

24. L. Vician et al., *Proc. Natl. Acad. Sci. U.S.A.* **92**, 2164 (1995).
25. M. Yoshihara et al., data not shown.
26. W. Li, J. T. Ohlmeyer, M. E. Lane, D. Kalderon, *Cell* **80**, 553 (1995).
27. Y. Zhong, V. Budnik, C.-F. Wu, *J. Neurosci.* **12**, 644 (1992).
28. G. W. Davis, C. M. Schuster, C. S. Goodman, *Neuron* **19**, 561 (1997).
29. D. O. Hebb, *The Organization of Behavior* (Wiley, New York, 1949).
30. U. Frey, R. G. Morris, *Nature* **385**, 533 (1997).
31. K. C. Martin et al., *Cell* **91**, 927 (1997).
32. We are grateful to S. Waddell for fruitful discussions, providing *Drosophila* strains, and critical reading of the manuscript; M. Sheng, M. Bear, S. Tonegawa, C. Quinn, M. Sur, and E. Montana for critical reading of the manuscript; K. Ikeda for useful discussion; C.-F. Wu for technical advice; G. Miesenböck for providing pHluorin DNA; N. Harden and E. Buchner for antibodies; and A. DiAntonio, D. Kalderon, J. Kiger, S. Thor, K. Schulze, J. Phillips, G. Davis, C. S. Goodman, and the Bloomington Stock Center for providing *Drosophila* strains. This work was supported by grants from the NIH, the Human

Frontiers Science Program Organization, the Packard Foundation, and the Searle Scholars Program.

Supporting Online Material

www.sciencemag.org/cgi/content/full/310/5749/858/DC1

Materials and Methods

Figs. S1 and S2

References and Notes

18 July 2005; accepted 5 October 2005
10.1126/science.1117541

Fast Readout of Object Identity from Macaque Inferior Temporal Cortex

Chou P. Hung,^{1,2,4*} Gabriel Kreiman,^{1,2,3,4*} Tomaso Poggio,^{1,2,3,4} James J. DiCarlo^{1,2,4}

Understanding the brain computations leading to object recognition requires quantitative characterization of the information represented in inferior temporal (IT) cortex. We used a biologically plausible, classifier-based readout technique to investigate the neural coding of selectivity and invariance at the IT population level. The activity of small neuronal populations (~100 randomly selected cells) over very short time intervals (as small as 12.5 milliseconds) contained unexpectedly accurate and robust information about both object "identity" and "category." This information generalized over a range of object positions and scales, even for novel objects. Coarse information about position and scale could also be read out from the same population.

Primates can recognize and categorize objects as quickly as 200 ms after stimulus onset (1). This remarkable ability underscores the high speed and efficiency of the object recognition computations by the ventral visual pathway (2–5). Because the feed-forward part of this circuitry requires at least eight or more synapses from the retina to anterior IT cortex, it has been proposed that the computations at each stage are based on just one or very few spikes per neuron (6, 7). At the end of the ventral stream, single cells in IT cortex show selectivity for complex objects with some tolerance to changes in object scale and position (2–4, 6, 8–16). Small groups of neurons in IT cortex tuned to different objects and object parts might thus provide sufficient information for several visual recognition tasks, including identification, categorization, etc. This information could then be "read out" by circuits receiving input from IT neurons (17–19).

Although physiological and functional imaging data suggest that visual object identity and category are coded in the activity of IT neurons (2–6, 8–16, 20), fundamental aspects

of this code remain under debate, including the discriminative power in relation to population size, temporal resolution, and time course. These questions must be understood at the population level to provide quantitative constraints for models of visual object recognition. We examined these issues by obtaining independent recordings from a large unbiased sample of IT neuronal sites and using a population readout technique based on classifiers. The readout approach consists of training a regularization classifier (21) to learn the map from neuronal responses to each object label (Supporting Online Material), as in recent studies in the motor system [e.g., (22)]. Instead of making strong assumptions about the prior probability distribution of the training examples, the classifier learns directly from them and generalizes to novel responses (21). The input consists of the neuronal responses from the independently recorded neurons; different input representations allow quantitative comparisons among neural coding alternatives (10, 13, 22–28). After training, the classifier can be used to decode the responses to novel stimuli. We used a one-versus-all approach whereby for each class of stimuli (8 classes for categorization, 77 classes for identification, 3 classes for scale and position readout; see below), one binary classifier was trained. The overall classifier prediction on test data was given by the binary classifier with the maximum activation. The performance of such classifiers

constitutes a lower bound on the information available in the population activity, but is a meaningful measure that could be directly implemented by neuronal hardware.

We used the classifier approach to determine the ability of more than 300 sequentially collected IT sites from two passively fixating monkeys to "categorize" 77 gray-scale objects as belonging to one of eight possible groups (29) (Fig. 1A). Figure 1B (red curve) shows the cross-validated performance of classifiers in performing this categorization task as a function of the number of recording sites (30). The spiking activity of 256 randomly selected multi-unit activity (MUA) sites was sufficient to categorize the objects with $94 \pm 4\%$ accuracy (mean \pm SD; for 100 sites, interpolated performance = 81%; chance = 12.5%). Similarly, we tested the ability of the IT population to identify each of the 77 objects (Fig. 1B, blue curve). Even small populations of IT neurons were capable of performing this identification task at high accuracy (for 256 sites, $72 \pm 3\%$ correct; for 100 sites, interpolated performance = 49%; chance = 1.3%), although at lower performance than categorization for the same number of sites (31). Classifier performance increased approximately linearly with the logarithm of the number of sites, which is indicative of a distributed representation in contrast to a grandmother-like representation (13, 28, 32, 33). Very similar levels of performance were obtained when single unit activity (SUA) was considered [Fig. 1C, (28)]. The local field potentials also contain information about object category [Fig. 1C, (28)]. Examination of the classification errors suggests that some objects and categories were easier to discriminate than others (Fig. 1D). All the results reported here were obtained using a linear (regularized) classifier. Classification performance was similar for several different types of classifiers, and the performance of linear classifiers—among the simplest classifiers—could not be substantially improved upon (28, 34).

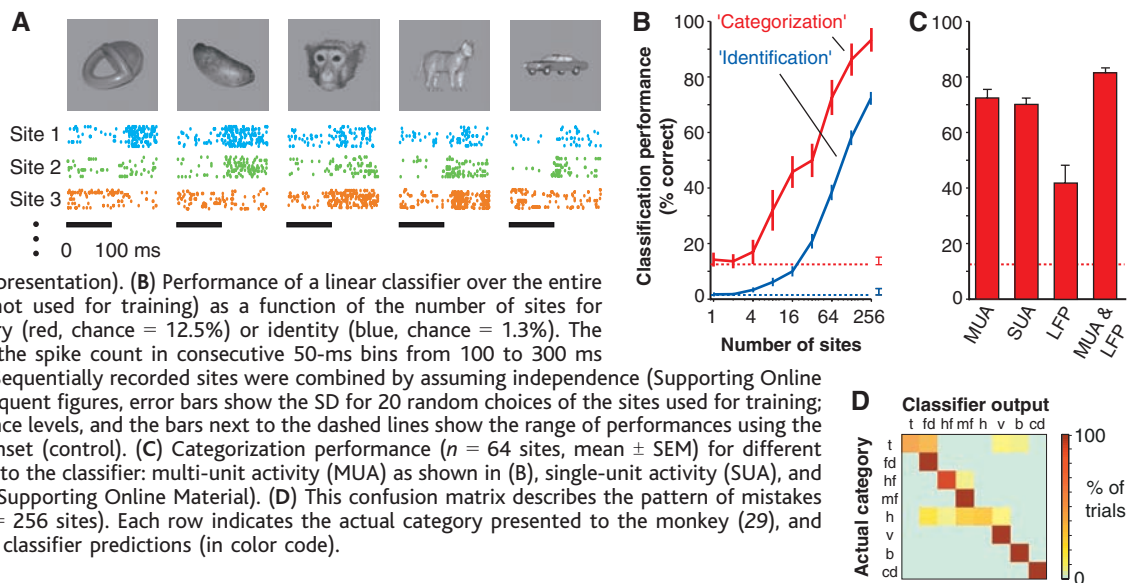
The performance values in Fig. 1, B to D, are based on the responses of single stimulus presentations that were not included in the classifier training. Thus, the level of recognition performance is what real downstream neurons could, in theory, perform on a single trial by simply computing a weighted sum of

¹McGovern Institute for Brain Research, ²Center for Biological and Computational Learning, ³Computation and Systems Biology Initiative (CSBI) at Massachusetts Institute of Technology (MIT), ⁴Department of Brain and Cognitive Sciences, MIT, Cambridge, MA 02139, USA.

*These authors contributed equally to this work.

†To whom correspondence should be addressed.
E-mail: chouhung@mit.edu

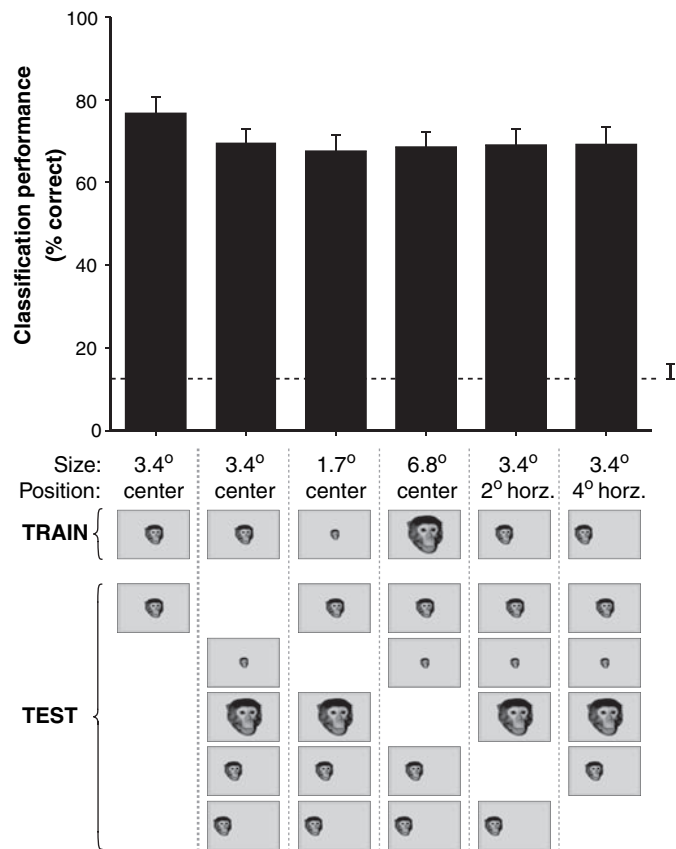
Fig. 1. Accurate readout of object category and identity from IT population activity. (A) Example of multi-unit spiking responses of 3 independently recorded sites to 5 of the 77 objects. Rasters show spikes in the 200 ms after stimulus onset for 10 repetitions (black bars indicate object presentation). (B) Performance of a linear classifier over the entire object set on test data (not used for training) as a function of the number of sites for reading out object category (red, chance = 12.5%) or identity (blue, chance = 1.3%). The input from each site was the spike count in consecutive 50-ms bins from 100 to 300 ms after stimulus onset (28). Sequentially recorded sites were combined by assuming independence (Supporting Online Material). In this and subsequent figures, error bars show the SD for 20 random choices of the sites used for training; the dashed lines show chance levels, and the bars next to the dashed lines show the range of performances using the 200 ms before stimulus onset (control). (C) Categorization performance ($n = 64$ sites, mean \pm SEM) for different data sources used as input to the classifier: multi-unit activity (MUA) as shown in (B), single-unit activity (SUA), and local field potentials (LFP, Supporting Online Material). (D) This confusion matrix describes the pattern of mistakes made by the classifier ($n = 256$ sites). Each row indicates the actual category presented to the monkey (29), and each column indicates the classifier predictions (in color code).



spikes over a short time interval (100- to 300-ms interval divided into bins of 50 ms in this case) (11, 23, 24, 28). This is notable considering the high trial-to-trial variability of cortical neurons (27). The IT population performance is also robust to biological noise sources such as neuronal death and failures in neurotransmitter release [fig. S1, (35)]. Although Fig. 1 (and most other decoding studies) assumes precise knowledge about stimulus onset time, this is not a limitation because we could also accurately read out stimulus onset time from the same IT population [fig. S5, (28)].

A key computational difficulty of object recognition is that it requires both selectivity (different responses to distinct objects such as one face versus another face) and invariance to image transformations (similar responses to, e.g., rotations or translations of the same face) (8, 12, 17). The main achievement of mammalian vision, and one reason why it is still so much better than computer vision algorithms, is the combination of high selectivity and robust invariance. The results in Fig. 1 demonstrate selectivity; the IT population can also support generalization over objects within predefined categories, suggesting that neuronal responses within a category are similar (36). We also explored the ability of the IT population to generalize recognition over changes in position and scale by testing 71 additional sites with the original 77 images and four transformations in position or scale. We could reliably classify (with less than 10% reduction in performance) the objects across these transformations even though the classifier only “saw” each object at one particular scale and position during training (Fig. 2). The “identification” performance also robustly generalized across position and scale (28). Neurons also showed scale and position invariance for novel objects not seen before (fig. S6). The IT population

Fig. 2. Invariance to scale and position changes. Classification performance (categorization, $n = 64$ sites, chance = 12.5%) when the classifier was trained on the responses to the 77 objects at a single scale and position (depicted for one object by “TRAIN”) and performance was evaluated with spatially shifted or scaled versions of those objects (depicted for one object by “TEST”). The classifier never “saw” the shifted/scaled versions during training. Time interval = 100 to 300 ms after stimulus onset, bin size = 50 ms. The left-most column shows the performance for training and testing on separate repetitions of the objects at the same standard position and scale (as in Fig. 1). The second bar shows the performance after training on the standard position and scale (scale = 3.4° , center of gaze) and testing on the shifted and scaled images of the 77 objects. Subsequent columns use different image scales and positions for training and testing.



representation is thus both selective and invariant in a highly nontrivial manner. That is, although neuronal population selectivity for objects could be obtained from areas like V1, this selectivity would not generalize over changes in, e.g., position (Supporting Online Material).

We studied the temporal resolution of the code by examining how classification per-

formance depended on the spike count bin size in the interval from 100 to 300 ms after stimulus onset (Supporting Online Material). We observed that bin sizes ranging from 12.5 through 50 ms yielded better performance than larger bin sizes (Fig. 3A). This does not imply that downstream neurons are simply integrating over 50-ms intervals or that no useful object information is contained in smaller time

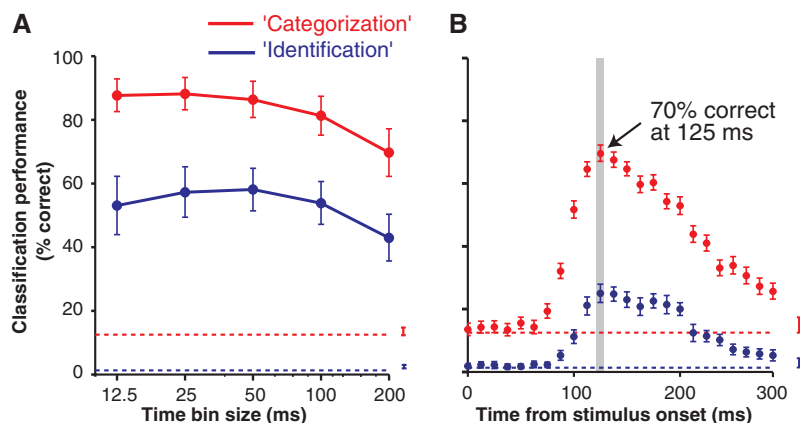


Fig. 3. Latency and time resolution of the neural code. (A) Classification performance ($n = 128$ sites) as a function of the bin size (12.5 to 200 ms, i.e., temporal resolution) to count spikes within the 100- to 300-ms window after stimulus onset for categorization (red) and identification (blue). The same linear classifier as in Figs. 1 and 2 was used. (B) Classification performance ($n = 256$ sites) using a single bin of 12.5 ms to train and test the classifier at different latencies from stimulus onset (x axis). The colors and conventions are as in Fig. 1B.

intervals. Indeed, we could decode object category at $70 \pm 3\%$ accuracy using only the spikes contained in one single bin of 12.5-ms duration at 125-ms latency (Fig. 3B). Notably, this time bin typically contained zero to two spikes (0.18 ± 0.26 spikes/bin, mean \pm SD). This shows that a few spikes from a small number of neurons (essentially a binary vector with either ones or zeros) are sufficient to encode “what” information in IT neurons within behaviorally relevant time scales.

What other “types” of information are carried in the IT population? Using the readout method, we compared the information available for “categorization” versus “identification” (18, 37, 38). The time course and temporal resolution did not depend strongly on the classification task (Fig. 3); the best sites for categorization overlapped the best sites for identification; the signal-to-noise ratios for categorization and identification were strongly correlated ($r = 0.54$, $p < 10^{-10}$); and the same randomly selected sites could be used for both tasks (28). The same IT neuronal population can thus be used by downstream neurons to perform tasks traditionally considered to be different (e.g., “categorization” versus “identification”).

Although anterior IT cortex is generally regarded as the brain area at the top of the ventral “what” stream, the readout approach allowed us to examine the possibility that the IT population might contain useful information about object scale and position (“where”). Our observation that IT populations convey scale- and position-invariant object category and identity information (Fig. 2) might seem to suggest that object position information is lost in IT neurons. However, it is also possible to read out—at least coarsely—both object scale and position (“where” information) based on the activity of the same population, independent of identity or category, by training

the classifier to learn the map between neuronal responses and scale or position, irrespective of object identity (fig. S4A). Reading out object position or scale had a similar time course to the readout of object category (fig. S4B). There was little correlation between the ability of each IT site to signal scale/position versus object category information, suggesting that IT neurons encode both types of information (fig. S4C).

Our observations characterize the available information in IT for object recognition, but they do not necessarily imply that the brain utilizes exclusively the IT neurons (39) or the same coding schemes and algorithms that we have used for decoding. However, a linear classifier—which we found to be very close to optimal (34)—could be easily implemented in the brain by summing appropriately weighted inputs to downstream neurons. Thus, targets of IT [such as prefrontal cortex (PFC)] could decode information over brief time intervals, using inputs from small neuronal populations (e.g., ~ 100 neurons). It is conceivable that the dynamic setting of the synaptic weights from IT to PFC may switch between different tasks in PFC, reading out information from the same neuronal population in IT cortex (18). In this perspective, some neurons in IT cortex would be similar to tuned units in a learning network, supporting a range of different recognition tasks including “categorization” and “identification” in PFC (40).

The approach described here can be used to characterize the information represented in a cortical area such as object identity in IT cortex (2–6, 8–11). Classifiers can be trained on any stimulus property and then tested to systematically examine putative neural codes for that stimulus information. Our results quantitatively show how targets of IT cortex may rapidly, accurately, and robustly perform tasks of categorization, identification, and readout

of scale and position based on the activity of small neuronal populations in IT cortex.

References and Notes

1. M. Fabre-Thorpe, G. Richard, S. J. Thorpe, *Neuroreport* **9**, 303 (1998).
2. D. Perrett, J. Hietanen, M. Oeom, P. Benson, *Philos. Trans. R. Soc.* **355**, 23 (1992).
3. N. K. Logothetis, D. L. Sheinberg, *Annu. Rev. Neurosci.* **19**, 577 (1996).
4. K. Tanaka, *Annu. Rev. Neurosci.* **19**, 109 (1996).
5. K. Grill-Spector, R. Malach, *Annu. Rev. Neurosci.* **27**, 649 (2004).
6. C. Keyers, D. K. Xiao, P. Foldiak, D. I. Perret, *J. Cogn. Neurosci.* **13**, 90 (2001).
7. R. vanRullen, S. Thorpe, *Vision Res.* **42**, 2593 (2002).
8. E. Rolls, *Curr. Opin. Neurobiol.* **1**, 274 (1991).
9. R. Desimone, T. Albright, C. Gross, C. Bruce, *J. Neurosci.* **4**, 2051 (1984).
10. P. M. Gochin, M. Colombo, G. A. Dorfman, G. L. Gerstein, C. G. Gross, *J. Neurophysiol.* **71**, 2325 (1994).
11. B. Richmond, R. Wurtz, T. Sato, *J. Neurophysiol.* **50**, 1415 (1983).
12. N. K. Logothetis, J. Pauls, T. Poggio, *Curr. Biol.* **5**, 552 (1995).
13. L. F. Abbott, E. T. Rolls, M. J. Tovee, *Cereb. Cortex* **6**, 498 (1996).
14. S. L. Brincat, C. E. Connor, *Nat. Neurosci.* **7**, 880 (2004).
15. M. Ito, H. Tamura, I. Fujita, K. Tanaka, *J. Neurophysiol.* **73**, 218 (1995).
16. G. Wallis, E. T. Rolls, *Prog. Neurobiol.* **51**, 167 (1997).
17. M. Riesenhuber, T. Poggio, *Nat. Neurosci.* **2**, 1019 (1999).
18. D. Freedman, M. Riesenhuber, T. Poggio, E. Miller, *Science* **291**, 312 (2001).
19. H. Tomita, M. Ohbayashi, K. Nakahara, I. Hasegawa, Y. Miyashita, *Nature* **401**, 699 (1999).
20. A. Ishai, L. G. Ungerleider, A. Martin, J. L. Schouten, J. V. Haxby, *Proc. Natl. Acad. Sci. U.S.A.* **96**, 9379 (1999).
21. T. Poggio, S. Smale, *Notices Am. Math. Soc.* **50**, 537 (2003).
22. K. V. Shenoy et al., *Neuroreport* **14**, 591 (2003).
23. L. F. Abbott, *Q. Rev. Biophys.* **27**, 291 (1994).
24. R. deCharms, A. Zador, *Annu. Rev. Neurosci.* **23**, 613 (2000).
25. A. S. Tolias, A. G. Siapas, S. M. Smirnakis, N. K. Logothetis, *Soc. Neurosci. Abstr.* **557.5** (2002).
26. A. Pouget, P. Dayan, R. Zemel, *Annu. Rev. Neurosci.* **26**, 381 (2003).
27. C. Koch, *Biophysics of Computation* (Oxford Univ. Press, New York, 1999).
28. <http://cbcl.mit.edu/software-datasets/kreiman/fastreadout.html>.
29. The eight groups used for categorization [toys (t), foods (fd), human faces (hf), monkey faces (mf), hand/body parts (h), vehicles (v), box outlines (b), cats/dogs (cd)] were defined before the experiments. Unsupervised clustering of neuronal responses yielded similar groups (28). Categorization became substantially worse upon arbitrarily defining these groups as sets of random objects (28). The discriminability for individual sites for passive and active viewing were similar (fig. S7, Supporting Online Material).
30. We assumed independence among neurons; this assumption should be revisited upon recording simultaneously from many neurons because correlations may contain additional information. Our estimate represents a lower bound on the information represented by small neuronal populations. However, even under these conditions, we obtain a high degree of accuracy [see also (41)].
31. Throughout the paper we randomly selected a given number of sites for decoding. The brain could be selectively wired such that targets of IT receive stronger input from the most relevant features. A simple feature selection step before the input to the classifier to select the sites with the highest signal-to-noise ratio (Supporting Online Material) showed that high performance levels could be achieved using a much smaller number of sites (fig. S2).
32. R. Quiroga, L. Reddy, G. Kreiman, C. Koch, I. Fried, *Nature* **435**, 1102 (2005).
33. M. P. Young, S. Yamane, *Science* **256**, 1327 (1992).
34. A linear classifier is given by $y = \text{sign}[f(x)]$, where y

is a binary label, x is the input vector (coding neural activity), and $f(x)$ is a linear function of the form $f(x) = w \cdot x + b = \sum_i c_i(x \cdot x_i) + b$. Training means estimating the vector of coefficients w and the scalar b from the training set of m (x_i, y_i) pairs, where x_i is the "input" part of each example and y_i is its associated label or "output." More complex classifiers of the form $f(x) = \sum_i c_i k(x, x_i)$ had very similar performance and were no better than the regularized linear classifiers for $n > 64$ sites. The estimated coefficients depend on regularization and are different for different regularization techniques (21).

35. Multiple sources of noise can affect the encoding of information. The performance of the classifier was very robust to deletions of substantial numbers of neurons during testing, simulating neuronal or syn-

aptic death (fig. S1A), and also to large proportions of deleted spikes (simulating failures in spike transmission or neurotransmitter release; fig. S1B).

36. We trained the classifier for the categorization task with 70% of the pictures and then tested it on the remaining 30% of the pictures. The performance was quite good and only slightly below the performance levels reported above (fig. S3; compare to Fig. 1).
37. Y. Sugase, S. Yamane, S. Ueno, K. Kawano, *Nature* **400**, 869 (1999).
38. R. Vogels, *Eur. J. Neurosci.* **11**, 1239 (1999).
39. B. Tjan, *Adv. Neural Inf. Processing Syst.* **13**, 66 (2001).
40. T. Poggio, E. Bizzi, *Nature* **431**, 768 (2004).
41. N. C. Aggelopoulos, L. Franco, E. T. Rolls, *J. Neurophysiol.* **93**, 1342 (2005).
42. We thank M. Kouh for the object recognition model; R. Quiroga and A. Kraskov for spike sorting; J. Mayo and J. Deutsch for technical support; CSBI for com-

puter cluster usage; and R. Desimone, N. Kanwisher, C. Koch, and T. Serre for comments on the manuscript. This research was sponsored by grants from NIH, NSF, and especially from the Defense Advanced Research Projects Agency and Office of Naval Research. Additional support was provided by Eastman Kodak Company, Daimler Chrysler, Honda Research Institute, The Pew Charitable Trusts, Whiteman fellowship (G.K.), and the McDermott chair (T.P.).

Supporting Online Material

www.sciencemag.org/cgi/content/full/310/5749/863/DC1

SOM Text

Figs. S1 to S7

References

19 July 2005; accepted 5 October 2005
10.1126/science.1117593

Neuronal Activity Regulates Diffusion Across the Neck of Dendritic Spines

Brenda L. Bloodgood and Bernardo L. Sabatini*

In mammalian excitatory neurons, dendritic spines are separated from dendrites by thin necks. Diffusion across the neck limits the chemical and electrical isolation of each spine. We found that spine/dendrite diffusional coupling is heterogeneous and uncovered a class of diffusionally isolated spines. The barrier to diffusion posed by the neck and the number of diffusionally isolated spines is bidirectionally regulated by neuronal activity. Furthermore, coincident synaptic activation and postsynaptic action potentials rapidly restrict diffusion across the neck. The regulation of diffusional coupling provides a possible mechanism for determining the amplitude of postsynaptic potentials and the accumulation of plasticity-inducing molecules within the spine head.

In mammalian excitatory neurons, synaptic stimulation triggers the flow of ions across the dendritic spine membrane, as well as the production of second messengers within the spine head. Buildup of signaling molecules, such as calcium or activated CaMKII (calcium/calmodulin-dependent protein kinase II), within the spine head activates regulatory cascades that lead to the modification of the enclosed synapse (1–4). Furthermore, stimulus-induced transport of proteins across the spine neck, such as CaMKII, protein translation initiation factors, and β -catenin, plays a role in synapse regulation and plasticity (5, 6). Thus, the regulation of diffusion across the spine neck offers a potentially powerful mechanism to control the efficacy and modulatory state of individual synapses.

We examined the regulation of the diffusional barrier posed by spine necks in rat hippocampal pyramidal neurons. Organotypic slice cultures were biolistically transfected with the photoactivatable green fluorophore PAGFP (7)

and the red fluorophore dsRed. Two-photon laser scanning microscopy (2PLSM) with illumination at 910 nm readily excites dsRed without photoactivation of PAGFP, revealing dendrites and spines that fluoresce in the red spectrum (Fig. 1). Focal illumination with a second laser tuned to 720 nm triggers two-photon activation of PAGFP (8), and the resulting green fluorescence can be subsequently monitored with 910-nm illumination. Photoactivation of PAGFP within individual spines triggers increases in fluorescence within the head that dissipate as activated PAGFP (PAGFP*) diffuses into the dendrite. The decay of the fluorescence transient in the spine head is well fit by a single exponential, yielding a time constant of equilibration (τ_{equ}) (9) of PAGFP* across the spine neck (Fig. 1, A to C). Repeated measurements (at 0.1 Hz) in individual spines over ~ 1.5 min yielded consistent values of τ_{equ} (fig. S1) with coefficients of variation (CVs) of ~ 15 to 20% (Fig. 1D). Conversely, τ_{equ} varied over a broad range from spine to spine (Fig. 1E, $n = 11/572$ cells/spines), with the majority of values ranging from 140 to 350 ms.

In a subset of spines, fluorescence did not decay appreciably in the sampling period of

1.9 s. For these spines, the barrier to PAGFP* movement across the neck was bidirectional, so that PAGFP* within the dendrite is able to diffuse away from the site of photoactivation but does not enter the spine head (Fig. 2, A and B; similar findings in 11 of 11 comparable spine/dendrite pairs). Conversely, PAGFP* diffuses from the dendrite into the heads of spines with less restrictive spine necks (Fig. 2, C and D; similar findings in 8 of 8 comparable spine/dendrite pairs). Thus, the lack of PAGFP* movement in a subset of spines results from a severe diffusional isolation imposed by the spine neck and not from aggregation or cross-linking of PAGFP within the head. Repeated measurements of τ_{equ} in these diffusionally isolated spines over prolonged periods revealed that the diffusional barrier is reversible and that large, apparently spontaneous reductions in τ_{equ} occur (Fig. 2, E and F; similar findings in 4 of 15 diffusionally isolated spines that were monitored repeatedly for >5 min).

We hypothesized that the heterogeneity of τ_{equ} results from active regulation of diffusional coupling in response to variability in neuronal and synaptic activity. Chronic manipulations of activity trigger homeostatic changes in synaptic parameters such as the number and composition of AMPA-type glutamate receptors (AMPARs) at the synapse (10, 11). Consistent with our hypothesis, 24 hours of incubation in the AMPAR antagonist NBQX shifted the distribution of τ_{equ} toward faster values (8/367 cells/spines; $P < 0.01$), whereas block of GABA_A receptors (GABA_ARs) with bicuculline shifted the distribution toward slower values (8/556 cells/spines; $P < 0.01$) (Fig. 3A). Similar results were obtained with measurements of dsRed diffusion by fluorescence recovery after photobleaching (fig. S2). In contrast, block of voltage-sensitive sodium channels (VSSCs) (6/438 cells/spines) or NMDA-type glutamate receptors (NMDARs) (7/449 cells/spines) by incubation in tetrodotoxin (TTX) or carboxypiperazin-4-yl-propyl-1-phosphonic acid (CPP), respectively, had no effect on the cumulative distribution of τ_{equ}

Department of Neurobiology, Harvard Medical School, 220 Longwood Avenue, Boston, MA 02115, USA.

*To whom correspondence should be addressed.
E-mail: bsabatini@hms.harvard.edu



Fast Readout of Object Identity from Macaque Inferior Temporal Cortex

Chou P. Hung, Gabriel Kreiman, Tomaso Poggio and James J. DiCarlo (November 3, 2005)
Science Translational Medicine **310** (5749), 863-866. [doi: 10.1126/science.1117593]

Editor's Summary

This copy is for your personal, non-commercial use only.

- | | |
|----------------------|--|
| Article Tools | Visit the online version of this article to access the personalization and article tools:
http://science.sciencemag.org/content/310/5749/863 |
| Permissions | Obtain information about reproducing this article:
http://www.sciencemag.org/about/permissions.dtl |

Science (print ISSN 0036-8075; online ISSN 1095-9203) is published weekly, except the last week in December, by the American Association for the Advancement of Science, 1200 New York Avenue NW, Washington, DC 20005. Copyright 2016 by the American Association for the Advancement of Science; all rights reserved. The title *Science* is a registered trademark of AAAS.