Rapid elbow movements in patients with torsion dystonia

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SUMMARY Rapid, self-paced and self-terminated elbow flexion movements were studied in a group of 10 patients with dystonia affecting the arms. The movements were slower and for small amplitude movements, more variable than those recorded in normal subjects. The duration of the first agonist burst was prolonged, even when compared with normal subjects deliberately moving slowly. Cocontraction of agonist and antagonist muscles during ballistic movements was common and may contribute to the bradykinesia. These findings are compared with similar studies of other diseases of the motor system. Unlike many other conditions which also reduce the speed of ballistic voluntary movements, the patients with dystonia in the present study showed a normal symmetry of acceleration and deceleration times. One interpretation of this finding is that aspects of the basic motor programmes are relatively preserved in this condition and account for the surprising retention of motor skills shown by some patients with dystonia.

The abnormal patterns of muscle activation that produce dystonic limb postures in patients with torsion dystonia are usually worse during voluntary movement.1-4 Indeed, in its early stages dystonia may be evident only during certain actions such as walking or handwriting. Muscle spasms, jerks and cocontraction of agonist and antagonist muscles distort the limb posture and interrupt the intended sequence of movement. Despite these problems, surprising degrees of motor skill are often retained by patients with advanced torsion dystonia. For example, they may demonstrate great proficiency in the use of hand-held communicators, manually operating tools, or as in the case of two of our patients, become accurate marksmen.4 It is therefore of some interest to examine the performance of simple motor tasks in patients with upper limb dystonia and the extent to which the above abnormal patterns of muscle activity interfere with the normal simple ballistic movement EMG pattern. Rapid, self-paced and self-terminated movements at a single joint are achieved by a triphasic pattern of muscle activity in the agonist, antagonist and agonist muscles.5-7 This pattern has been studied for various disorders of movement10-16 and the results from this study in patients with torsion dystonia are compared with those described in patients with other diseases of the basal ganglia, cerebellum, and upper motor neurone pathways.

Patients and methods

The patients comprised three females and seven males, aged 15 to 79 (mean age 47), all of whom had dystonia affecting one or both limbs. Nine patients had primary (idiopathic) torsion dystonia; three of these had segmental dystonia affecting the neck and arms, five had generalised dystonia and one had focal dystonia of an arm. The remaining patient had symptomatic hemidystonia due to an arteriovenous malformation of the contralateral basal ganglia. Six patients were receiving medication (clonazepam, baclofen, benzhexol) at the time of the study. All patients and normal subjects gave informed consent for the studies which were approved by the local ethical committee.

Elbow flexion movements were examined with the subjects seated in a chair and their arm abducted to 90°. The forearm was semipronated and rested on a horizontal manipulandum pivoted about the axis of rotation of the elbow joint. The elbow position was measured by a potentiometer and displayed on an oscilloscope screen in front of the subject. Electromyographic activity (EMG) was recorded by Ag/AgCl surface electrodes placed on the following muscles;
biceps, triceps, posterior deltoid, pectoralis major and the flexors and extensors of the fingers and wrist in the forearm.

Position, velocity (electronically differentiated from the position) and rectified EMG activity were recorded for each elbow flexion movement with a PDP12 computer, (sampling rate 250 Hz per channel), using programmes devised by HB Morton. EMG signals were amplified using a Devices 3160 preamplifier with high and low pass filters set at 80 Hz and 2.5 kHz (3 dB points) respectively, and a Devices 3120 amplifier.

All subjects were asked to make elbow flexion movements of 15° and 30° as fast as they could from a starting angle of 120°. After some practice, 10 to 15 trials for each kind of movement were recorded. Results were compared with values obtained from nine normal subjects aged 29 to 67 (mean age 44). Normal subjects were then asked to perform these movements at roughly half their maximal speed, imitating the velocity of the movements of the dystonic patients.

Each single trial was inspected on the computer display unit. The amplitude, peak velocity of movement and the duration of the first agonist (biceps) EMG burst were measured. The first agonist EMG burst in normal subjects was relatively well defined with clear onset and offset. However, in some patients this was not the case. The duration of the first agonist EMG burst was often difficult to measure since the point at which it terminated was poorly defined. In these patients, the criterion for determining the end of the agonist burst was that the EMG should decline to less than 10% of the peaksize for at least 20 msec. This decrease was expected before the peak velocity of movement. If the first burst of EMG activity continued for more than 50 msec after the peak velocity for any individual movement, then the burst was said to be "tonic" and not to have a measurable duration. The first antagonist EMG burst was not analysed in these trials. The duration of the movement was measured from the velocity trace and was defined as the time between the onset to the first zero crossing. Movement time was divided into acceleration time (time from the beginning of the movement until the peak velocity) and deceleration time (time from moment of peak velocity until zero velocity). The coefficient of variation for the movement amplitudes were calculated from the mean and standard deviation for each size of movement to give an indication of movement variability. Statistical analysis of the data was performed using unpaired Student's t-tests and a mixed model analysis of variance (ANOVA).

Results

A typical elbow flexion movement through an angle of 30° performed by a normal subject moving as fast as possible is illustrated in figure 1a. When a patient with dystonia attempted to perform a similar movement of comparable size, as fast as possible (figure 1c), two major differences were evident: i) the movement of the patient was slower than the fast movement of the normal subject and ii) the duration of the first agonist EMG burst (Ag1) in the biceps of the patient was prolonged. The question which arises is whether or not the patients' movements are similar to those of normal subjects moving at a comparable velocity (that is, slowly).

To investigate this question we asked the normal subjects to perform a series of movements to the same 15° and 30° targets when moving at half their maximal speed. An example of a deliberate slow movement in a normal subject is illustrated in figure 1b (the same subject shown in figure 1a). The duration of Ag1 was still shorter than that of the patient moving at a similar speed. Data were analysed from all subjects and patients for both 15° and 30° movements and the following comparisons were made. First, movements of normal subjects moving at their maximal speed were compared with their movements at half their maximal speed. Second, the movements of the patients with dystonia were compared with the slow and fast movements of normal subjects.

Normal subjects

All nine normal subjects were able to perform the 15° and 30° movements with ease, whether moving at maximum speed, or about half that value. There was a tendency for the fast movements to overshoot the target distance before settling down to the final end.
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Fig 2. Comparison of various parameters of elbow flexion movements made through an angle of 15° or 30° by normal subjects moving as fast as possible and moving at about half their maximal speed and patients with dystonia. Histograms show the mean values and SE. Statistical details are given in the text (a). The maximum velocity of the elbow flexion movements measured as degrees per second. The dystonic patients performed the movements at the same speed as normals moving slowly. (b) The duration of the 15° and 30° movements in the normal group and patients with dystonia was similar except when the normal subjects moved slowly (means and SE are shown). (c) The variability from trial to trial is expressed as the coefficient of variation of the movement amplitude. The coefficient of variation for the 15° movements of the dystonic patients was large (means and SE are shown). (d) The first agonist EMG burst (Ag1) in biceps was measured according to the criteria described in Methods. Although the maximum velocity of the movements of the dystonic patients was similar to that of the slow moving normal subjects (fig 2a), the duration of the Ag1 EMG bursts was prolonged compared to fast and slow moving normal subjects.

The duration of the movement was obviously longer in the slow movements. Comparison of the duration of fast and slow movements in the 15° and 30° tasks yielded a significant interaction term in the ANOVA (F(1,8) = 82.98, p < 0.05), indicating that the movement duration increased more between 15° and 30° movements in the slow than in the fast task (figure 2b). This was a result of the prolonged deceleration phase of the 30° slow movement. The variability of movement amplitude (expressed as coefficient of variation) was the same whether subjects moved fast or slowly (F(1,8) = 6.15, p > 0.05) (figure 2c). The duration of Ag1 also was the same in both movements (F(1,8) = 0.06, p > 0.05) (figure 2d).

In normal subjects the first burst of EMG activity in the agonist muscle was followed by a discrete burst of EMG activity in the antagonist muscle. The behaviour of the antagonist burst was similar to that previously described by Marsden, et al but has not been
forearm and forearm flexors and elbow flexion inappropriate for the target.

Patients with dystonia

Nine of the 10 patients were able to perform the required elbow flexion movements and flex their elbow to approach the desired target. The mean (SD) amplitudes of movement in the patient group were slightly smaller [15° (4°) for the 15° target and 29° (5°) for the 30° target] than those achieved by the normal subjects moving slowly (F(1,16) = 4.76; p < 0.05) or as fast as possible (F(1,16) = 23.66; p < 0.001).

The fastest elbow flexion movements of the patients with dystonia were slower than the fastest movements of normal subjects (F(1,16) = 68.96, p < 0.001) analysed in detail here because of the difficulties in measuring data from the patients (see Methods). EMG activity in other muscles of the arm (forearm flexors, extensors, pectoralis major, deltoid) showed an alternating pattern of EMG activity. The forearm flexors, pectoralis major and posterior deltoid were activated at the same time as the biceps; the forearm extensors were activated later or at the same time as the triceps (figure 3a).

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Fig 3 Single trials of an elbow flexion movement to a 30° target in a normal-subject (a) and a patient with dystonia (b). The top two traces show the position and velocity records while the remainder show rectified EMG recordings from biceps, triceps, forearm flexors and extensors, pectoralis major and deltoid muscles. The burst duration of the agonist muscle (biceps) in the elbow flexion movement is prolonged in the patient with dystonia, and the timing of agonist and antagonist muscle contractions is inappropriate for the movement. Note also the prolonged activation of pectoralis major (overflow) and cocontraction of the forearm and shoulder muscles.
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(figure 2a). Although the velocity of 30° movements was greater than that of 15° movements in both groups, the increment in velocity was much smaller in the patients, giving rise to a significant interaction in the ANOVA (F(1,16) = 9.04, p < 0.05). In contrast, the slow movements of the normal subjects were executed at the same speed as the fastest movements of the patients (F(1,16) = 0.34, p > 0.05) (see figure 2a).

Acceleration and deceleration times were the same as in the slow normal group except for the 30° movement. When normals performed a slow 30° movement, the deceleration time was much longer than in the patients (table 1). This was because when the normal group moved slowly over a long distance, they tended to glide into the final target whereas the patients did not. The result of this can be seen in the comparison of movement durations. The duration of 15° movements was the same as in the patients as in the slow normal group, but in the 30° task, the patients took less time than the normal group moving slowly (figure 2b). This difference gives rise to the interaction term in the comparison of movement duration (F(1,16) = 29, p < 0.001).

The amplitude of attempted 15° movements was more variable from trial to trial in the patients than in normal subjects, whether the latter were moving slow or fast (Student’s t test p < 0.01) (figure 2c). The variability of 30° movements was the same in all groups (Student’s t test p > 0.05). The duration of the first agonist EMG burst was longer in the patients for both 15° and 30° tasks whether compared with normal subjects moving fast (F(1,16) = 16.43, p < 0.001), or slow (F(1,16) = 16.15, p < 0.001) (figure 2d).

Several patterns of antagonist EMG activity were observed. These varied from a normal appearance to cocontraction often with prolonged bursts or tonic EMG activity without recognisable bursts. The patterns of EMG activity in the forearm flexors and extensors, and in the pectoralis major and posterior deltoid muscles, also varied from relatively normal patterns, with appropriate alternating activity in these agonist—antagonist muscle pairs, to various patterns of prolonged bursts in these muscle groups. An example of a movement in a patient with arm dystonia is shown in figure 3b. Cocontraction of the agonist (biceps) and antagonist (triceps) muscles can be seen, and there abnormal activation of the pectoralis major muscle (overflow).

Among the 10 patients studied, the findings in the patient with symptomatic hemidystonia were similar to those in patients with idiopathic dystonia. One severely affected individual with generalised idiopathic torsion dystonia was unable to perform the elbow flexion task. On attempting to flex the elbow he could only produce movements of 40–60° and was unable to make movements of smaller amplitude. The accompanying EMG bursts were greatly prolonged and frequently had to be classified as tonic (see Methods), while the maximum velocity was 232° (SD 40°) per second was even slower than the other patients.

Discussion

Rapid self paced and self terminated movements at a single joint have now been studied in patients with a wide variety of disorders of voluntary and involuntary movement.10-16 Our results extend this analysis to patients with torsion dystonia. Their elbow flexion movements differed from normal in the following ways: 1) the peak velocity was slower, 2) the extent of movement was more variable for the small, but not the large amplitude task, 3) the duration of the first burst of agonist EMG activity was prolonged even when compared with the normal group who had been purposely instructed to move slowly, at a similar speed to the patients, 4) there often was cocontraction of the antagonist muscle.

The movements of dystonic patients, however, were normal in one respect. The acceleration and deceleration times were approximately equal indicating that the velocity profile was bell shaped.17 The same profile was seen in the fast movements of the normal subjects, although it became skewed towards longer deceleration times when the same normal individuals were instructed to move slowly, particularly in the 30° task. The bell shaped velocity profile of the dystonic patients indicates that despite the abnormal pattern of EMG activity, at least this aspect of the ballistic motor programme appears to remain intact.

Table 2 compares the deficits in the present dystonic patients with those reported in other neurological conditions. Interestingly the movements of all groups are slower than normal, perhaps indicating that rapid movements of normal subjects are optimal and that their velocity cannot be exceeded in any pathological state. Although all groups move more slowly than normal, combinations of increased movement variability, asymmetric velocity profile and prolonged and/or cocontracting EMG activity distinguish between several of the groups. Not surprisingly, the voluntary movements of patients with dystonia appear very similar to those of patients with athetosis12 and patients with Huntington’s disease,14 although there is a lack of data on the velocity profiles of the latter condition. Despite the clinical differences between the involuntary sustained muscle spasms which produce relatively fixed postures in dystonia (and athetosis) and the continual flow of movement in chorea, the control of simple voluntary movements exhibits similarities in the two conditions. The motor symptoms of dystonia, athetosis and chorea are...
thought to be due to abnormal function of the basal ganglia, so the common deficits of voluntary movement in all three conditions may reflect disruption of the normal basal ganglia contribution to rapid simple movements. The prolonged, excessive and variable patterns of voluntary muscle activity seen in patients with dystonia, athetosis and chorea may indicate, as Hallett suggests, that the basal ganglia play a role in selectively activating only those muscles appropriate for a particular task and inhibiting those which are not required. Other physiological studies in dystonia also point to excessive muscle activation and/or lack of normal inhibition of muscle activity. The R2 component of the blink reflex is prolonged and its recovery curve hyperexcitable in patients with blepharospasm, there is a reduction in the amount of Ia presynaptic inhibition in patients with dystonia affecting the arm, the duration of the long-latency stretch reflex in the wrist flexor muscles is prolonged. The EMG pattern responsible for the slow movements of the patients with Parkinson’s disease is quite different from that of dystonic, athetoid and choreic patients. The agonist and antagonist bursts are timed correctly and appropriate muscles are activated in Parkinson’s disease, but the EMG bursts are smaller than necessary to produce the same rapid movements shown in the normal group. Why then does this classic disease of the basal ganglia produce different abnormalities to those seen in dystonia, athetosis, and chorea? We have suggested previously that the basal ganglia circuitry as presently conceived (for example 21) allows for the possible coexistence of bradykinesia and hyperkinesia. In essence there may be at least two circuits directly concerned with movement within the basal ganglia; abnormalities of the one involved with the initiation of a voluntary movement may result in bradykinesia; abnormalities in the other, which may be involved in minimising unwanted or extraneous movement could lead to dyskinesias. One or both circuits may be differentially affected in different disease states. Thus, in Parkinson’s disease, loss of the dopaminergic nigrostriatal input may affect the former pathway whereas both might be affected in dystonia, athetosis and chorea.

Finally, why are the movements of patients with dystonia slower than normal subjects? The patients are not weak and their EMG bursts are not shorter than normal. If we assume that the EMG-force relation of the muscles is normal, three other possibilities emerge: 1) peak velocities may be reduced because the agonist EMG burst is prolonged; a large agonist burst may be capable of producing normal peak velocities, if excessively prolonged would take the limb beyond the target point. To prevent overshoot, the burst would have to be selected to be smaller and, hence, peak velocity inevitably would be reduced; 2) action of the agonist muscle might be opposed by inappropriate cocontracting activity of the antagonist; 3) there may be a primary difficulty in producing maximum bursts of phasic muscle force even though voluntary strength is intact. In the most rapid contractions, moto-neurones discharge phasically at rates of up to 200 Hz. Presumably a very large descending input is needed to produce these rates and the pathology of dystonia may make this impossible; the 30–40 Hz rate for a normal fused voluntary contraction might be achieved relatively easily, but the resulting contraction would be slower.

In conclusion, these experiments have shown that rapid, self-paced and self-terminated movements of patients with dystonia are slower and more variable than normal. Deficits in the duration of the first agonist EMG burst and in control of antagonist and synergist muscles are similar to those seen in patients with Huntington’s disease or athetosis. Such deficits may reflect a common contribution of the basal ganglia to the control of ballistic movements, perhaps involving the grading of EMG agonist burst size and duration and the selection and timing of appropriate agonist and antagonist muscle activity.
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