reasons, understanding DS is an important first step towards understanding how the cortex processes motion signals.

Direction selectivity in cortical neurons was first documented in the cat (3). Since then, it has been found in neurons all along the visual dorsal stream (an area associated with motion processing) in primates like Macaque monkeys (4-7), whose vision is like that of humans. DS neurons are in fact present across species; they are widespread among visual mammals, an experimental fact that testifies to their biological significance.

In the visual pathway of Macaques, DS appears first in the primary visual cortex (V1), in the Simple cell population of the input layer 4Cα (8, 9). These neurons provide feedforward DS signals to subsequent cortical layers and brain regions in the dorsal pathway. Thus to discover the origin of DS, one is led to examining how neurons in layer 4Cα acquire their DS --- and that is where it gets interesting: The neurons that provide visual signals to layer 4Cα, the Magnocellular cells in the Lateral Geniculate Nucleus (LGN), are not direction-selective (10-13). Yet many of the cells in the input layer of V1 to which they project are DS. A fundamental scientific question, therefore, is how 4Cα neurons acquire their DS. That is the question we would like to answer in this paper.

Though many papers have been written on DS since its discovery over half a century ago, and there is continued interest in the subject (14-17), no satisfactory mechanistic explanation for the origin of DS has been proposed before now: Early conceptual models of how DS may arise, such as the Reichardt multiplier (18) or the motion-energy model (19), were not concerned with biological mechanisms. Some DS models have invoked intra-cortical inhibitory mechanisms for DS (20, 21, 17), in particular cortical inhibition tuned to the non-

preferred direction, but intracellular-recording (22) showed such models to be implausible. Other results are for different animals. For example, in their recent papers, Lein and Scanziani (15) and Billeh et al (16) adopted the Sustained-Transient theory from earlier studies in cat (23) to explain DS in mouse, but this theory is implausible for Macaque because for Macaque there are not clearly separate classes of Sustained and Transient Magnocellular LGN neurons.

It is widely accepted that the essence of the DS computation has to do with spatio-temporal inseparability (19). What was lacking were biological mechanisms that could produce such biases, and a clear understanding of why such biases would necessarily result in a meaningful notion of direction selectivity. These are the issues we would like to address. In this paper we hypothesize that a plausible mechanism is the interplay between (a) the different dynamics of ON and OFF LGN cells and (b) the specific wiring that connects them to V1. The latter is well known: OFF and ON LGN cells are wired specifically to segregated V1 receptive field subregions, an idea that goes all the way back to Hubel and Wiesel (24, 25). As for (a), we will identify in **Results** specific differences in the ON/OFF pathways that together with (b) lead to distinct response time-courses in separated subregions. The mechanisms we propose are biologically grounded, and as we will show, they are sufficient for initiating DS in the feedforward LGN input to cortical cells.

To constrain our theory, we present also experimental results on the responses of a group of 4Cα Simple cells to drifting gratings. Our data show that most of the cells recorded were unambiguously DS, preferring consistently the same direction over their entire visible ranges of spatial frequency (SF) and temporal frequency (TF); about half of the cells recorded had high DS. Our data reveal also an important characteristic of DS neurons, namely the approximate invariance of DS with SF and TF. Explaining the broad-band character of DS (in TF and SF) is a challenge for all previous theories. Our theoretical predictions are in excellent agreement with data, and it includes a complete analysis of how stimulus features like SF and TF interact with ON/OFF dynamics and LGN-to-cortex wiring to explain the broad-band character of DS.

With regard to broader implications, although the theory as described in this paper is specifically about DS, an important message is that when combining information from multiple

channels, slight biases in their temporal filters can greatly enhance the capability of a system. Thus it may be possible to exploit further the temporal axis in signal processing biological and nonbiological, especially in the neural processing of sensory inputs and possibly in computer vision.

Results

I. Experimental data

First, we must show that most Simple cells in Macaque 4Cα are DS cells. The data that show it are the responses of a population of Simple cells that were recorded with microelectrodes. The visual receptive fields of the cells were located at 1-6^0 eccentricity. For each cell, responses were recorded for high-contrast, drifting sinusoidal gratings in different orientations (ori), spatial frequencies (SF) and temporal frequencies (TF). All measurements of Simple cell responses are in terms of the amplitude of the f1 component of neural firing rate in response to the drifting grating (15, 26). We also recorded visual responses of Complex cells in layer 4Cα but they did not exhibit DS and are not considered in this paper.

In the experiments, after the optimal orientation of each cell was determined, responses to the two opposite directions at the optimal orientation were measured with what we thought were stimuli that would produce the Maximum firing rate. If response in one direction was larger, that direction was called Preferred (abbreviated as Pref) and the other was the Opposite (to Preferred, or Opp) direction. Then we studied Pref and Opp responses over a range of TF and SF.

Using the ratio of Pref and Opp responses as a measure of DS, we present below statistics for two groups of cells: a “highly DS” group consisting of cells with Pref/Opp > 3, and a group with Pref/Opp > 2 which we refer to as “DS”. We required Pref/Opp > 2 to ensure that the preferred direction could be meaningfully identified.

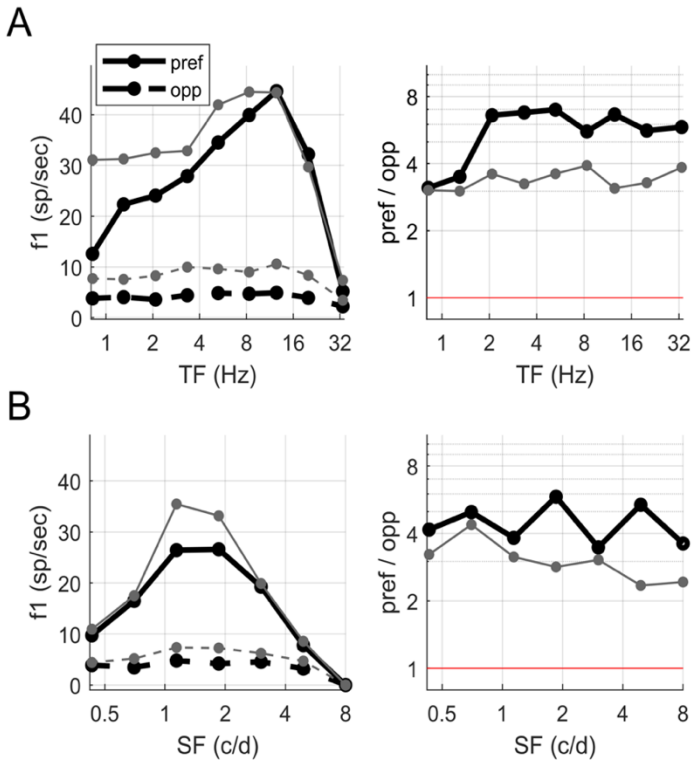

Figure 1. Data on firing rates and DS as functions of TF and SF. A. Data for 25 Simple cells with Pref/Opp > 3 (thick black) and 40 simple cells with Pref/Opp > 2 (grey) are presented. Left: Median Pref and Opp-firing rates as function of TF. Right: pref/opp as function of TF, for the same two groups of cells. Only cells for which firing rate at Max(pref,opp) > 4 spikes/s were included; below 10 Hz this condition was met by all the cells; by 32 Hz, # cells (n) dropped to 13 for Pref/Opp > 3 and to 23 for Pref/Opp > 2. **B.** Corresponding figures with SF in place of TF, for a collection of 26 Simple cells with Pref/Opp > 3 (thick black) and 42 simple cells with Pref/Opp > 2 (grey). Excluding low firing cells as in **A**, for Pref/Opp > 3, n = 24, 15, 3 at 3, 5, 8 c/d respectively; for Pref/Opp > 2, n = 39, 27, 8 at 3, 5, 8 c/d.

DS as function of TF

We have complete TF data for 53 Simple cells in layer 4Cα. Of these 53 cells, about half (25/53) were highly DS, while 75% (=40/53) were DS. The left panel of Figure 1A shows the medians of the Pref and Opp-responses for these two groups as functions of TF. Observe that for both groups, the median Pref-response is larger than median Opp-response consistently across the entire TF range. The Pref-response for the highly DS cells is temporally bandpass, with a much smaller response at 1Hz than at 10 Hz. Notice also that response amplitude drops off steeply at high TF so that at 32 Hz both Pref and Opp responses are very small and often below the noise level.

202 The right panel of Fig. 1A shows, for the same two groups of cells, the medians of the
203 pref/opp ratio as functions of TF. For both groups, these ratios are roughly constant throughout
204 the visible range of TF, demonstrating that in Macaque layer 4Cα, DS is broad-band in TF.
205
206 **DS as function of SF**
207 We have complete SF data for 54 Simple cells, of which 26 were highly DS and 42 were
208 DS. Figure 1B shows the same plots as in Figure 1A but as functions of SF, where the TF was at
209 or near optimal for the neuron. The pref/opp ratios were uniformly away from 1 throughout
210 the visible range of SF, implying that DS in layer 4Cα neurons is broad-band in SF, though
211 pref/opp declined gradually as SF increased from 1 to 5 c/d in the DS group.
212
213 To summarize: Our data show that most of the Simple cells recorded in layer 4Cα had
214 some DS, and roughly half had high DS, with Pref/Opp > 3. DS was found to be broad-band in
215 both TF and SF; each cell had a consistent preferred direction and roughly constant pref/opp-
216 ratios over the entire visible ranges of both TF and SF.
217
218
219 ## II. Proposed mechanisms for DS for ON-OFF LGN pairs
220 **Results II** considers the summed responses of two LGN cells, one ON and one OFF. We
221 propose mechanisms for how DS is produced, giving a complete analysis in this simplified
222 setting. Predictions from the theory developed here are then verified in **Results III** against
223 typical configurations of LGN afferents to cortical cells.
224
225 **ON-OFF pair setup**
226 We study here the sum of the responses of two LGN cells, one ON and one OFF,
227 representing two rows of ON and OFF LGN cells. Throughout the analysis, the OFF-cell is placed
228 on the left, the ON-cell is on the right, and they are separated spatially by d degrees, as shown
229 in Figure 2A. The stimuli L(x,t) considered are moving sinusoidal grating patterns; most
230 experimental data about DS in cortical neurons have been obtained with such stimuli, including

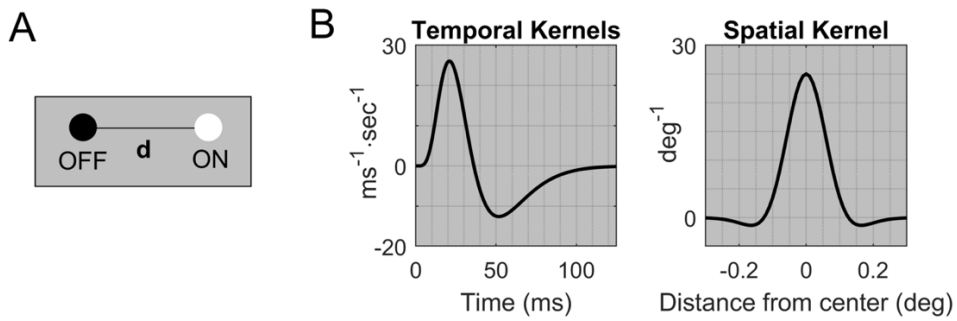

Figure 2. Setup for theory. A. A pair of ON and OFF LGN cells is separated by d°, with ON to the right of OFF. **B.** Reference temporal kernels K(t), and spatial kernel A(x,y). LGN response is assumed to be the convolution of the stimulus function with these two kernels.

those in **Results I.** Our theory generalizes to drifting bars and edges, but that will not be discussed here.

We consider only gratings aligned with the rows of ON and OFF cells, so that in the simplified setting of an ON-OFF pair they are either left or right-moving. Drifting gratings are **Right** (from left to right) when L(x,t) = C sin(2π(-gx+ft)), and **Left** (from right to left) when L(x,t)=C sin(2π(gx+ft)); here g (units: c/deg) is spatial frequency (SF), f (units: Hz) is temporal frequency (TF), and C is contrast. Throughout the discussion, we assume the responses $R_{ON}(t)$ and $R_{OFF}(t)$ of the ON and OFF-cells are given by convolutions of L(x,t) against the cell's spatial and temporal kernels, i.e.,

$$\mathrm{R_{ON}}(t), \mathrm{R_{OFF}}(t) = \pm \iiint K(s)A(x_0 - x, y)L(x, t - s)dxdyds,$$

where K(s) is the temporal kernel and A(x,y) is the spatial kernel. The cells are located at $(x_0,0)$; $x_0 = 0$ for OFF and $x_0 = d$ for ON. The forms of the temporal and spatial kernels are illustrated in Figure 2B; exact formulas and other details are given in **Materials and Methods**. For the reference temporal kernel K(s), we have chosen the cross-over point from positive to negative to lie at 36 ms so that the peak response occurs at about TF=10 Hz following the data reported in Figure 1A, left.

Of interest is $R(t) = R_{ON}(t) + R_{OFF}(t)$, the sum of the ON and OFF responses.

255 **DS via differential in phase differences**

256 We start with the following questions: Does the summed response R(t) of the ON-OFF pair
257 have a preferred direction, and what is the preferred direction? The answers are clear in the
258 frequency domain, an insight that goes back to Watson and Ahumada (27, 28). DS depends on
259 the phase difference between $R_{ON}(t)$ and $R_{OFF}(t)$, denoted $\Delta\varphi$, which takes values between 0
260 and π. The closer $\Delta\varphi$ is to 0, the larger the amplitude of R(t) because then ON and OFF
261 responses are in phase and reinforce each other. The closer $\Delta\varphi$ is to π, the smaller is R(t)'s
262 amplitude because then ON and OFF are in antiphase and cancel each other.

263 If ON and OFF cells have the same response time-course, then there is no DS (27, 28, 19).
264 In this case, $\Delta\varphi(\rightarrow) = \Delta\varphi(\leftarrow)$ where $\Delta\varphi(\rightarrow)$ is the ON-OFF phase difference for a right-moving
265 grating, and $\Delta\varphi(\leftarrow)$ for motion to the left. But if the symmetry of ON and OFF is broken and
266 $\Delta\varphi(\rightarrow) \neq \Delta\varphi(\leftarrow)$, the phase difference of one of the directions is closer to 0, and that is the
267 preferred direction.

268

269 ***Mechanism #1. Delay of ON-response relative to OFF***

270 A delay in the ON-response with respect to OFF is consistent with retinal circuitry required
271 for the sign inversion in the ON pathway (29) and with direct measurements (30, 31). We
272 consider a delay of t_0 ms in the response of the ON-cell, and explain how that leads to unequal
273 phase differences for the right and left-moving gratings.

274 To illustrate how the introduction of a delay breaks the symmetry between $\Delta\varphi(\rightarrow)$
275 and $\Delta\varphi(\leftarrow)$, we show in Figure 3 numerical simulations of the response of an ON-OFF pair to
276 drifting gratings. For those simulations, f=10 Hz, around the peak TF for Magnocellular cells (13)
277 and g=2.5 c/deg, at or near an optimal SF for many cells in V1 representing the parafovea (32)
278 which is the region we are modeling. The distance d (Fig. 2A) between ON and OFF is set at 0.1°,
279 a first approximation based on retinal topography (33).

280 In Figure 3A, there is zero delay. ON and OFF responses are shown separately (in white
281 and black), and their sum is shown in red. The responses to left and right motion are identical
282 with the roles of ON and OFF interchanged, resulting in the same summed response amplitude
283 for Left and Right. In Figure 3B, a delay t_0 between ON and OFF is introduced (left panel). Data
284 (30, 31) suggest that t_0 is likely between 5-15 ms. Here we use t_0 = 10 ms. The summed

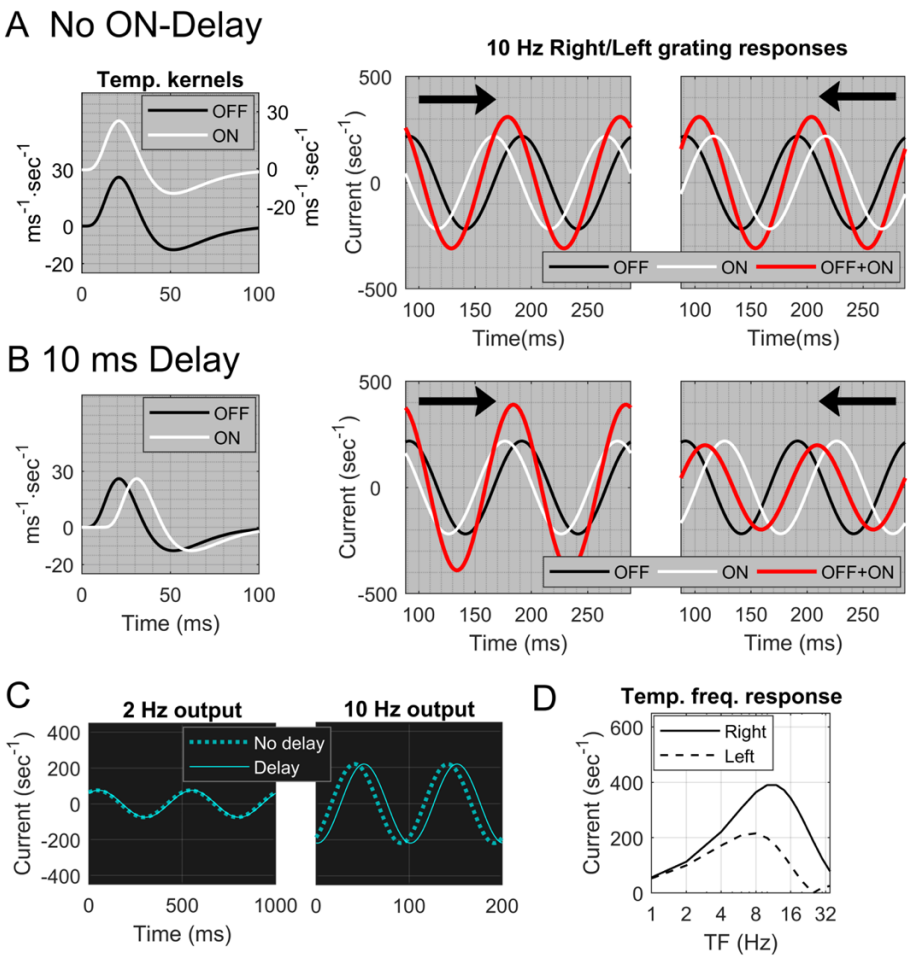

Figure 3. Effect on DS from time delay between ON and OFF LGN cells. **A.** Two identical temporal kernels, one OFF (black) and one ON (white) (leftmost panel). The ON kernel is displaced vertically to avoid overlap with OFF. The two panels on the right show responses of an ON-OFF pair separated by d=0.1^0 to gratings at 10Hz and 2.5 c/deg. Waveforms of ON (white), OFF (black) and summed response ON+OFF (red), with arrows indicating the direction of motion. **B.** Two identical kernels, one OFF (black) and one ON (white), with a 10 ms delay for the ON-kernel, are shown in the leftmost panel. The two panels on the right show results analogous to those in **A**, with an ON-delay of 10 ms. **C.** The two panels show the responses of the ON-kernel with a 10 ms delay (solid) compared to no-delay (dashed) at TF=2Hz and 10 Hz respectively. **D.** Dependence on temporal frequency (TF) of the effect of delay on DS. Plotted is the F1 component of the ON+OFF sum as a function of TF for both Right (solid) and Left (dashed) directions.

response R(t) is larger for Right motion, for which the ON and OFF responses are closer to being in-phase (Figure 3B); Right is the preferred direction.

To understand Figure 3B we analyze ON/OFF phase differences in the Right and Left directions. For Right, ON lags OFF because of the separation d and the delay t_0; there is also a

phase difference of π $(=180^0)$ because of the sign difference between ON and OFF. Thus, $\Delta\varphi(\rightarrow) = [2\pi(- gd - ft_0) + \pi]$ where we use the formalism that [x] denotes the shortest distance between x and the set $\{0, \pm 2\pi, \pm 4\pi, \ldots\}$. (The phase difference is distance between two points on a circle and all multiples of 2π are equivalent to 0.) For Left, $\Delta\varphi(\leftarrow) = [2\pi(gd - ft_0) + \pi]$; in the Left direction, the ON cell has a phase lead due to the separation d, but a phase lag due to the delay. Response amplitudes in the two directions are predictable from the phase differences: here $\Delta\varphi(\rightarrow)$ is smaller so Right is the preferred direction.

Because the delay enters $\Delta\varphi$ as $2\pi f t_0$, ON time-delays are effective in producing DS at higher TF, but not at lower TF (Figure 3D). For instance, a grating with TF=10 Hz travels 1/10 of a cycle in 10 ms, whereas at TF=2 Hz, 10 ms is only 1/50 of a cycle. Figure 3C shows the phase shifts of an ON-cell due to a 10-ms delay in response to gratings with TF = 2 and 10Hz. The 10Hz grating produced significant DS (Figure 3B). However, for the 2Hz grating, the phase shift is negligible and the picture is closer to that of no-delay; it does not produce much DS (Figure 3D).

The data in Figure 1A show, however, that DS is present both at high and low TF in the Macaque. We therefore need another mechanism in addition to the time-delay.

Mechanism #2. Modified structure of ON-kernels

Another suggestion from LGN data (30) is that the waveforms of ON and OFF time-kernels have different temporal structure. We use the following notation: Let the OFF-kernel $K_{OFF}(t)$ = K(t) where K(t) is as in Figure 2B (see **Online Methods** for more detail), and say a temporal kernel $K_{a,b}(t)$ has structure (a, b) if

$$K_{a,b}(t) = \begin{cases} a\,K(t) & t < 36\ ms \\ b\,K(t) & t > 36\ ms \end{cases} .$$

In this notation, $K_{OFF}(t)$ has the structure (1,1). By analysis and computer simulations we established that a kernel $K_{a,b}(t)$ with $a \neq b$ produces a phase shift relative to $K_{1,1}(t)$. For $a>b$, it is a <u>phase lag</u>, and the lag is more prominent at low TF and declines as TF increases (Figure 4A). Other kernels $K_{a,b}(t)$ with $a>b$ produce similar results. In general, the larger a/b, the larger the shift. Ratios of a/b ~ 2 are consistent with data for ON-kernels (30).

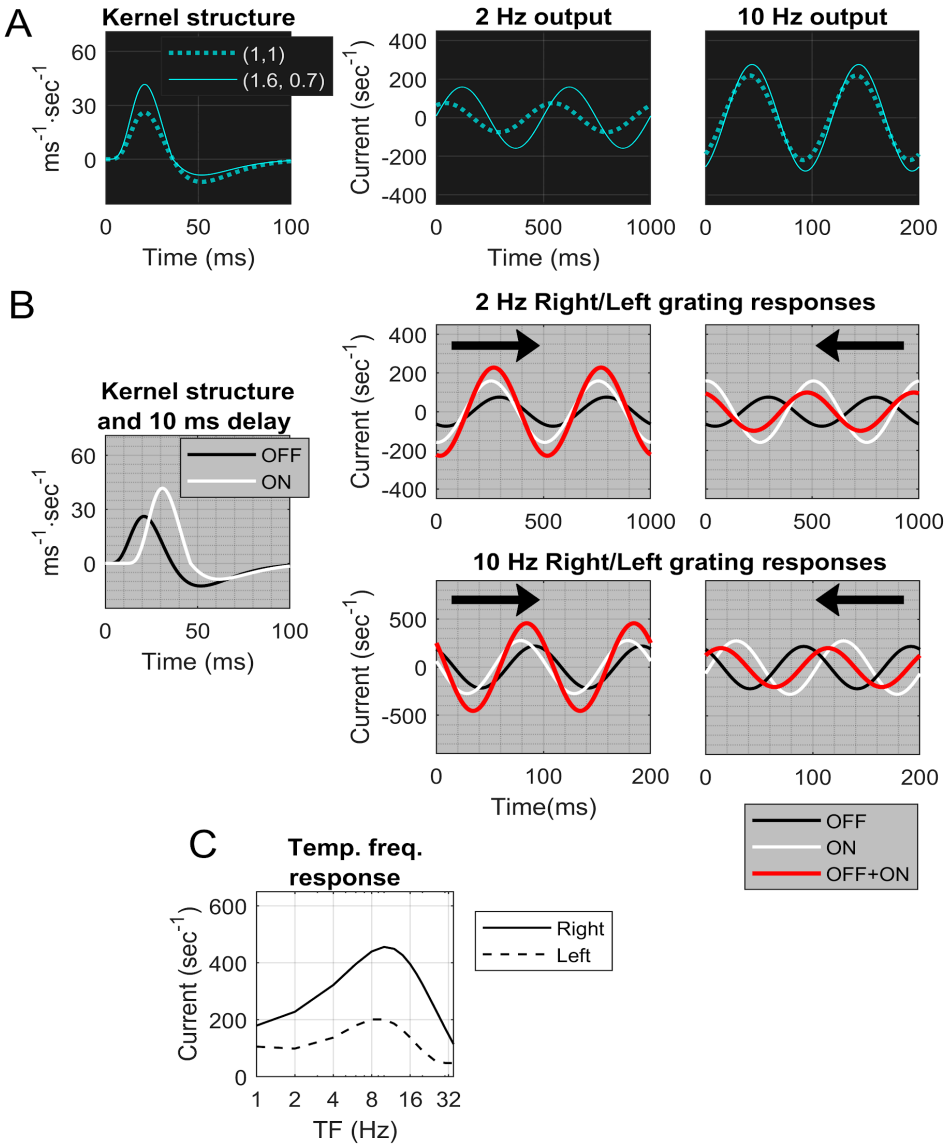

Figure 4. Effect on DS of time delay and modified temporal kernel structure. **A.** The left panel shows a kernel of type (1.6, 0.7) (solid) and one of type (1,1) (dashed). The two panels on the right show the phase lags of the (1.6, 0.7)-kernel (solid) relative to one of type (1,1) (dashed) at TF = 2 Hz and 10 Hz. **B.** The setup is like that of Figure 1; d = 0.1^0 and SF=2.5 c/deg. The ON-kernel is of type (1.6,0.7) in addition to having a 10 ms delay with respect to OFF. In the righthand panels the ON responses (white) and OFF responses (black) have greater phase differences for Left than for Right. Summed waveforms (red) in Right(-->) and Left(<--) directions for TF=2Hz and TF=10 Hz reveal Right preference at both TFs. **C.** Right vs Left responses as functions of TF, at SF=2.5 c/d. It is the analogue of Fig 1D but with a (1.6, 0.7) ON-kernel in addition to the 10 ms delay.

Combining Mechanisms #1 and 2, i.e., letting $K_{OFF}(t)= K_{1,1}(t)$, and letting the ON-kernel $K_{ON}(t)= K_{a,b}(t)$ for some a>b *with* a delay of t_0 ms, we may write $\Delta\varphi\ (\rightarrow)$ and $\Delta\varphi\ (\leftarrow)$ as

$$\Delta\varphi(\rightarrow) = [2\pi(-gd - ft_0 - p(f,K_{ON})) + \pi] \quad \text{and} \quad \Delta\varphi(\leftarrow) = [2\pi(gd - ft_0 - p(f,K_{ON})) + \pi]$$

where $p(f,K_{ON})>0$ is the phase lag contributed by the temporal structure of the ON kernel, and [ ] has the same meaning as before.

From the discussion above, it follows that Mechanisms #1 and 2 combined would produce DS in the summed response of an ON-OFF pair across a wide range of TF with a consistent preferred direction. Simulations confirm the correctness of this reasoning. Figure 4B shows the individual and summed responses for left and right-moving gratings, where the ON-kernel has temporal structure (1.6, 0.7) in addition to a time delay of 10 ms, for both f=2Hz and f=10Hz. The response shows a clear preference for Right motion.

Figure 4C confirms these results across the full range of TF. We compare this panel to Figure 1A: Figure 4C is our theoretical prediction of the summed *feedforward inputs* to a cortical cell (for left and right gratings), while Figure 1A, left, shows data on the *responses* of DS cortical cells. Similarities in the shapes of the input and output curves as functions of TF and the fact that pref/opp for both are roughly constant over the entire range of TF (Figures 4C and 1A, right) lend credibility to the theory proposed.

Preferred direction depends on d and SF (equivalently RF size)

In Figures 3 and 4, the preferred direction is from OFF to ON (Right), but is that always the case? If not, then is there a direction that is consistently preferred by an ON-OFF pair, or does the preference vary with stimulus? To answer these questions systematically involves examining the relation between the spatial frequency (SF) of the grating and the physical separation d between the ON and OFF pair.

A crucial observation is that if we fix d and vary SF, then the preferred direction switches at gd = 1/2, 1, 3/2 … where d is ON/OFF separation and g is SF. Specifically, in our OFF-ON configuration with OFF on the left, when gd < 1/2, i.e., when d < 1/2 the spatial cycle of the grating, Right motion is preferred (this was the case in the examples shown in Figures 3 and 4).

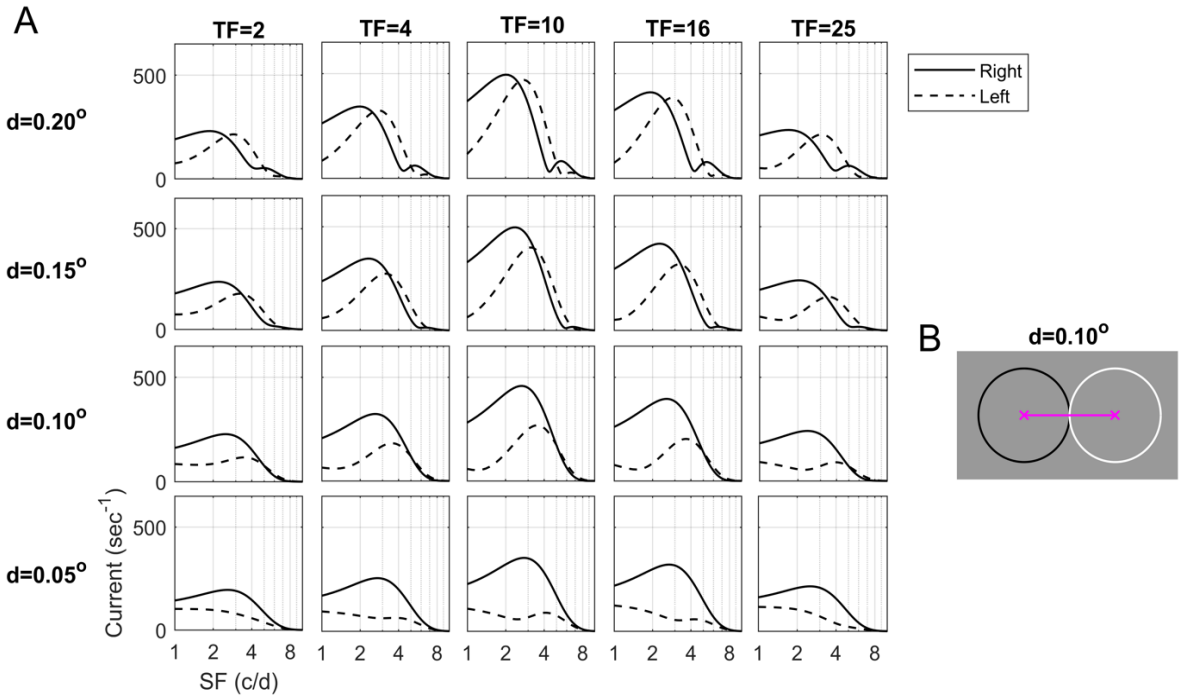

Figure 5. Dependence of the DS of the ON-OFF pair on SF and d. **A**. Matrix of response vs SF plots is drawn parametric in d and TF. Rightward and Leftward responses are plotted vs TF for 4 different values of d: 0.05, 0.1, 0.15, 0.20^0. **B.** Two LGN cells separated by d=0.1^o. The circles are 1 SD of their spatial kernels.

For 1/2 < gd < 1, Left motion is preferred, and directional preference is reversed again as gd increases past 1. We found that preferred directions are determined solely by the relation between d and SF, and are independent of the TF of the grating, provided that the temporal shift $ft_0 + p(f,K_{ON})$ is < π, a condition easily satisfied by the mechanisms proposed. A mathematical proof for the assertions in this paragraph is given in **Materials and Methods**.

Figure 5A illustrates the consequences of these predictions by showing Right and Left responses as functions of SF, for four different values of d, and five TFs. In the rows in which d=0.15 or d = 0.2 deg, the predicted directional switch can be seen as SF increases in each plot: When g < 1/(2d), the preferred direction is from OFF to ON (Right), and when g > 1/(2d), it is from ON to OFF (Left), and this is true across the entire range of TF. Reversals for d=0.1 are less visible because ON/OFF separation of d=0.1^o corresponds to reversals at 5 c/d, where the

curves have steep slopes. As for d=0.05°, reversals are predicted to occur at 10 c/d, a high enough SF that the response is too small to be visible.

The following points are illustrated in Figure 5A. One is that while, mathematically, direction reversals occur at gd = 1/2, 1, 3/2 …, as a visual response a reversal is registered only if the SF at which it occurs falls within the animal's range of visible SF, which is known to scale with rf size (34). A second point is that as important as the phenomenon of reversal itself is the loss of DS near reversal points. The lowering of Pref/Opp for an entire region of (d, SF) near reversal occurs as a result of the gradual interchange of preferred directions.

That leads to our third point, the prediction that DS is stronger for smaller ON/OFF separation. In Figure 5A, peak firing rate occurs at TF=10 Hz and SF = 2-3 c/d, a good approximation to data (Figure 1). Observe that at these values of TF and SF, Pref/Opp is close to 1 at d=0.2°, a little larger at d=0.15°, and continues to increase as d gets smaller. The Pref/Opp ratio is larger for smaller d because, with decreasing d, reversal occurs farther from SF=2-3 c/d.

Implications for the Macaque

In Macaque V1, we have not observed evidence for reversals of Preferred Direction (Figure 1B, right), suggesting that reversal occurs outside of the range of SF that produces a response. This range can be seen in Figure 1B, left. The discussion above together with the results in Figure 1 predict therefore that to provide feedforward input with robust DS across the full range of SF, ON-OFF rows that project to the same cortical cell are likely < 0.1° apart at the eccentricities of the cells recorded. The decline in DS for SF between 1 and 5 c/d seen in Figure 1B suggests also that we may be heading for a reversal point, so this ON-OFF separation is likely > 0.05°.

Comparing theory to data, our theoretical prediction of the summed LGN feedforward inputs to cortical cells at TF=10 Hz (the optimal TF) and d = 0.05-0.1° as shown in Figure 5 shows a remarkable resemblance to the data on cortical responses shown in Figure 1B, left.

423 ## III. DS in feedforward LGN inputs

424 In **Results II**, we developed a theory for the DS of an ON-OFF pair. We now extend this
425 theory to typical feedforward LGN inputs to layer 4Cα, assuming that the set of LGN cells that
426 project to a simple cell in layer 4Cα is organized in two or three rows of ON and OFF cells,
427 alternating in ON/OFF in the case of three rows. These widely accepted rules of LGN-to-cortex
428 connections were proposed by Hubel and Wiesel and have been confirmed in experiments (35,
429 25, 36, 37). The setup is as described at the beginning of **Results II**, but for arbitrary collections
430 of LGN cells. The response of each LGN cell is computed as a convolution of the light intensity
431 function with spatial and temporal kernels as before, and DS for the summed responses from
432 the entire group is evaluated.

433 LGN sampling of visual space is known to be very sparse; two independent estimates
434 based on M-retinal ganglion cell density (33) and on Macaque LGN data (38) point to an
435 average of about 9 LGN cells projecting to an area of 0.25^o x 0.25^o in the visual field at 5°
436 eccentricity. Respecting this density, we modeled a sheet of LGN cells using either a randomly
437 perturbed hexagonal lattice (Figure 6 A-C) or a randomly perturbed rectangular lattice (Figure 6
438 D-E). Once the ON and OFF mosaics were fixed, the temporal kernel of each LGN cell was drawn
439 randomly and independently following the prescription from **Results II**.

440 We consider in Figure 6 some typical configurations of LGN cells that form the inputs to a
441 cortical cell. Following (39), it is likely that no more than 4-6 LGN cells project to a Simple cell,
442 and we learned from **Results II** that ON/OFF separation d should be $< 0.1^o$, with stronger DS
443 predicted for smaller**.**

444 The configurations in Panels A and B translate easily to the ON-OFF setting studied in
445 **Results II:** projecting along a direction orthogonal to the rows of ON and OFF cells, we arrive at
446 precisely the picture treated in **Results II** with near-optimal spacing between the two cells. As
447 expected, DS of the summed responses of these configurations are strong, and as predicted,
448 the preferred direction is from OFF to ON.

449 Panels C and D require extensions of the ideas developed for ON-OFF pairs in **Results II**. In
450 Panel C, the projected configuration has three cells, an ON-cell flanked by two OFFs. DS for this

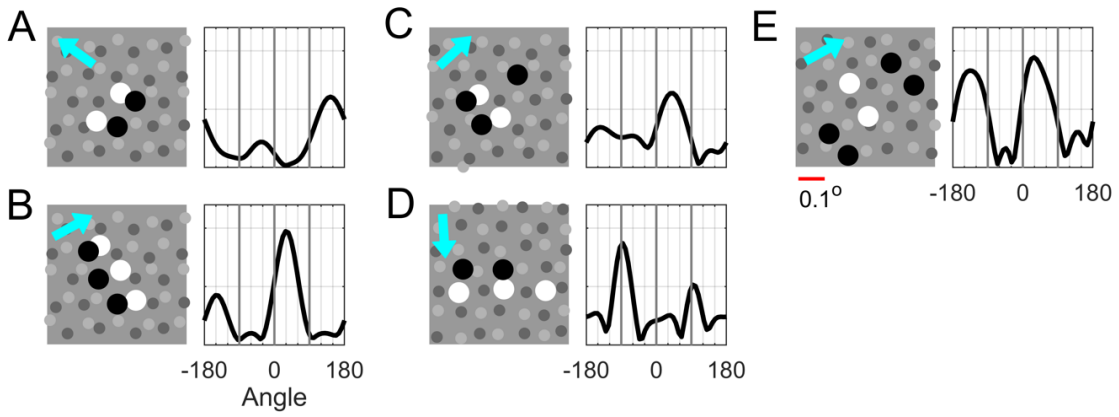

Figure 6. Sample LGN configurations and orientation (ori) tuning of their summed responses. Shown are 5 examples (**A-E**) of configurations of LGN cells that project to a simple cell in layer 4Cα (left) and ori tuning curves of their summed responses (right). Grey dots indicate locations of LGN cells present: ON in lighter grey, OFF in darker grey. Larger black and white dots are the OFF and ON cells in the configuration. LGN kernels are randomly drawn, and are of type (a,b): for OFF, a,b ∈ (0.9,1.1), and for ON, a ∈ (1,2) with a/b ∈ (1.4, 2.2); ON-delay time ∈ (9,11) ms, also randomly drawn. In ori tuning curves for summed responses, cyan arrow indicates preferred direction according to tuning curves. Scale bar of 0.1 deg shown below Panel E.

3-cell configuration can be thought of as competition for DS-dominance between the OFF-ON pair on the left and the ON-OFF pair on the right; one should also expect cancellation as these two pairs will prefer opposite directions. In the configuration shown in Panel C, one would predict that the left OFF-ON pair should prevail, because it carries more weight and ON/OFF distance is smaller. The orientation tuning curve shown confirms these predictions: the left ON-OFF pair is dominant, and DS is weaker due to partial cancellation. Panel D projects to an ON-OFF pair, with the ON-cell carrying significantly more weight. That weakens the DS, though the preferred direction is still quite evident.

The configuration in Panel E projects to an OFF-ON-OFF triplet that is left-right symmetric; no DS can be expected for such a configuration and little is observed.

Figure 6 and other examples not shown illustrate the fact that the mechanisms proposed in **Results II** lead to variable amounts of DS in feedforward LGN inputs depending on the spatial alignment of the group of LGN cells (and their temporal kernels). The conclusion about variable amounts of DS is consistent with the data presented in **Results I**, which show

476 that not all 4Cα neurons are DS, and among those that are DS, Pref/Opp can take on a range of
477 values; see Figure 1.
478
479
480

481 # Discussion

482 The main contributions of this paper can be summarized as follows:
483
484 (1) We presented experimental data to document systematically the dependence of DS on
485 TF and SF for neurons in layer 4Cα of Macaque V1, and proposed a theory the
486 predictions of which are consistent with these data.
487
488 (2) Our theory asserts that DS could originate from dynamic differences between ON and
489 OFF cells and should therefore already present in feedforward LGN inputs to cortex.
490 Two mechanisms the combined action of which is sufficient for initiating DS across the
491 full spectrum of TF were proposed.
492
493 (3) We made precise how stimulus features such as TF and SF, the dynamics of ON/OFF LGN
494 cells, and LGN-to-cortex wiring interact to determine the preferred direction and
495 magnitude of DS in feedforward LGN inputs.
496
497 We do not claim that the mechanisms proposed are the only mechanisms that contribute
498 to DS. The presence of specific intra-cortical connections that enhance or modify DS, for
499 example, would not contradict our theory, especially in layers beyond 4Cα where longer-range
500 connections are present; see e.g. (14). Our theory asserts only that direction selectivity could
501 *originate* from dynamic differences between ON and OFF LGN cells.
502 Previous theoretical work has proposed spatio-temporal inseparability in x-t plots as a
503 characterization of DS (19). Findings of such inseparability have been obtained for V1 neurons
504 in experimental studies (40, 22). While x-t plots in and of themselves are not intended to shed
505 light on mechanisms, spatio-temporal bias is an indication of DS. In **Supplemental Information**

(SI) we will show that when the mechanisms proposed in **Results II** are implemented, the summed responses of ON-OFF pairs of LGN cells generate x-t plots resembling those seen in data in the papers above. The converse need not be true, however: x-t inseparability need not imply DS under certain stimulus conditions, a fact we will also demonstrate in **SI**.

We finish with the following comparison of our theory with theories of DS in mouse V1 cortex. A neural mechanism for DS proposed in previous modeling studies was for specific connections between V1 cells and LGN cells that are either Sustained or Transient in their step responses (23, 41). The Sustained-Transient concept was revived recently in models of DS in mouse V1, by Lien and Scanziani (15) and Billeh et al (16). We consider the Sustained-Transient theory to be implausible for Macaque because there are not clearly separate classes of Sustained and Transient Magnocellular LGN neurons. In our proposed theory, ON vs OFF LGN cells have different temporal kernels, but these differences are much smaller and subtler than those observed in Sustained and Transient neurons in rodent or cat retinal and LGN cells (16). Even accepting Sustained/Transient differences as underlying mechanisms of DS for mouse, the story there is incomplete without information on the wiring of these two subpopulations and how that leads to the DS properties observed. In the mouse, estimates are that there are either ten (16) or forty (15) LGN cells converging into one subfield. The problem of how to place all (or most) of the Sustained inputs into one subfield and Transient inputs into the other remains to be clarified. Our theory solves this problem by postulating that ON and OFF cells have different temporal impulse responses. That together with the consensus view on how to segregate ON and OFF inputs from LGN into different subfields (24, 25) is sufficient for generating DS.

Materials and Methods

Modeling LGN response

We model the input current to an LGN cell the center of whose receptive field is located at $(x_{0,}\, y_0)$ by

$$I_{\pm}(t) = [I_B \ \pm Q(t)]^{+}$$

where

$$Q(t) = \iiint K(s)A(x_0 - x, y_0 - y)L(x, y; t - s)dxdyds \ .$$

Here the $\pm$ sign is for ON and OFF-cells respectively, I_B is background current, $K(s)$ and *A(x,y)* are the temporal and spatial kernels of the LGN cell, *L(x,y;t)* is the light intensity map, and C is a constant to be adjusted, and [ ]$^+$ denotes the maximum of the bracketed value and 0.

In **Results II**, the OFF cell is placed at $(x_{0,}\, y_0) = (0,0)$, the ON cell is placed at $(x_{0,}\, y_0) = (d, 0)$, and L(x,y;t) depends only on x and t. The more general formulas above are used in **Results III**. In both **Results II and III,** we have used $\pm Q(t)$ as our ON/OFF LGN *response*, conflating the input current to LGN with its output (in addition to omitting the background firing and the [ ]$^+$ part of the operation). This conflation was on purpose: the two quantities are roughly proportional, and the exact sinusoidal form of $\pm Q(t)$ makes the analysis much more transparent.

Formulas for the spatial and temporal kernels are as follows: The spatial kernel is a difference of Gaussians

$$A(x,y) = (a/(\pi\sigma_a^2)) * \exp(-(x^2+y^2)/\sigma_a^2) - (b/(\pi\sigma_b^2)) * \exp(-(x^2+y^2)/\sigma_b^2)$$

with a=1.0, b=0.74, σ_a=0.0894, σ_b=0.1259 (42); σ_a and σ_b are chosen to make the center part of the receptive field correspond to a Gaussian with std. dev. ~ 0.05 deg (13). The preferred spatial frequency of the spatial kernel *A(x,y)* is 2.5 c/deg. The reference temporal kernel K(t), from which ON- and OFF-kernels are derived, has the form (adapted from Zhu et al (42))

$$K(t) = (t^6/\tau_0^7) * \exp(-\, t\,/\tau_0) \ - \ (t^6/\tau_1^7) * \exp(-\, t\,/\tau_1)$$

with τ_0 = 3.66 ms and τ_1 = 7.16 ms. Parameters are chosen following guidance from temporal kernels derived in experiments (30) and the temporal frequency response of LGN cells, ensuring the peak response occurs at ~10 Hz (13). Note that K_{OFF} is positive before the zero crossing, as the polarity of the LGN cell is implemented in the sign of +/- in the expression for I(t), not in the kernel.

Mathematical analysis of reversal of directional preference

In **Results II,** we asserted that preferred direction switches at gd = 1/2, 1, 3/2 … where d is ON/OFF separation and g is the SF of the grating. A mathematical proof goes as follows. Look at the quantities inside the brackets [ ] in the formal expression for phase differences derived in **Results II,** and recall that without the temporal phase lag, the spatial phase lead for ON is $-2\pi gd + \pi$ for Right-moving gratings and $2\pi gd + \pi$ for Left-moving gratings. Consider first the case gd < ½ (Figure 7A), where $-2\pi gd + \pi$ is between 0 and π, and $2\pi gd + \pi$ is between π and 2π. These phase leads are depicted as dashed red arrows. The effect of the temporal phase lag for ON is to shift these arrows to the left, to the solid red arrows depicting the net phase leads for ON. Provided that the temporal phase shift ft_0+p(f) is less than π (the mechanisms proposed produce shifts < $\pi/2$), we have $\Delta\varphi(\rightarrow) < \Delta\varphi(\leftarrow)$, because the red arrow for Right is closer to 0 than the red arrow for Left is to 2π. Hence Right motion is preferred.

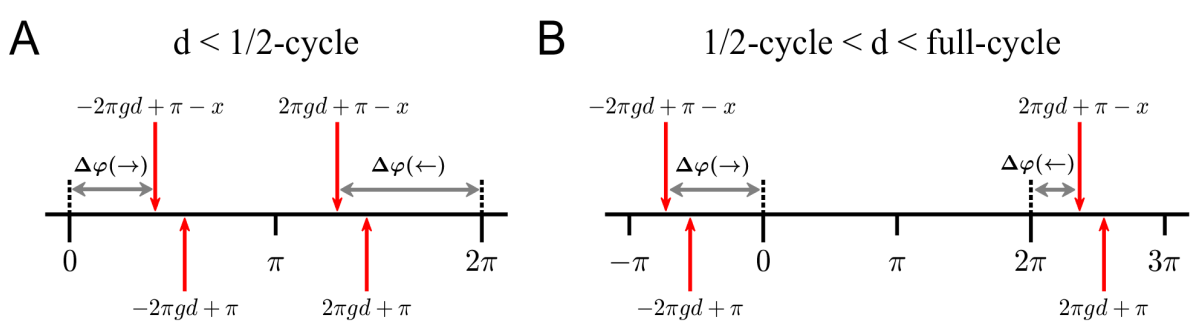

Figure 7. Mathematical analysis of interplay between d and SF. **A**. Phase diagram when separation d < 1/2 cycle of the stimulus grating. Dashed red arrows are spatial phase leads for ON without taking into account temporal phase lags; solid red arrows take into temporal phase lags. Red arrows between 0 and π are for Right gratings; arrows between π and 2π are for Left. The quantity $x = ft_0$+p(f,K_{ON}) is temporal phase shift, and grey lines with arrowheads are $\Delta\varphi(\rightarrow)$ and $\Delta\varphi(\leftarrow)$. **B.** Phase diagram when 1/2-cycle < d < full cycle.

586 The picture is different for ½ < gd < 1 (Figure 7B). For this range of d, $-2\pi gd + \pi$ lies
587 between $-\pi$ and 0, and $2\pi gd + \pi$ lies between 2π and 3π. Here the temporal phase lag causes
588 $\Delta\varphi(\rightarrow) > \Delta\varphi(\leftarrow)$, resulting in a preference for Left motion.
589 A similar argument shows that preference is reversed again at gd = 1.
590
591
592 **Experimental Methods**
593 *Animal Preparation.* Adult male old-world monkeys (M. fascicularis) were used in acute
594 experiments in compliance with National Institutes of Health and New York University Animal
595 Use Committee regulations. The animal preparation and recording were performed as
596 described in detail previously (43, 44). Anesthesia was initially induced using ketamine (5-20
597 mg/kg, i.m.) and was maintained with isofluorane (1-3%) during venous cannulation and
598 intubation. For the remainder of the surgery and recording, anesthesia maintained with
599 sufentanil citrate (6-18 μg/kg/h, i.v.). After surgery was completed, muscle paralysis was
600 induced and maintained with vecuronium bromide (Norcuron, 0.1 mg/kg/h, i.v.) and anesthetic
601 state was assessed by continuously monitoring the animals' heart rate, EKG, blood pressure,
602 expired CO_2, and EEG.
603 After the completion of each electrode penetration, 3–5 small electrolytic lesions (3 μA
604 for 3 s) were made at separate locations along the electrode track. At the end of the
605 experiments, the animals were deeply anesthetized with sodium pentobarbital (60 mg/kg, i.v.)
606 and transcardially exsanguinated with heparinized lactated Ringer's solution, followed by 4 L of
607 chilled fresh 4% paraformaldehyde in 0.1 M phosphate buffer, pH 7.4. The electrolytic lesions
608 were located in the fixed tissue and electrode tracks were reconstructed to assign the recorded
609 neurons to cortical layers as described previously (8). The data from the current study were
610 obtained during the course of the basic characterization of neuronal receptive field properties
611 that is standard for many visual cortical experiments. In 49 animals, more than 700 neurons
612 were recorded in oblique penetrations. For this paper, a full data set was obtained for 94
613 neurons localized to layer 4Cα.
614 *Characterization of visual properties of V1 Neurons***.** We recorded action potentials
615 (spikes) extracellularly from single units in V1 using glass-coated tungsten microelectrodes.

Each single neuron was stimulated monocularly through the dominant eye (the non-dominant eye occluded). Receptive fields were located in the visual field between 1 and 6^0 from the center of gaze. Stimuli were displayed at a screen resolution of 1024x768 pixels, a refresh rate of 100 Hz, and a viewing distance of 115 cm on either a Sony Trinitron GDM-F520 CRT monitor (mean luminance 90-100 cd/m^2) or an Iiyama HM204DT-A CRT monitor (mean luminance 60 cd/m^2). Monitor luminance was calibrated using a Photo Research PR-650 spectroradiometer and linearized via a lookup table.

The response to drifting gratings was used to characterize visual response properties. In this paper, we report measurements of orientation tuning, spatial frequency (SF) and temporal frequency (TF) tuning. We used the f1 response for neurons (traditionally called simple cells) with f1/f0 ratios > 1.

Orientation tuning: The responses of each neuron were recorded to different orientations between 0 – 360^0, either in 20 or 15^0 steps. The stimuli were achromatic gratings at the preferred SF and TF, at a contrast of 64% or greater. All stimuli were presented in a circular window confined to the classical receptive field (CRF).

SF tuning: Each neuron was presented with a range of spatial frequencies, usually in ½ octave steps from 0.1 c/deg to around 10 c/deg. For some neurons, the upper limit was extended if the neurons had responses to higher spatial frequencies. For almost all neurons that were orientation selective we measured SF tuning at the preferred orientation and drift-direction as well as at the non-preferred drift direction.

TF tuning: Each neuron was presented with a range of temporal frequencies, usually in 1 octave steps from 0.5 Hz to 32 Hz. Measurements were made for drifting gratings at the optimal orientation and in the preferred and non-preferred directions.

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