

Corollary Discharge Provides Accurate Eye Position Information to the Oculomotor System

Barton L. Guthrie, John D. Porter, David L. Sparks

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Abstract. The saccadic system accurately compensates for perturbations of eye position produced by microstimulation of the superior colliculus. This requires that information about the stimulation-induced change in eye position be provided by an extraretinal source—either proprioceptive endings in extraocular muscles or a centrally generated corollary discharge. It is shown that compensation remains intact after elimination of extraocular muscle proprioception, demonstrating that corollary discharge provides accurate eye position information.

Precise information about the position of the eyes in the orbit is required for localization of visual targets (1) and is an essential component of current models of the oculomotor system (2). The question of whether eye position signals originate from a central copy of the oculomotor command or arise peripherally from extraocular muscle proprioceptors is a fundamental issue in oculomotor physiology.

A century ago Helmholtz concluded that knowledge of eye position was derived from a "measure of the effort of will required to move the eyes" (3). Such an internal signal representing an intended change in eye position was referred to as corollary discharge by Sperry (4). Recently, physiological studies have shown that a number of brain areas contain neurons which have tonic firing rates correlated with eye position (5). These neuronal populations are functionally close to the final oculomotor output and project to other centers involved in the control of eye movements. Such physiological and anatomical properties suggest that these neurons generate a corollary discharge which could be used to represent eye position. Unlike most other structures under motor control, the eyes are not subject to changing external loads, so a given oculomotor output always produces the same movement. Because of this tight coupling of motor output with the resulting movement, a corollary of the motor command would indeed accurately reflect eye position.

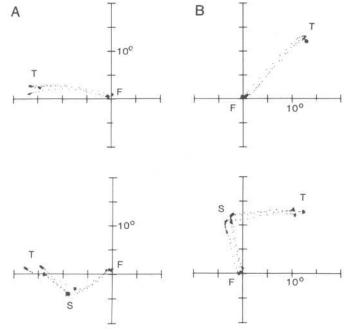
On the other hand, mammalian extraocular muscles are endowed with welldeveloped proprioceptors (6). Noting the presence of these receptors, Sherrington (7) proposed that extraocular muscle proprioception was the source of eye position information. Recent physiological studies have demonstrated that afferent fibers from extraocular muscles carry signals corresponding to muscle length and tension and are distributed to almost every known visuomotor center (8).

In this study we tested the hypothesis that corollary discharge alone provides accurate eye position information. After elimination of extraocular muscle proprioception (9), we used a saccadic tracking task (10) which required rapid and accurate feedback about eye position. A monkey looked at a lighted fixation target in an otherwise dark room. As this target was extinguished a saccadic target was presented for 50 msec at a selected location in the visual field. On randomly selected trials, after the sac-

Fig. 1. (A) and (B) represent postoperative trials at two different collicular stimulation sites, performed in the absence of extraocular muscle proprioception and plotted in a plane Cartesian coordinate system. Trials were performed in total darkness, with initial fixation at lighted target Coincident with F_{-} the offset of F, saccadic target T was presented for 50 msec. Saccades FT, shown on the upper axes, are three saccades to target location T without collicular stimulation. Shown on the lower axes are three stimulation-compensation trials using the same

cadic target was extinguished, but before the onset of the saccade to it, eye position was changed in a direction away from the target by microstimulation of the superior colliculus. After a brief pause at the new eye position, the animal made a compensatory saccade to the original saccadic target location. The compensatory saccade was different from the saccade that would have been generated based solely upon a retinal error signal (distance and direction of the target retinal image from the fovea). For compensation to occur, the saccadic system must be informed of the stimulationinduced change in eye position by an extraretinal source. If extraocular muscle proprioception is an essential source of eye position information, without it the oculomotor system would not be capable of generating an accurate compensatory saccade. On the other hand, if corollary discharge is used as an accurate eye position signal, the animals would continue to compensate in the absence of extraocular muscle proprioception.

Two rhesus monkeys were used in the experiment. Each was implanted with scleral search coils (11) and trained to make saccades to visual targets (12). Skull cylinders were implanted over appropriately placed small craniectomies for microelectrode stimulation of the su-



target locations. In these trials, after the offset of target T but before saccade FT could be initiated, the eyes were driven away from the target to location S by collicular stimulation. After a brief pause at S, compensatory saccades ST were made to the location of the (now absent) target T. This compensation could have occurred only if the saccadic system received information about the stimulation-induced change in eye position from F to S. Since retinal and proprioceptive feedback about these eye movements were not available, a centrally generated corollary discharge must have provided the eye position information. If the saccadic system had not been informed of the stimulation-produced movement from F to S, saccades with the same vector as FT but starting from eye position S would have been expected.

perior colliculus. Display of visual targets, monitoring of eye position, delivery of reward, and stimulation of the colliculus were computer-controlled (13). Prior to surgery, each animal was tested in the stimulation-compensation task by using systematically varied target locations. Then, with a subtemporal approach, the ophthalmic nerves were transected bilaterally at their junction with the trigeminal ganglion, eliminating extraocular muscle proprioception. Complete transection was ascertained postoperatively by loss of the corneal blink reflex, then confirmed postmortem by the absence of labeling of trigeminal ganglion sensory neurons after injection of horseradish peroxidase into the extraocular muscles. Intraoperative damage to the oculomotor nerves was ruled out by normal action of the extraocular muscles postoperatively (14). After surgery each monkey was tested in the stimulation-compensation task.

Preoperatively, compensation was tested for three collicular stimulation sites using two to five target locations per site. Each compensatory saccade exhibited a normal magnitude-velocity relation and followed a 20- to 60-msec fixation at the end of the stimulationinduced saccade. Compensation was accurate, showing an average error of 4.5 degrees, ranging from 1.6 to 6.8 degrees, varying with target location and stimulation vector. This performance is comparable to that described in an earlier, more extensive study of stimulation-compensation behavior (13). After elimination of extraocular muscle proprioception, compensation remained intact (Fig. 1). Four collicular sites were stimulated using three to six target locations per site. As was the case preoperatively, each compensatory saccade had a normal waveform and followed a brief fixation after the stimulation-produced saccade. Accuracy of compensation was not significantly different, showing an average error of 4.0 degrees, ranging from 2.1 to 7.3 degrees.

Postoperative compensation is possible only if information about the stimulation-induced eye movement is still available. Three possible sources of this eye position signal are the retina, extraocular muscle proprioception, and a centrally generated corollary discharge. In this study the dark experimental environment and the absence of the saccadic target during and after the stimulationinduced saccade prevented retinal feedback about change in eye position. Transection of the ophthalmic nerves eliminated extraocular muscle proprioception. Thus, eye position information must have been provided by a centrally generated corollary discharge. This experiment, while demonstrating that corollary discharge provides accurate eye position information, does not address the possibility that extraocular muscle proprioception may provide information for some oculomotor functions (15).

Although previous investigations have suggested that motor systems distribute corollaries of their output (16), none has unequivocally demonstrated a functional role for these internal signals. This study, however, clearly demonstrates that the oculomotor system generates a corollary discharge which accurately represents the intended motor act and is used in generating further movements.

BARTON L. GUTHRIE

JOHN D. PORTER DAVID L. SPARKS

Department of Physiology and **Biophysics and Neurosciences** Program, University of Alabama in Birmingham, Birmingham 35294

References and Notes

- 1. I. P. Howard, Human Visual Orientation (Wiley, New York, 1982), p. 275. 2. D. A. Robinson, in Their Basic Mechanisms of
- Ocular Motility and Their Clinical Implications, D. Bach-y-Rita and G. Lennerstrand, Eds. (Per-gamon, Oxford, 1975), p. 337; J. van Gisbergen, D. A. Robinson, S. Gielen, J. Neurophysiol. 45. 417 (1982); D. Zee, L. Optican, J. Cook, D. A. Robinson, W. Engel, Arch. Neurol. 33, 243 (1976); S. Yasui and L. R. Young, Science 190, 906 (1975).
- 3. H. von Helmholtz, Treatise on Physiological Optics (Dover, New York, 1962); D. M. Weiter, New Fork, 1962), D. M. Mackay, in Handbook of Sensory Physiology VII (3A), R. Jung, Ed. (Springer-Verlag, New York, 1973), p. 307.
 R. W. Sperry, J. Comp. Physiol. Psychol. 43, 482 (1950); E. V. Evarts, Neurosci. Res. Prog. Bull. 9, 86 (1971).
- Among sites containing neurons with tonic firing rates proportional to eye position are the abdu-cens internuclear neurons [S. M. Highstein, A. cens internuclear neurons [S. M. Highstein, A. Karabelas, R. Baker, R. A. McCrea, J. Comp. Neurol. 208, 369 (1982)], cerebellar vermis [H. Noda and D. A. Suzuki, J. Physiol. (London) 294, 349 (1979)], medial vestibular nucleus [R. A. McCrea, K. Yoshida, A. Berthoz, R. Baker, Exp. Brain Res. 40, 468 (1980)], nucleus prepositus hypoglossi [R. McCrea and R. Baker, Anat. Rec. 196, 122A (1980)], oculomotor nucleus [C. Evinger, R. Baker, R. A. McCrea, R. Spencer, in Progress in Oculomotor Research, A. Fuchs and W. Becker, Eds. (Elsevier/North-Holland, New York, 1981), p. 263], frontal eye fields [E. Bizzi, Exp. Brain Res. 6, 69 (1968)], and paramedian pontine reticular formation [E. L. Keller, J. Neurophysiol. 37, 316 (1974)]. Neurophysiol. 37, 316 (1974)].
- T. Greene and R. Jampel, J. Comp. Neurol. 126, 547 (1966); S. Cooper and P. M. Daniel, Brain 72, 1 (1949); G. L. Ruskell, J. Neurocytol. 7, 693 (1978)
- C. S. Sherrington, Brain 1, 332 (1918).
- C. S. Sherrington, Brain 1, 332 (1918).
 Areas involved in visuomotor function which receive extraocular muscle proprioceptive signals are the cerebellum [A. F. Fuchs and H. H. Kornhuber, J. Physiol. (London) 200, 713 (1969)], superior colliculus [P. K. Rose and V. C. Abrahams, Brain Res. 97, 95 (1975)], mesencephalic tegmentum and posteroventral thalamus [E. Manni, G. Palmieri, R. Marini, Exp. Neurol. 37, 412 (1972)], lateral geniculate [I. M. L. Donaldson and R. A. Dixon, Exp. Brain Res. 38, 245 (1980)], visual cortex [P. Buisseret and L. Maffei, *ibid.* 28, 421 (1977)], frontal cortex [H. Barbas and B. Dubrovsky, Exp. Neurol. 74, 67 (1981)].
- (1981)].
- In the monkey we have determined the location of cell bodies providing extraocular muscle sen-sory receptors by using the technique of retrograde transport of horseradish peroxidase after its injection into the muscles [J. D. Porter, B. L.

Guthrie, D. L. Sparks, Soc. Neurosci. Abstr. 8, 156 (1982), J. D. Porter, B. L. Guthrie, D. L. Sparks, J. Comp. Neurol. 218, 208 (1983)]. After examining the oculomotor nerves, trigeminal ganglia, and brainstem of three species of ma-caque (Macaca mulatta, M. nemistrina, and M. fascicularis), labeling was found only in ipsilat-eral trigeminal ganglia. Section of the ophthal-mic nerve at its junction with the trigeminal ganglion prior to injection of horseradish peroxi-dase into ipsilateral extraocular muscles eliminated all labeling of trigeminal sensory neurons

- 10. L. E. Mays and D. L. Sparks, Science 208, 1163 (1980).

- (1980).
 A. F. Fuchs and D. A. Robinson, J. Appl. Physiol. 21, 1068 (1966).
 D. L. Sparks and R. Holland, Behav, Res. Methods Instrum. 7, 115 (1975).
 D. L. Sparks and L. E. Mays, J. Neurophysiol. 49, 45 (1983); D. L. Sparks and J. D. Porter, ibid. p. 64 *ibid.*, p. 64. 14. We have found that conjugate eve movements
- are unaffected by the loss of extraocular muscle

proprioception [B. L. Guthrie, J. D. Porter, D.

- L. Sparks, Soc. Neurosci. Abstr. 8, 156 (1982)]. Psychophysical studies suggest that extraocular muscle proprioception may function in ocular stabilization, but probably is not significant in the localization of visual targets [A. A. Skavenski, in Eye Movements and Psychological Processes, R. A. Monty and J. W. Senders, Eds. (Erlbaum, Hillsdale, N.J., 1976), p. 277; G. S. Brindley and P. A. Merton, J. Physiol (London) 153, 127 (1960)].
- 153, 127 (1960)].
 D. I. McCloskey, in Handbook of Physiology, vol. 2, The Nervous System, V. B. Brooks, Ed. (Williams & Wilkins, Baltimore, 1981), p. 1451.
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