Matters arising

A double-blind controlled trial of high dose methylprednisolone in multiple sclerosis

Sir: In relation to the observation of Milligan et al on the antispastic effect of steroids in chronic progressive multiple sclerosis, we would like to point out that 10 years ago we published in Chile, a report on the subject. We reported the use of prednisone [1 mg per kg weight], in seven patients with spastic paraparesis. The patients had the following diagnosis: one case of chronic multiple sclerosis, one case of cervical rachitostenosis, and five cases of unknown aetiology without the clinical and pathological characteristics of multiple sclerosis. A method was designed for the evaluation of the spasticity. Using this method we observed clinical improvement of the spasticity score in all these patients. We think that the effect reported by Milligan et al and our own report may be due to a specific antispastic effect of steroids. On this respect we agree with Compston et al, on the possibility of a direct action of steroids on muscle fibres. One possibility would be an effect upon the intrasural muscle fibre, because no important change was observed on striated muscle fibres.

We believe that further clinical and experimental studies are necessary to define specific roles and mechanisms of the antispastic effect of corticosteroids.

Luis Cartier R
Renato Verdugo L
Neurosciences Department,
School of Medicine
Universidad de Chile
Leonardo da Vinci 7221
Las Condes, Santiago
Chile

References


Nifedipine and the treatment of myotonia

Sir: The claim by Grant et al of significant improvement in the myotonia of 10 patients with myotonic dystrophy following nifedipine treatment seems exaggerated. Firstly, in the crossover study, the valid statistic compares placebo with nifedipine 10 mg and by the authors' own admission there was no significant difference between these two treatments in any of several objective tests. Yet, they attempt to convince us that there are significant differences between nifedipine and initial groups but not between placebo and initial groups in the finger extension time test as presented in their fig 3. This figure is appalling in its lack of essential information (paired data are not indicated) and it contains erroneous information. Although it is stated that 20 results are used in the analysis only 19 are visible in the placebo group and a recalculation of the mean places it at about 1.5 s, not 1.7 s as stated and illustrated. This puts the mean for placebo on a level with those of the nifedipine treatments. Alternatively, if the mean plotted in their figure is correct then the missing 20th point must be an outlier at greater than 5 s and off scale above the figure. In either case it appears highly doubtful that there is anything to the results other than a possible slight improvement in finger extension times during the course of the study due to familiarity with the testing procedure. In any case the trial has been flawed by its single blind methodology. The possibility of differential encouragement to perform well in the finger extension test has not been controlled and could not be excluded even if there were any significant differences between placebo and nifedipine treatment groups.

Furthermore, there is no theoretical basis for the use of calcium channel blockers in the treatment of myotonia. As far as is understood, the prolonged contractions (delayed relaxations) in myotonia are due to myogenic action potential after-discharge. Such after-discharges would be indistinguishable from voluntary tetanic action potentials at the level of calcium channels involved in excitation-contraction coupling. Therefore at dosages where therapeutic depression of myotonic contractions would be achieved, equal depression of voluntary contraction would be apparent as an intolerable side effect. Weakness is already a debilitating feature of most myotonic conditions and particularly so, due to the dystrophic process, in myotonic dystrophy.

In vitro experimentation provides us with no expectation of antmyotonic action from calcium channel blockers. Nifedipine (up to 30 μM), and nicardipine (up to 50 μM) have proven ineffective as antmyotonic agents in the rat diaphragm myotonic by blocking CI- channels with either 2,4-dichloroephenoxyacetate or with antracene-9-carboxylate (unpublished observations). It can be inferred that Ca++ channels were blocked in these experiments without effect on the myotonia since nifedipine (5 μM) is known to block Ca++ channels in rat extensor digitorum longus muscle. As added proof, Ca++ channels are similarly blocked by Cd++ (0-1-1M) but Cd+ at these concentrations also blocks CI- channels and is a powerful inducer of myotonia, the attendant absence of Ca++ conductance notwithstanding.

Finally, it needs to be remembered that myotonic stiffness is seldom a major concern of patients with myotonic dystrophy who are much more likely to be debilitated by other aspects of their disease. Treatments for myotonia, then, must prove less of a nuisance than the myotonia itself and must not aggravate other symptoms of the disease. Cardiac conduction abnormalities are frequent in myotonic dystrophy and calcium channel blockers are known to have adverse effects on conduction. Other adverse reactions to these agents are also common. Considerably better clinical, experimental and theoretical evidence of nifedipine's efficacy as an antmyotonic agent would need to be presented before I was convinced that it had any benefit to weigh against its disadvantages. Other treatments, backed up by success in double blind trials, supported by in vitro studies and soundly theoretically based, are available.

A H BRETAG
School of Pharmacy,
South Australian Institute of Technology,
Adelaide, SA 5000, Australia.

References

A double-blind controlled trial of high dose methyl prednisolone in multiple sclerosis.
L Cartier and R Verdugo

*J Neurol Neurosurg Psychiatry* 1988 51: 316
doi: 10.1136/jnnp.51.2.316

Updated information and services can be found at:
http://jnnp.bmj.com/content/51/2/316.1.citation

**Email alerting service**
Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

**Notes**

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/