A comparison of koniocellular, magnocellular and parvocellular receptive field properties in the lateral geniculate nucleus of the owl monkey (Aotus trivirgatus)

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1. By analogy to previous work on lateral geniculate nucleus (LGN) magnocellular (M) and parvocellular (P) cells our goal was to construct a physiological profile of koniocellular (K) cells that might be linked to particular visual perceptual attributes.

2. Extracellular recordings were used to study LGN cells, or their axons, in silenced primary visual cortex (V1), in nine anaesthetized owl monkeys injected with a neuromuscular blocker. Receptive field centre–surround organization was examined using flashing spots. Spatial and temporal tuning and contrast responses were examined using drifting sine-wave gratings; counterphase sine-wave gratings were used to examine linearity of spatial summation.

3. Receptive fields of 133 LGN cells and 10 LGN afferent axons were analysed at eccentricities ranging from 2.8 to 31.3 deg. Thirty-four per cent of K cells and only 9% of P and 6% of M cells responded poorly to drifting gratings. K, P and M cells showed increases in centre size with eccentricity, but K cells showed more scatter. All cells, except one M cell, showed linearity in spatial summation.

4. At matched eccentricities, K cells exhibited lower spatial and intermediate temporal resolution compared with P and M cells. K contrast thresholds and gains were more similar to those of M than P cells. M cells showed lower spatial and higher temporal resolution and contrast gains than P cells.

5. K cells in different K LGN layers differed in spatial, temporal and contrast characteristics, with K3 cells having higher spatial resolution and lower temporal resolution than K1/K2 cells.

6. Taken together with previous results these findings suggest that the K cells consist of several classes, some of which could contribute to conventional aspects of spatial and temporal resolution.

In the primate lateral geniculate nucleus (LGN), three principal types of relay cells have been identified: the koniocellular (K), the parvocellular (P) and the magnocellular (M) cells. These relay cell classes can be distinguished based upon a number of criteria including laminar location, morphology, connections and neurochemistry in several primate species including bush babies, owl monkeys, marmosets and macaque monkeys (see Casagrande, 1984, 1999, Hendry & Reid, 2000 for review). K LGN cells are, on average, the smallest relay cells. They form thin layers that lie below the M and P layers, they contain the calcium-binding protein calbindin-D28k, and they send their axons to the cytochrome oxidase (CO) blobs in cortical layer III and to cortical layer I of the primary visual cortex (V1) (Lachica & Casagrande, 1992; Hendry & Yoshioka, 1994; Johnson & Casagrande, 1995; Ding & Casagrande, 1997). In contrast, M and P LGN cells are large and medium in size, are located in the ventral and dorsal layers of the LGN, contain the calcium-binding protein parvalbumin and send their axons principally to the upper and lower tiers of cortical layer IV of Brodmann (1909), respectively (Jones & Hendry, 1988; Lachica et al. 1992; Hendry & Yoshioka, 1994; Johnson & Casagrande, 1995; Ding & Casagrande, 1997; Goodchild & Martin, 1998).

The physiology of only two of these three LGN cell classes, the M and P cells, has been studied in detail across primate species (e.g. bush baby: Norton & Casagrande, 1982; Irvin et al. 1986, 1993, Norton et al. 1988; macaque monkey:...
Kaplan & Shapley, 1982; Derrington & Lennie, 1984; Hubel & Livingstone, 1990; Reid & Livingstone, 1992; Spear et al.
1994; owl monkey: Sherman et al. 1976; O'Keefe et al.
White et al. 1998; squirrel monkey: USrey & Reid, 2000). It
has been hypothesized that M and P cells support distinct
extrastriate visual pathways, based upon differences in
physiological signatures and upon their separate projection
pathways to V1 and within V1 (see DeYoe & Van Essen,
1988; Livingstone & Hubel, 1988; Merigan & Maunsell,
1993; Casagrande & Kaas, 1994 for review). For example,
the greater sensitivity of P cells to chromatic contrast (in
macaque monkeys and marmosets) and to higher spatial
frequencies (in all primate species examined) has been
linked to the processing of detail and colour while the
greater sensitivity of M cells to higher temporal frequencies
has been linked to motion perception (DeYoe & Van Essen,
1988; Livingstone & Hubel, 1988).

Few studies, however, have examined the response properties
of K LGN cells, especially in simian primates. K LGN cells
have been studied in the most detail in the nocturnal
prosimian bush baby (Galago crassicaudatus) (Norton &
Casagrande, 1982; Irvin et al. 1986, 1993; Norton et al.
1988). In contrast to M and P cells, K cells in bush babies
were found, on average, to have larger receptive fields and
slower orthodromic and antidromic conduction velocities.
Like cat W cells, bush baby K cells (referred to earlier as
W-like) also appeared to be heterogeneous as a group; some
could not be driven well by grating stimuli or were only
doorly driven by such stimuli. Other bush baby K cells,
however, were unlike the 'sluggish' cat W cells, and
responded briskly to gratings; they exhibited contrast
sensitivity functions whose resolution levels lay intermediate
between those of the average M or P cell (Norton et al. 1988).

Only recently has there been an effort to examine the
response properties of K cells in any simian primate (Martin
of K cells in the diurnal New World marmoset monkey
revealed one population of K cells that appeared to receive
input from blue-ON ganglion cells suggesting involvement
of this pathway in the processing of chromatic information
(Martin et al. 1997, White et al. 1998). Since K cells send
axons to the CO blobs of V1, the latter result is consistent
with the proposed role of CO blob cells in chromatic
processing (Livingstone & Hubel, 1988), although recent
studies in primate V1 conflict with earlier reports and
suggest that CO blob cells are not unique in their selectivity
for chromatic stimuli (Lennie et al. 1999; DeBruyn et al.
1993; Edwards et al. 1995; Leventhal et al. 1995). Because
bush babies have only a single cone type, lack blue cones
and also have well-defined CO blobs, K cells and their CO
blob targets probably perform more universal visual
function than the processing of chromatic signals or perform
more than one role across species (Wilder & Rakic, 1990;
Casagrande & Kaas, 1994; Jacobs et al. 1996). Regardless,
given the paucity of information available on K cell

Our aim for the present study was to characterize fully the
receptive field properties of K LGN cells in the nocturnal
simian owl monkey and compare K cell properties to those
of M and P cells. Owl monkeys, like prosimian bush babies,
have only one cone type and lack blue cones entirely, and
like marmosets, are New World simian primates (Wikler &
us several other advantages. First, with the exception of
macaque monkeys, the visual systems of owl monkeys have
been studied in the most detail (for review see Casagrande
& Kaas, 1994). Second, owl monkeys have well-developed
LGNs with simple laminar patterns consisting of two
Layers, two M layers and at least three well-defined
K layers. Finally, the axon structure and cortical target cells
of K LGN cells in owl monkeys have been studied in detail
(Ding & Casagrande, 1997). Some of the results reported here
were presented previously in abstract form (Xu et al. 1999).

METHODS

General preparation

Conventional extracellular recording techniques were employed to
examine the receptive field properties of LGN neurones in nine adult
owl monkeys (Aotus trivirgatus). These monkeys were a gift from
the USAID Malaria Vaccine Development Program. We recorded
directly from LGN cells in seven owl monkeys. In the remaining
two monkeys recordings were made from LGN cell afferent axons
in V1 where intrinsic neuronal activity was inhibited (silenced) by
the GABA agonist muscimol (Chapman et al. 1991, Boyd et al.
1998). All monkeys were handled according to the National
Institute of Health Guide for the Care and Use of Animals under
an approved protocol from the Vanderbilt University Animal Care
and Use Committee. Animals were pre-medicated with injectable
atropine (0.1 mg kg−1), acepromazine (0.5–1 mg kg−1) and
dexamethasone (2 mg kg−1). Anaesthesia was induced with an
intramuscular injection of ketamine HCl (8–12 mg kg−1) and mask
inhalation of Isoflurane. Animals were maintained with these
anaesthetics while a cannula was inserted into the femoral vein
of one hind-limb for subsequent delivery of anaesthetic and
neuromuscular blocking agents. Anaesthesia was then maintained
with injections of ketamine for the remainder of the surgical
manipulations. After a cannula was placed in the trachea, the
animals were mounted in a stereotactic apparatus, the scalp was
reflected on the midline and stainless-steel screws were inserted in
the skull over the frontal lobe for recording EEGs to monitor
general levels of arousal. Neuromuscular blockade was initiated by
i.v. injection of 1–1.5 mg kg−1 vecuronium bromide (Norcuron).
Animals were artificially ventilated with a mixture of 75% N2 O,
23.5% O2 and 1.5% CO2 delivered at a rate of 28–35 strokes
min−1 with a volume of about 15 ml to maintain the peak end tidal
CO2 level at 4%. Anaesthesia and neuromuscular blockade were
maintained by intravenous infusion of sufentanil citrate (Sufenta;
12–15 mg kg−1 h−1) and vecuronium bromide (0.2 mg kg−1 h−1)
mixed in 5% dextrose lactated Ringer solution delivered at a rate
of about 27 ml h−1. In order to ensure that adequate levels of
anaesthesia were maintained, heart rate, end-tidal CO2 and EEGs

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were monitored continuously in the presence of the neuromuscular blocker. If the animal exhibited any signs that the anesthetic levels were inadequate (fluctuating heart rate or low voltage fast EEG records), the percentage of Sufenta in the infusion line was immediately increased.

For LGN recording, a small craniotomy (4 mm × 5 mm) was made over the location of LGN according to stereotaxic coordinates established previously and the dura reflected. The brain was protected with a layer of agar. After the electrode was inserted into the brain, the opening in the skull was covered with an additional layer of paraffin wax to ensure recording stability. For recordings that were to be made from 'silenced' V1, holes were drilled directly over the area centralis representation in V1 in the posterior lateral cortex; the dura in the hole was incised. Initially, the receptive field properties of one or two cortical cells in V1 were analyzed. The electrode was then withdrawn and 25–50 µl of 50 mm muscimol was applied to the exposed cortex for 5–10 min to silence cortical activity. The cortical surface was then rinsed with saline before the craniotomy was sealed (Chapman et al., 1991; Boyd et al., 1996). Muscimol was reapplied to the cortical surface as soon as there were signs of recovery of neural activity. Generally cortical activity did not show any signs of recovery during the typical duration, less than 4 h, of one penetration in cortex. LGN axons could be distinguished from cortical cells based upon their generally higher spontaneous activity, monocular responses and vigorous responses to flashing spots and rapidly moving stimuli. At the end of each penetration electrolytic lesions were made to mark cells so that their locations could be reconstructed relative to laminar and compartmental (CO blob or interblob) borders.

Pupils were dilated with atropine eye drops (0.5% ophthalmic atropine sulfate). Individually fitted clear gas-permeable contact lenses were used to render the retina conjugate with the viewing screen 57 cm distant. In some animals, lenses with 3 mm pupils were used. Retinal landmarks (optic disc and area centralis) were projected onto the plotting screen with the aid of a reversible ophthalmoscope. The electrode was then lowered into the brain using a microdrive, and responses to visual stimuli were monitored until characteristic LGN responses were found. During physiological recording, EEG, ECG, end tidal CO₂ level and rectal temperature were monitored and maintained at appropriate levels.

**Recording, stimulation and data acquisition**

Commercially made Parylene-coated tungsten electrodes (FHC Inc., Bowdoinham, ME, USA) with an impedance of 5–10 MΩ were used to record from LGN cells. Well-isolated units were used to trigger standard pulses, which could be played over the audio monitor and counted by a computer that also controlled the presentation of stimuli.

The receptive field of each unit was initially plotted by manually controlled stimuli displayed on the tangent screen, with eye dominance determined and the receptive field boundaries drawn. The receptive field centres and surround were identified as either ON, OFF or ON–OFF. In addition, any strong suppressive effect of the surround on the centre was noted. The horizontal and vertical extent of the receptive field centre was measured on a plotting table using a small flashing spot. The average horizontal and vertical extent of the receptive field centre was taken as the diameter of the receptive field centre. We also qualitatively differentiated cells into sustained and transient categories according to their response to stationary contrast stimuli presented for 5–10 s within the centre of the receptive field. Units that exhibited above spontaneous maintained discharge during this period were categorized as sustained. Stimuli consisted of drifting sine-wave and counter-phase gratings presented at different spatial and temporal frequencies, contrasts and orientations, and phases in the case of counter-phase gratings. Stimuli were generated by an image-processing board (Pepper PRO 1280) with a capacity of 1024 pixels × 1280 pixels by 8 bits of modulation and presented on a cathode ray tube (CRT) screen that subtended an angle of 10 deg with a background luminance of 110 cd m⁻². Given that owl monkeys have a rod-dominated retina, we examined to see if our measurements using these luminance levels differed depending upon whether we used a 3 mm pupil or not. No differences were found. For cells with strong suppressive responses, stimuli were presented within a 4 deg window instead of the full 10 deg screen. Cells were tested with spatial frequencies ranging from 0.1 to 8.6 cycles deg⁻¹, temporal frequencies ranging from 1 to 32 Hz at the optimal spatial frequencies, and contrasts ranging from 3 to 56% at optimal spatial and temporal frequencies. Linearity was tested using different phase angles of the counter-phase sine-wave gratings stepped through the receptive field such that the cell's responses were sampled at all positions within the centre and surround at least once. The initial linearity test was run with spatial, temporal and orientation parameters optimized for the cell at moderate contrast (25%). Next the cell was re-tested at double its preferred spatial frequency. If the cell still responded adequately it was re-tested at 3 times the optimal spatial frequency, and so forth, until the cell no longer responded. By increasing the spatial frequency we could ensure that non-linearities in spatial summation would be reliably detected if present (Hochstein & Shapley, 1976; Derrington & stone, 1984).

Data were collected by a generic GC-386. The primary data analysis tool was construction of 2-s, 128 times post-stimulus time (PST) histograms. The interleaved histogram technique of Henry et al. (1973) with randomization was adopted to reduce artifacts from the inherent non-stationarity of the visual system. A stimulus set was specified and comprised each measuring condition as well as a null condition (a blank screen at the mean luminance of the gratings) to assess the maintained discharge. Each element in the stimulus set was presented once in a random order with a 1 s interval of blank screen between each presentation. Presentation of the set was then repeated in a random order until each stimulus condition had been tested completely (5–10 times). With 4 s presentation periods, data are based on 20–40 s of averaging for each condition. The PST histograms for each cell were Fourier transformed, and the spatial (F1) and second harmonic (F2) components of the responses were analyzed. Data were plotted with Igor 3.1 (WaveMetrics, Inc.) software. The following receptive field properties were measured for each cell: peak spatial frequency and cut-off, peak temporal frequency and cut-off, response to stimulus contrast and linearity of spatial summation. In addition, we determined orientation selectivity for each cell tested and all remaining tests were done with the gratings set to the preferred orientation if one existed. Data on the orientation selectivity of LGN cells in owl monkey will be submitted for publication separately. If the cell exhibited no orientation selectivity then all tests involved vertically oriented drifting gratings.

We examined linearity of spatial summation with counter-phase gratings in all the cells that responded well to drifting gratings. The ratio of the mean amplitude of the second harmonic (averaged across all spatial phases) to the amplitude of the fundamental at the best phase was used as an index of non-linearity (Hochstein & Shapley, 1976; Derrington & Lennie, 1984). An index value greater than 1.0 indicates a substantial non-linearity. In the present study, cells with 2nd harmonic (F2)/1st harmonic (F1) ratios of < 1.0 at all spatial frequencies tested were classified as linear. Cells with
F2/F1 ratio > 1.0 at any one of the spatial frequencies tested were classified as non-linear cells.

Spatial and temporal frequency peaks were defined as the frequencies at which cells exhibited the highest first harmonic response. Cut-offs were determined by extrapolating the high-frequency limb of the curves (i.e., log frequency) to control levels determined from responses to the blank control screen (baseline).

Several measures of contrast sensitivity were compared. Threshold contrast was determined by extrapolating to baseline (i.e., log contrast). Contrast gain was determined based upon the slope of the linear portion of the contrast–response curve (i.e., log contrast) where responses were well above threshold. In addition, we attempted to fit contrast–response curves with a hyperbolic function in the form of:

$$\text{Response}(c) = \frac{R_{\text{max}}}{C_a + (C_a - C)^n},$$

where $R_{\text{max}}$ is the maximum response rate, $C_a$ is the contrast required for half-maximal response, and the exponent $n$ is the rate of change (Albrecht & Hamilton, 1982). Such hyperbolic fits were useful only in those cells that exhibited response saturation. In this group we also compared $C_a$ values obtained from these curve fits.

**Statistical analysis**

Statistical comparisons of receptive field properties across the K, P and M groups were done by one-way ANOVA with post hoc mean difference tests (Tukey and LSD tests), provided that the data did not violate the prerequisite of variance homogeneity across groups. For groups with both unequal variances and unequal samples, we used instead the Kruskal–Wallis test, which is a non-parametric procedure commonly advocated as an alternative of the ANOVA (Zar, 1999). In the latter case, a Mann–Whitney U test was used for between group comparisons. Alpha levels of $P \leq 0.05$ were considered significant.

**Histological procedures**

The position of each recorded cell or axon was noted by the depth indicated on the microdrive. Electrolytic lesions (5 μA × 5 s) were made to mark the location of electrode tracks (see Fig. 1). At least two electrolytic lesions were made on each electrode penetration to aid in reconstruction of the track and to calculate tissue shrinkage. At the termination of each experiment, the animal was deeply anesthetized with an overdose of pentobarbital (sodium pentobarbital) and perfused transcardially with a saline rinse followed by fixation with 0.1% glutaraldehyde and 3% paraformaldehyde in 0.1 M phosphate buffer. The brain was removed and cryoprotected overnight in a solution containing 30% sacrose in 0.1 M phosphate buffer. It was then frozen and cut into 50 μm slices. To locate the LGN cells recorded, alternate sections of the LGN were stained for Nissl bodies, CO or immunostained for calbindin-128k. CO staining was performed using the method described by Boyd & Matsubara (1996).

For LGN afferent axons in V1, K axons can be distinguished anatomically since they terminate within cortical layers III and I, while M and P axons terminate within the upper and lower halves of cortical layer IV, respectively (see Ding & Casagrande, 1997). To identify cortical layers alternate V1 sections were stained for Nissl bodies and CO, respectively. LGN axon types were classified based upon the laminar position established via reconstructions of the electrode tracks.

Calbindin immunostaining was performed using a rabbit polyclonal antibody (Swant CB-38). Before being placed in the primary antibody, the sections were placed in 0.3% H2O2 for 30 min followed by three rinses in 0.1 M Tris-buffered saline (TBS, pH 7.4). Sections were then incubated for 1 h in blocking buffer consisting of 10% normal donkey serum, 1% bovine serum albumin and 0.1% Triton X-100 in TBS. Then the sections were placed in primary antibody diluted 1:5000 in antibody buffer consisting of 10% normal horse serum, 0.2% bovine serum albumin and 0.2% Triton X-100 in TBS. Sections were incubated overnight in the primary antibody and then rinsed 3 times in TBS before being placed in the secondary antibody. The secondary antibody (biotinylated donkey anti-rabbit (Chromogen) was diluted at 1:200 in antibody buffer. Sections were incubated in secondary antibody for 2 h, rinsed 3 times in TBS and placed in an avidin–biotin complex (Standard Elite ABC kit, Vector) prepared in TBS. After 2 h the sections were rinsed 3 times with TBS and the calbindin immunoreactivity was visualized by placing the sections into a solution containing 50 mM imidazole, 25 mM nickel ammonium sulfate, 0.01–0.02% 3,3-diaminobenzidine and 0.0003% H2O2 in 0.05 M TBS. Sections remained in this solution until labelled cells were visible and the reaction product was quite dark (usually 15–20 min).

**RESULTS**

We recorded from 133 LGN cells and 10 LGN afferent axons in silenced V1 with eccentricities ranging from 2.8 to 31.3°. All LGN axons recorded from silenced V1 had eccentricities of less than 10°. From this total population 38 K units (36 cells and 2 axons), 45 P cells and 34 M units (29 cells and 5 axons) could be assigned with confidence to specific layers based upon histological reconstructions.

**Receptive field structure and size**

Twenty (53%) of the 38 K units analyzed gave a sustained response to a stationary stimulus held in the receptive field center for at least 5 s. The remaining 18 (47%) responded transiently. Twenty-three K units exhibited typical center–surround receptive fields either ON center with an OFF surround or the reverse. Fifteen of the K units had either strong suppressive surrounds, ON–OFF surrounds or no clear surrounds. Thirteen K units (34%) were poorly driven by grating stimuli, but responded well to flashing spots or single light bar stimuli moved manually. Of the 13 K cells that did not respond to gratings, 2 appeared to respond only to changes in luminance and, although this impression was not tested quantitatively, these cells seemed to respond very much like the ‘luminance units’ described in the cat retina by Barlow & Levick (1969). In addition, two other K cells seemed to have unusually long onset latencies to flashing spots of light.

Thirteen (38%) of the 34 M units analyzed responded in a sustained manner and the remainder responded transiently to a stationary stimulus of appropriate contrast. In contrast to the K and M populations, the majority (71%) of P cells responded in a sustained fashion. Ninety per cent of the P and M units showed standard center–surround receptive fields; the remainder showed either weak or unclear surrounds or suppressive surrounds. Also, in contrast to the
K population, only four P cells (9%) and two M cells (6%) responded poorly to grating stimuli although all of these cells responded briskly to flashing spots or moving bars of light.

Since one property reported to characterize cat W cells is that of sluggishness (see Sur & Sherman, 1982) we compared the peak response rates with optimal stimulus conditions for those K, P and M cells that responded to gratings. As reported by Usrey & Reid (2000), owl monkey LGN cells tend to show low overall peak responses. No statistical differences ($P = 0.27$) were found between K, M and P cells in peak responses to spatial and temporal frequencies at moderate contrast (28%): K cells, $10.6 \pm 1.0$ spikes s$^{-1}$ (mean $\pm$ s.e.m.; $N = 24$), M cells, $12.4 \pm 1.0$ spikes s$^{-1}$ ($N = 32$), and P cells $10.2 \pm 1.0$ spikes s$^{-1}$ ($N = 33$).

In all three LGN cell classes receptive field centre size tended to increase with eccentricity, but this relationship was least clear for the K population where there was a large

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**Figure 1. Histological reconstruction of recording sites**

Photomicrographs of adjacent parasagittal LGN sections showing lesions in sections stained for Nissl bodies (A), CO (B) and calbindin-D28k (C and D). Calbindin-D28k labels cells mainly in layers K1, K2 and K3. Calbindin-labelled cells in K4 are sparse. The arrow and arrowhead in B indicate a pair of lesions located in the contralateral M layer (arrowhead) and in layer K2 (arrow) that mark a single electrode penetration. The arrows in $A$ and $B$ point to the same lesion; arrowheads in $B$ and $C$ point to the same lesion. $D$ shows a higher power photomicrograph of the distribution of K cells immunostained for calbindin-D28k in K layers K1–K3 in another case. Scale bar in $B$, 200 $\mu$m and also applies to $A$ and $C$, in $D$, 100 $\mu$m.
degree of scatter at all eccentricities. Figure 2 shows the relationship between receptive field centre size and eccentricity for the subset of the K (14), P (19) and M (18) cells where centre boundaries were unambiguous. For this population, average receptive field centre diameter was 1.05 ± 0.25 deg for K cells, 0.87 ± 0.11 deg for P cells and 0.92 ± 0.13 deg for M cells.

**Linearity of spatial summation**

In cats the major feature that is used to distinguish X and Y retinal ganglion and LGN cells is linearity or non-linearity of spatial summation (Enroth-Cugell & Robson, 1966; Hochstein & Shapley, 1976). According to Hochstein & Shapley (1976) the identification of a cell as an X cell on the basis of linear summation requires not only a strong dependence on spatial phase and response at the fundamental modulation frequency, but also that spatial phase dependence be demonstrated at higher than the cell’s preferred spatial frequency. This is because Y cells can exhibit a strong spatial phase dependence and respond quite well at the fundamental modulation frequency if the grating is presented at a low spatial frequency. Therefore, we also examined for spatial phase dependence both at the preferred spatial frequency of the cell and at least 2 x the preferred frequency. Cells were considered to respond linearly if they showed a clear null F1 response. Also, if the F2 response became dominant over the F1 response we classified the cell as non-linear (Hochstein & Shapley, 1976). In Y cells the F2 component was found to be phase insensitive. We calculated the F2/F1 response ratio as an index of non-linearity: In cats, X cells were always found to have a non-linearity index of less than 1·0 while for Y cells this index was found to be greater than 1·0 (Hochstein & Shapley, 1976).

We found that all K (N = 17), P (N = 32) and all but one M (N = 27) unit could be classified as linear according to the above described criteria. Figure 3 shows examples of the spatial phase dependence in the responses of different LGN cells and provides a summary histogram showing the non-linearity indices of all cells.

The K cell shown in Fig. 3A was tested at its optimal spatial frequency of 0·8 cycles deg\(^{-1}\). The peaks of the F1 responses are much higher than average F2 responses at all phases outside of the null points. This cell exhibited a non-linearity index of 0·21. Nulls are clearly evident at phase positions −120 and 60 deg. Figure 3B shows the responses of the same cell tested at 2 x the optimal spatial frequency. As would be expected the F1 responses decrease as the spatial frequency is increased beyond the cell’s preferred frequency but at no point is the average F2 response higher than the F1 response. Even at 2 x the preferred spatial frequency, this cell still exhibits clear evidence of a null. All P cells and all but one M cell showed clear evidence of spatial linearity. The spatial phase responses of a typical M cell are shown in Fig. 3C. In this example the cell was tested at 2 x its preferred spatial frequency. The F1 response curve exhibits nulls at −90 and 90 deg. At the peak responses of this cell the F1 curve is always higher than the F2 curve. Only one M cell showed any indication of non-linearity (Fig. 3D). In this cell the peak F2 responses are higher than the F1 response. However, unlike cat Y cells this cell shows evidence of phase dependence.

Figure 3E shows a summary distribution of the non-linearity indices for all cells in the population tested at their optimal or close to optimal spatial frequencies. Only one M cell (also shown in Fig. 3D) exhibited an index of greater than 1·0 suggesting spatial non-linearity. The distributions of the non-linearity indices for the other cells show no clear trends that correlate with cell class. The majority of K, M and P cells show non-linearity indices of < 0·4 with a peak for each of the populations around 0·2.

**Spatial and temporal resolution**

Spatial and temporal frequency tuning curves and responses to contrast are shown for representative K, P and M cells in...
Fig. 4A. A peak spatial frequency at about 0.8 cycles deg\(^{-1}\) and a high spatial frequency cut-off at around 6 cycles deg\(^{-1}\). Typical of all K cells tested, this K cell showed a broad band-pass tuning curve with a sharper drop-off in response to higher than lower frequencies. As shown in Fig. 4B, this K cell responded to temporal frequencies over a broad range from 1.0 Hz (the lowest temporal frequency tested) to 6.0 Hz with a clear peak at 2 Hz. The contrast–response function for this same K cell is shown in Fig. 4C. The extrapolated contrast threshold for this cell is about 2.5%, and its contrast gain (the slope for the linear segment of the rising phase) is 29 spikes s\(^{-1}\) per log contrast %.

Spatial and temporal tuning curves for a representative P cell are shown in Fig. 5. Unlike K and M cells most P cells exhibited symmetrical band-pass spatial frequency tuning curves. This cell showed a peak spatial frequency response at 0.8 cycles deg\(^{-1}\) and a cut-off around 6.4 cycles deg\(^{-1}\) (Fig. 5A). Like the K cell shown in Fig. 4, this P cell had a peak temporal frequency at 2.0 Hz but a higher cut-off at 10 Hz. The contrast–response curve for this P cell is shown

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**Figure 3. Linearity of spatial summation**

The first harmonic component (F1) is shown by ・ and the second harmonic component (F2) by □. A, an example of a K cell tested with a phase angle range (from −180 to 180 deg) in 12 steps at its optimal spatial frequency (SF 0.8 cycles deg\(^{-1}\)), optimal temporal frequency (TF 2 Hz) and moderate contrast (28%). The F1 curve had null positions around −120 and 60 deg. B, the same K cell tested at 2 × the optimal SF (SF 1.6 cycles deg\(^{-1}\)). Its F1 responses decreased, but the peaks of the F1 curve were still higher than the average F2 curve across all phase angles. C, phase tuning curve for an M cell at 2 × its optimal spatial frequency (SF 0.8 cycles deg\(^{-1}\)), optimal temporal frequency (TF 2 Hz) and moderate contrast (28%). The peaks of the F1 curve were higher than the average F2 curve at all phase angles. The F1 curve had null positions around −90 and 90 deg. D, only one M cell exhibited any evidence of spatial non-linearity. This cell was tested with a global phase angle range (from −180 to 180 deg) in 12 steps at its optimal spatial frequency (0.6 cycles deg\(^{-1}\)), optimal temporal frequency (2 Hz) and moderate contrast (28%). Average F2 responses across all phase angles except at the centre of the field are higher than F1 responses with a ratio of F2/F1 greater than 1.0 at the peaks. E, non-linearity index (F2/F1) histograms of K, P and M cells. Most K, P and M cells have a non-linearity index of less than 0.4. Out of 76 cells (K = 17, P = 32, M = 27), one M cell had an index greater than 1.0 suggestive of spatial non-linearity.
in Fig. 5. It shows a sigmoidal shape and an extrapolated threshold of around 5% contrast and a contrast gain of 29.2 spikes s⁻¹ per log contrast %. Most P cells had contrast gains of less than 15 spikes s⁻¹ per log contrast %, which was lower than that found in most M and K cells.

Spatial and temporal tuning curves for a representative M cell are shown in Fig. 6. Unlike K and P cells, some M cells did not exhibit a low spatial frequency roll-off as shown in Fig. 6A. This cell had a peak spatial frequency of 0.2 cycles deg⁻¹ and spatial frequency cut-off of 6 cycles deg⁻¹. This cell also responded well to all temporal frequencies from 1.0 Hz (the lowest tested) to 8 Hz but still responded above background at 20 Hz. This cell's peak temporal frequency is the same as that for the K and P cells shown earlier, 2 Hz (Fig. 6B). The contrast–response curve for this M cell is shown in Fig. 6C. It shows an extrapolated threshold of 2.5% contrast and a contrast gain of 18.9 spikes s⁻¹ per log contrast %.

Figures 4 and 5. K and P cell spatial frequency, temporal frequency and contrast tuning curves

The F1 response curves for one K cell are shown for different spatial frequencies (in cycles deg⁻¹) in A and different temporal frequencies (in Hz) in B. This cell had a SF peak at 0.8 cycles deg⁻¹ and a cut-off around 6 cycles deg⁻¹. The peak TF was 2 Hz and the cut-off was around 6 Hz. The contrast response curve is shown in C.

The F1 response curves for one P cell are shown for different spatial frequencies (in cycles deg⁻¹) in A and different temporal frequencies (in Hz) in B. This cell had a SF peak at 0.8 cycles deg⁻¹ and a cut-off above 6.4 cycles deg⁻¹. The peak TF was 2 Hz and the cut-off was 10 Hz. The contrast response curve is shown in C.
Table 1 provides the mean ± s.e.m. spatial and temporal resolution values for each population of cells which included a total of 25 K, 41 P and 32 M units. Statistical comparisons were confined only to the 15 K, 27 P and 16 M cells (Table 2) at roughly matched eccentricities (>10 deg). The histograms in Figs 7–9 compare the eccentricity-matched populations for each parameter measured. As can be seen in Table 2 and Fig. 7 K, P and M cells show broad overlapping ranges of peak and cut-off spatial frequencies. A one-way ANOVA comparison of the populations at matched eccentricities revealed significant differences in peak spatial frequencies and cut-offs (P = 0.008 and 0.05, respectively). A post hoc mean test showed that the P cells had significantly higher peak spatial frequencies than K or M cells (P = 0.02 and 0.03, respectively); the mean peak spatial frequencies for K and M cells did not differ (P = 0.99). P cells differed significantly from M cells in their spatial frequency cut-offs (P = 0.05). Although K cells tended to have lower mean spatial cut-offs than P cells, this trend did not reach significance (P vs. K, P = 0.26; M vs. K, P = 0.73).

Table 1. Summary of spatial and temporal resolution and contrast sensitivity values for the K, P and M populations

<table>
<thead>
<tr>
<th></th>
<th>Peak SF</th>
<th>SF cut-off</th>
<th>Peak TF</th>
<th>TF cut-off</th>
<th>Contrast threshold</th>
<th>Contrast gain</th>
</tr>
</thead>
<tbody>
<tr>
<td>K cell (n = 25)</td>
<td>0.56 ± 0.07</td>
<td>3.0 ± 0.3</td>
<td>2.19 ± 0.17</td>
<td>11.3 ± 0.9</td>
<td>4.5 ± 0.6</td>
<td>17.5 ± 2.0</td>
</tr>
<tr>
<td>P cell (n = 41)</td>
<td>0.72 ± 0.06</td>
<td>4.5 ± 0.6</td>
<td>2.13 ± 0.12</td>
<td>9.6 ± 0.7</td>
<td>4.7 ± 0.6</td>
<td>15.0 ± 1.1</td>
</tr>
<tr>
<td>M cell (n = 32)</td>
<td>0.59 ± 0.06</td>
<td>3.3 ± 0.3</td>
<td>2.00 ± 0.13</td>
<td>12.5 ± 1.2</td>
<td>3.2 ± 0.2</td>
<td>18.8 ± 1.4</td>
</tr>
</tbody>
</table>

All values are means ± s.e.m.

Table 2. Summary of spatial and temporal resolution and contrast sensitivity values for the K, P and M cells at matched eccentricity (>10 deg)

<table>
<thead>
<tr>
<th></th>
<th>Peak SF</th>
<th>SF cut-off</th>
<th>Peak TF</th>
<th>TF cut-off</th>
<th>Contrast threshold</th>
<th>Contrast gain</th>
</tr>
</thead>
<tbody>
<tr>
<td>K cell (n = 15)</td>
<td>0.43 ± 0.07</td>
<td>2.7 ± 0.4</td>
<td>2.25 ± 0.25</td>
<td>11.2 ± 1.2</td>
<td>3.8 ± 0.6</td>
<td>17.6 ± 2.8</td>
</tr>
<tr>
<td>P cell (n = 27)</td>
<td>0.69 ± 0.06</td>
<td>4.0 ± 0.4</td>
<td>1.96 ± 0.11</td>
<td>9.1 ± 0.8</td>
<td>5.0 ± 0.8</td>
<td>14.5 ± 1.3</td>
</tr>
<tr>
<td>M cell (n = 16)</td>
<td>0.43 ± 0.06</td>
<td>3.1 ± 0.51</td>
<td>2.06 ± 0.17</td>
<td>14.9 ± 2.0</td>
<td>2.9 ± 0.3</td>
<td>18.2 ± 1.4</td>
</tr>
</tbody>
</table>

Figure 6. M cell spatial frequency, temporal frequency and contrast tuning curves

The F1 response curves for one M cell are shown for different spatial frequencies (in cycles deg⁻¹) in A and different temporal frequencies (in Hz) in B. This cell had a SF peak at 0.2 cycles deg⁻¹ and a cut-off around 6 cycles deg⁻¹. The peak TF was 2 Hz and the cut-off was above 20 Hz. The contrast response curve is shown in C.
Interestingly, as shown in Fig. 7A, the K cells' peak spatial frequency distribution appeared to have two modes suggesting that K cells have subclasses.

Figure 8 compares the peak and cut-off temporal frequencies for K, P and M cells. Mean values are given in Table 2. As can be seen in Fig. 8A, K, P and M cells have very similar temporal frequency peak distributions with most cells preferring 2-0 Hz. Figure 8B shows that K cells have temporal frequency cut-offs that lie between those of P and M cells although here again the populations show broad areas of overlap. Temporal frequency cut-offs were found to differ significantly between the three groups (Kruskal-Wallis test, $P = 0.003$). This difference can be accounted for by differences between P and M cells ($P = 0.001$) since Mann-Whitney $U$ tests showed that the K cell population did not differ from either the M ($P = 0.22$) or P ($P = 0.07$) populations. As can be seen, the temporal resolution values of K cells lie between those of P and M cells.

It is also noteworthy that, as demonstrated for K cells in the peak spatial frequency domain, M cells exhibit double peaks indicative of two modes in both spatial and temporal frequency cut-offs (Figs 7B and 8B). The double peaks seen in the M distributions also may hint at the existence of subclasses.

Figure 9 compares contrast threshold values (A) and the contrast gain values (B) for each of the populations; mean values are shown in Table 2. Contrast thresholds of M cells were significantly lower than those of P cells (Mann-Whitney $U$ test, $P = 0.05$). K cells did not differ significantly from M or P cells ($P = 0.26$ and $0.45$). M cells exhibited significantly higher average contrast gains than P cells (Mann-Whitney $U$ test, $P = 0.03$).
test, \( P = 0.05 \), although K cells did not differ significantly from either P or M cells (\( P = 0.48 \) and 0.53). Thus, as with temporal resolution, the contrast characteristics of K cells seem to lie between those of M and P cells.

We also attempted to fit contrast–response curves for all the cells tested with a hyperbolic function in the form of:

\[
\text{Response (C)} = \frac{R_{\text{max}} \cdot (C^n)}{(C^n + C_{50}^n)},
\]

where \( R_{\text{max}} \) is the maximum response rate, \( C_{50} \) is the contrast required for half-maximal response and the exponent \( n \) is the rate of change or contrast gain index (Albrecht & Hamilton, 1982). The contrast–response curves of the majority of M (24/32) and K cells (15/23) were well fitted by a hyperbolic function; however, the curves of most P cells (25/37) showed little response saturation and thus could not be adequately fitted by this function. For those K, P and M cells where the fit was good, the mean \( C_{50} \) differed between populations. For K cells it was 25.9 ± 4.0, for P cells 39.0 ± 5.3 and for M cells 20.7 ± 2.6. The mean \( C_{50} \) differed significantly between the three groups (ANOVA, \( P = 0.005 \)). The mean \( C_{50} \) of P cells was significantly different from that of either M or K cells (P vs M, \( P = 0.001 \); P vs K, \( P = 0.03 \)). However, the mean \( C_{50} \) of K cells did not differ significantly from that of M cells (\( P = 0.29 \)).

**Properties of K cells in different layers**

There are four K layers in the owl monkey based upon the distribution of calbindin-labelled cells. Of these K layers, three are well developed, with K1 and K3 exhibiting the largest number of K cells (see Fig. 1C and D). Of our sample of K units, reconstructions indicated that 2 K axons in V1 were in layer 3B suggesting that they probably arise from LGN layer K3; 4 were in LGN layer K1, 11 were in LGN layer K2, 20 were in LGN layer K3 and 1 was in LGN layer K4. However, of the total sample of 38 K units, only 3 K1, 6 K2, 14 K3 cells plus the 2 K axons responded well enough to grating stimuli for quantitative measures to be made.

Since our previous anatomical studies in owl monkeys showed that the different K cell layers (K1/K2 vs. K3) have distinct axonal termination patterns in V1 (Ding & Casagrande, 1997), we asked whether these anatomical distinctions correlated with any differences in K cell receptive field properties. As shown in Fig. 10 differences were found between the properties of the cells in the different K layers. Cells in K1/K2 tended to be selective for lower spatial frequencies and higher temporal frequencies than cells in layer K3 (Fig. 10A and B). These trends were significant, however, only for the differences between temporal frequency cut-offs (Mann-Whitney \( U \) test, \( P = 0.05 \)) perhaps due to the small \( N \) value. In addition, K1/K2 cells consistently exhibited higher contrast thresholds and lower contrast gains than K3 cells (Fig. 10C and D), although this trend did not reach statistical significance. Overall, these K layer differences suggest that the ventral-most K layers, K1/K2, resemble M cells more than P cells in their spatial and temporal resolution characteristics whereas the resolution values of K3 cells tend to lie intermediate between those of P and M cells. Clearly, a larger \( N \) value will be required to confirm the trends seen.

**DISCUSSION**

By analogy to previous work on M and P cells, our goal was to construct a physiological profile of K cells that might be linked to particular visual perceptual attributes. We find that the spatial and temporal resolution of most K cells in the owl monkey overlap extensively those of M and P cells and that, unlike M and P cells, a large minority of K cells could not be driven by standard drifting gratings. The response of the K cell population to visual stimuli is therefore more heterogeneous than the response of either the M or P cell population. Finally, our data provide evidence that some of the heterogeneity within the K population can be accounted for by position in the LGN because cells in different K layers exhibit differences in spatial and temporal resolution.
resolution. We will consider each of these findings in light of studies conducted by others.

**Non-standard receptive field properties**

A third of the K cells in owl monkeys responded poorly or not at all to drifting gratings. Less than 8% of P and M cells displayed similar response behaviour. Compared with the M and P cells, a larger proportion of K cells also were difficult to characterize with other stimuli including manually moved light bars or flashing spots of various sizes. This description of K cell responses is consistent with earlier findings in the prosimian bush baby (Norton & Casagrande, 1982; Irvin et al. 1986) and the New World simian marmoset (Solomon et al. 1999; A. J. White, S. G. Solomon & P. R. Martin, unpublished observations). The existence in owl monkeys, bush babies and marmosets of many K cells that cannot be driven by standard stimuli supports the idea that K cells are quite distinct from M and P cells. Some of the non-standard characteristics that have been reported for both simian and prosimian K cells have also been reported for cat W cells. These properties include very large and difficult-to-plot receptive fields, and non-standard centre-surround organization. Some K cells may therefore be analogous to cat W cells (Sur & Sherman, 1982; see also Norton & Casagrande, 1982).

**Receptive field structure and size**

Consistent with an earlier study of LGN receptive field properties in owl monkeys (Sherman et al. 1976), we found that M cells were, on average, more transient in their responses to appropriate standing contrast stimuli than were P cells. Approximately half the K cells we tested responded in a transient manner. We did not find a correlation between the degree of response transience and other properties of K cells, but this possibility was not examined in detail. Sustained and transient responses have been previously examined for all three cell classes only in the bush baby by Norton & Casagrande (1982) who reported that most M cells exhibited transient responses, most P cells exhibited sustained responses, and about equal numbers of K cell were sustained or transient.

Receptive field centre sizes in owl monkey tend to increase with increasing eccentricity as in the LGNs of other primates (present study; for review see Casagrande & Norton, 1991). K cells differed from this pattern only in the

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**Figure 10. Spatial and temporal characteristics and contrast sensitivity of K cells in different layers**

*Panel A:* the mean peak SF was 0.42 ± 0.12 cycles deg⁻¹ for K1/K2 cells (n = 9) and 0.64 ± 0.10 cycles deg⁻¹ for K3 cells (n = 14). The mean SF cut-off was 2.5 ± 0.5 cycles deg⁻¹ for K1/K2 cells and 3.6 ± 0.5 cycles deg⁻¹ for K3. Peak SF and SF cut-off did not differ significantly between K1/K2 and K3 cells. *Panel B:* the average peak TF was 2.44 ± 0.29 Hz for K1/K2 cells and 1.96 ± 0.23 Hz for K3 cells. The average TF cut-off was 12.8 ± 1.7 Hz for K1/K2 cells and 9.5 ± 0.9 Hz for K3 cells. Peak SF did not differ significantly between K1/K2 cells and K3 cells, but K1/K2 cells had a significantly higher TF cut-off than K3 cells (Mann-Whitney U test; P < 0.05). *Panel C:* the mean contrast threshold was 4.7 ± 1.1 for K1/K2 cells and 4.3 ± 0.6 for K3 cells; values not significantly different. *Panel D:* the mean contrast gain was 14.2 ± 1.5 for K1/K2 cells and 20.9 ± 3.6 for K3 cells. Contrast gain did not differ significantly between K1/K2 cells and K3 cells.
overall degree of variation in centre size. Some K cells had significantly larger receptive field centre sizes than P or M cells at the same eccentricities, but other K cells had receptive field centres that fell in the range of the smallest we encountered. K cells in bush babies and marmosets have relatively larger receptive field centres than those of P and M cells at the same eccentricities and the variability in K cell size is also higher than that of P and M cells (Norton & Casagrande, 1982; Irvin et al., 1993; A. J. White, S. G. Solomon & P. R. Martin, unpublished observations). In marmosets, however, M cells also show high variability in size (A. J. White, S. G. Solomon & P. R. Martin, unpublished observations). Receptive field centre sizes for P cells are smaller than those of M cells at any eccentricity (Sherman et al. 1976; Norton & Casagrande, 1982; Derrington & Lennie, 1984; Irvin et al. 1993; O'Keefe et al. 1998; Urey & Reid, 2000). In owl monkeys two reports (O'Keefe et al. 1998; Urey & Reid, 2000) have shown that receptive field sizes of P cells are smaller than those of M cells at all eccentricities. P and M receptive fields in owl monkeys were also found to be larger than those of macaque monkeys by a factor of about 2 and smaller than those of the bush baby by a factor of 2 which fits with the differences in visual acuity of these three primate species (O'Keefe et al. 1998). Although our data are generally consistent with these results, we find more overlap between the P and M receptive field sizes than was reported previously in owl monkeys. Notably, although P cells in marmosac monkeys have been found to have smaller centres than M cells at matched eccentricities, significant overlap also has been reported (Derrington & Lennie, 1984; Spear et al. 1994; see Merigan & Maunsell, 1993 for review).

Spatial summation

Linearity of spatial summation is useful in distinguishing cell classes in cat LGN, but its usefulness in primate LGN cell classification remains controversial (see Casagrande & Norton, 1991 for review). We found that all K, P and M cells, with the exception of one M cell, were linear. Because we examined spatial linearity at the highest spatial frequency to which each cell would respond, we believe it is unlikely that we failed to drive non-linear subunits adequately and so missed a population of non-linear LGN cells. Consistent with our results, Urey & Reid (2000) report that all the M and P cells in owl monkey and squirrel monkey LGN (K cells were not examined) are linear as measured by a null test. Additionally, counterphase gratings revealed very few spatially non-linear LGN cells in bush babies that resemble Y cells in cats (Norton & Casagrande, 1982). In macaque monkeys and marmosets, however, variable percentages of non-linear LGN cells have been reported (Kaplan & Shapley, 1982; Marrocco et al. 1982; Derrington & Lennie, 1984; Blake more & Vital-Durand, 1986; Norton et al. 1988; O'Keefe et al. 1998; Urey & Reid, 2000; A. J. White, S. G. Solomon & P. R. Martin, unpublished observations). O'Keefe et al. (1998) report that P cells in owl monkey LGN have higher spatial frequency cut-offs, lower optimal temporal frequencies and cut-offs and lower levels of responsivity than M cells. Our results are consistent with these findings, except that our data suggest that the P and M cells preferred similar peak temporal frequencies. Both our work and that of O'Keefe et al. (1998) found that the differences between owl monkey M and P cells in contrast sensitivity and gain are markedly lower than those reported for macaque M and P cells (Kaplan & Shapley, 1982; Hicks et al. 1993; Derrington & Lennie, 1984; Spear et al. 1994). Physiological subclasses of K cells

In owl monkeys the different K layers (K1/K2 vs. K3) have distinct axonal termination patterns in V1. Axons from K3 cells mainly terminate within the cytochrome oxidase (CO) blobs in layer III of V1 and axons from cells in K1/K2 mainly terminate in cortical layer I (Ding & Casagrande, 1997). These anatomical differences suggest that some physiological differences may exist between these populations. Indeed, our results show that K cells in K1/K2 are more selective for lower spatial frequencies and higher temporal frequencies than cells in layer K3. Thus the K cells (K1/K2) that lie between or below the M layers tend to resemble M cells in resolution whereas the K3 cells have resolution values that lie between the average values for M and P cells (at least in terms of temporal resolution).
matching their anatomical position between the LGN M and P layers.

The distribution of the peak spatial frequencies of K cells also exhibits two modes, thus suggesting subclasses. Interestingly, M cells also showed double peaks within their distributions of spatial and temporal frequency cut-offs indicating the existence of M subclasses. The presence of M subclasses is consistent with anatomical work in owl monkeys (Boyd et al. 2000) and anatomical and modelling data in macaque monkeys (Lund et al. 1995; Bauer et al. 1999).

Additional support for the existence of K subclasses comes from the preliminary immunocytochemical work (Song et al. 2000) which shows that K cells in macaque monkeys are neurochemically diverse. Two marker proteins used to identify K cells in macaque monkeys, calbindin-D28k and the alpha form of calcium–calmodulin-dependent kinase II (αCaMKII), do not completely overlap in their distributions within the different LGN K layers. In fact, three subclasses of K cells could be identified by double-label immunocytochemistry. One subclass contained calbindin-D28k only, the second subclass contained αCaMKII only, and the third contained both proteins. The proportion of each type was found to vary within the different K layers suggesting that even within K layers different subclasses of K cells may coexist.

A role for K cells in vision

Our physiological results from owl monkeys cannot directly address the most interesting question; namely, what role do K cells play in the vision of primates? These data, however, do provide descriptions of K cell response characteristics from which we can draw a few conclusions. The first is that K cells in owl monkeys are a heterogeneous population. The idea that K cells contain subclasses is supported both by work in the nocturnal primate bush baby (Irvin et al. 1986) and work in the diurnal simian marmoset (Martin et al. 1997; White et al. 1998; Solomon et al. 1999; A. J. White, S. G. Solomon & P. R. Martin, unpublished observations). K cells in marmosets are heterogeneous with some responding to colour (blue-ON), others responding only to achromatic gratings and still others being unresponsive to gratings.

Our physiological results in owl monkey and comparable data in bush babies suggest that many K cells exhibit spatial and temporal resolution values in the range that would allow these cells to contribute to conventional aspects of spatial and temporal vision. If this is true, why is the anatomy of the K pathway so different from that of the P and M pathways? Why do K cells send their axons either to cortical layer I or to the CO blobs within cortical layer III and not to a subdivision of layer IV as is the case with M and P LGN cells? Casagrande (1994) suggested that clues about the K pathway might be gained by examining the role of the target cells (CO blobs) of the K pathway as well as any extraretinal inputs that are unique to K cells. What comes to mind first about the CO blobs is colour vision, based upon the famous papers by Livingstone & Hubel (1984, 1988). They suggested that cells in the CO blobs are tuned specifically to chromatic stimuli. The finding of blue-ON K cells in marmosets fits with this hypothesis, but is inconsistent with the fact that CO blobs are ubiquitously well developed in all primates and that K cell projects to CO blobs in all primates examined, but blue cones are absent in the nocturnal owl monkey and in the bush baby (Wikler & Rakic, 1990; Jacobs et al. 1993). Clearly K cells, CO blobs, and the anatomy of this pathway are highly conserved features across primates. Perhaps some K cells contribute uniquely to brightness contrast information but also colour contrast in species that have colour vision. Alternatively, K cells could contribute to a variety of other aspects of vision that might be tested adequately only in the awake behaving preparation. The subgroup of K cells that do not respond well to gratings could, for example, contribute to eye movement-related signals because K LGN cells in all primates receive a direct input from the superficial layers of the superior colliculus (input that P and M cells lack), and because a significant number of K cells project indirectly to the dorsal medial visual area (DM), an area concerned more with motion than with object vision (Harting et al. 1980, 1991; Beck & Kaas, 1998, 1999).

Finally, the neurochemistry and projection patterns of some K cells indicate that they are part of a neuro-modulatory pathway. LGN K cells and cells within the adjacent inferior pulvinar nucleus are similar in that they both contain calbindin (Jones & Hendry, 1989). Both K cells and the inferior pulvinar cells project to layer I, the most superficial layer of VI (Ogren & Hendrickson, 1977; Carey et al. 1979). Projections to layer I in all cortical regions are in a position to modulate signals within all cortical layers because the apical dendrites of the majority of cortical neurons extend into layer I (Vogt, 1991). Moreover, some investigators have proposed that the pulvinar nucleus is involved in visual attention (see Robinson & Petersen, 1992 for review). Perhaps some of the K cells that respond poorly in the anaesthetized animal project to layer I and are active along with the pulvinar cells in regulating visual attention. Future work in which K cells can be queried with an electrode in awake behaving monkeys may be able to more directly address these hypotheses.


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