Idiopathic polyneuritis: serial studies of nerve and immune functions

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SUMMARY The clinical features of idiopathic polyneuritis in 22 patients admitted to the University of Kentucky Medical Center between July 1965 and June 1970 are described. Serial studies of peripheral nerve function in 12 patients followed to clinical recovery showed changes in nerve conduction velocity, distal motor latency, and/or muscle action potential amplitude. Six children and one adult, showing a change in all three parameters, exhibited a strong correlation between the degree of slowing and the shortness of their illness. Assessment of immune status, by quantitative measurement of cerebrospinal fluid immunoglobulin G (11 patients) and by the degree of lymphocyte transformation on exposure to specific brain protein and/or a homogenate of sciatic nerve (10 patients), bore no consistent relation to the severity or course of the illness. A double-blind trial of short-term, high dose adrenocorticotrophic hormone has yielded no valid evidence to date for or against the effectiveness of such therapy in this illness.

The natural history of idiopathic polyneuritis has been a subject of controversy since the earliest descriptions of this illness (Landry, 1859; Guillain Barré, and Strohl, 1916; Wiederholt, Mulder, and Lambert, 1964). Some divide the disease into various clinical categories which reflect the duration and severity of illness in their patients (Ravn, 1967). Others have called for strict diagnostic criteria which, if violated, imply possibly a separate illness, certainly one which follows a different course (Osler and Sidell, 1966). Still others find no correlation between persistence of deficit and age, speed of progression, objective sensory loss, cerebrospinal fluid pleocytosis, steroid therapy, or nerve conduction studies done many years after the onset of symptoms (Pleasure, Lovelace, and Duvoisin, 1968).

It is important to settle this question, if for no other reason than to evaluate various forms of therapy. This will be a difficult task, as long as the cause of the disease remains obscure. Recent and convincing pathological evidence supports the concept that whatever the precipitating cause, the common pathological denominator is an intense immune response (Asbury, Arnason, and Adams, 1969).

This report deals with a group of 22 patients with idiopathic polyneuritis, followed at the University of Kentucky Medical Center during the five years since its Department of Neurology was started. Twelve of the patients were followed to clinical recovery with serial studies of peripheral nerve function. Two patterns of change in nerve function were observed. Patients who followed one pattern showed a high correlation between slowing of nerve conduction velocity and brevity of illness. Immune factors were studied in 13 patients, to assess the incidence and specificity of such factors and the degree of their correlation with the course and severity of the polyneuritis.

METHODS

During the period from 1 July 1965 through 30 June 1970, 23 patients were admitted to the Medical Center with a diagnosis of idiopathic polyneuritis. One of these patients is not included in this study—a child, seen at the age of 2 for slow motor development, who had a normal cerebrospinal (CSF) protein level (16 mg/100 ml) despite a slow motor nerve conduction velocity (11 m/sec) in his ulnar nerve. He was eliminated from the study not only because his CSF protein should have been elevated if he had idiopathic polyneuritis (Chambers and MacDermot, 1957), but also because nothing is known of the onset or subsequent clinical course of his problem.

Therefore, the patient group chosen for study comprises 22 subjects—eight adults and 14 children. During

1Supported in part by research grants from the American Cancer Society, Kentucky Division.
the five year period in question, there were 1,614 admissions to the adult neurology service (13,385 to adult medical beds in general), and 6,633 admissions to the children's service. This amounts to an incidence of 5 per 1,000 adult neurology admissions (0.6 per 1,000 adult medical admissions), and 2.1 per 1,000 paediatric admissions—or an overall incidence of 1.1 per 1,000 admissions.

Studies were made of peripheral nerve function in the patient group by standard methods previously described (McQuillen and Gorin, 1969). Where possible, these studies included measurement of motor and mixed nerve conduction velocities in an ulnar nerve; of distal latency—that is, the time from stimulus at the wrist to onset of the grouped muscle action potential in the hand—and of grouped muscle action potential amplitude. An ulnar nerve was chosen for study, in order that changes during serial studies might be compared with the normal variation known to occur in its function (McQuillen and Gorin, 1969). Random studies of other nerves (peroneal and tibial) did not yield data that were qualitatively different, even when involvement was more prominent in the lower extremities.

The motor nerve conduction velocities cited in this paper are modal velocities. This average velocity was felt to be more reliable for comparison in serial studies, since the amplitude of the evoked muscle action potential was often quite small. When the amplitude was small, it was difficult to judge accurately the time from stimulus to onset of the response, particularly with distal stimulation.

Quantitative measurement of immunoglobulin content in serum and CSF was performed by the single radial immunodiffusion method of Mancini, Carbonara, and Heremans (1965). Lymphocyte transformation was studied by culture of lymphocytes obtained from heparinized blood samples. The culture period was five days. Each series consisted of quadruplicate cultures containing approximately 1.5 × 10⁶ cells. Saline solution was added to control cultures. Either a homogenate of human sciatic nerve or specific brain protein was added as antigen to test cultures. Twelve hours before the end of the culture period 4 µc tritiated thymidine was added to each culture. At the end of the culture period the cells were washed to eliminate unincorporated isotope. The activity of incorporated isotope was determined in a scintillation counter. Disintegrations per minute were determined for each culture. Stimulation (or depression) is expressed as a per cent of controls.

RESULTS

CLINICAL FEATURES The characteristics of the patient group are outlined in Table 1. The categories are those of Ravn (1967). They are used because the duration of illness is felt to differ from one category

| TABLE 1 |

<table>
<thead>
<tr>
<th>Category</th>
<th>No.</th>
<th>Age (yr.)</th>
<th>Sex</th>
<th>Involvement at peak of illness</th>
<th>Duration (months)</th>
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<th>Therapy</th>
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<td></td>
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<td></td>
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<td>Extremity*</td>
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<td>M</td>
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<td>22</td>
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<td>M</td>
<td>0</td>
<td>+++</td>
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<td>0</td>
</tr>
</tbody>
</table>

*0 = no involvement; + = mild; ++ = moderate; +++ = severe; +++++ = no function.
†0 = absent; + = hypoactive; ++ = normal.
‡From peak weakness to recovery; patient lost to follow-up if no information is given; patient 5 is still recovering function.
to another, even in untreated patients. Each category is subdivided further by age, since idiopathic polyneuritis may be a milder disease in childhood than in adult life (Markland and Riley, 1967; Asbury et al., 1969; Paulson, 1970).

The most seriously ill patients are in category I. All patients in this category required tracheostomy and assisted respiration. Patients 2 and 3 could still swallow at the peak of their illness, but all others in this category required a nasogastric tube for nourishment. Patient 1 had a total external ophthalmoplegia, in addition to his other troubles.

Category II consists of patients with milder involvement of bulbar and spinal nerves, not requiring mechanical assistance to breathe or swallow. Children predominated in this category. Patients 18 and 19 had mild dysphagia and jaw weakness but could swallow effectively. Patient 8 had mild respiratory weakness but did not require assisted respiration.

In category III are patients with no bulbar weakness. Patient 22 had the onset of quadriplegia within less than a day, as did patient 3 in category I. Both of these patients had suggestively upgoing toes, during the acute phase of their illness only. Both had hyponatraemia that responded to severe restriction of fluids, much in the fashion of the syndrome of inapposite antidiuretic hormone secretion (Posner, Ertel, Kossman, and Scheinberg, 1967). Patient 3 developed a persistent cardiac arrhythmia that disappeared only when he was given high doses of prednisone. He had electrocardiographic changes interpreted by a cardiac consultant as evidence for a myocarditis.

Deep tendon reflexes were elicited despite the presence of weakness in two adults and three children. A mild stocking-glove hypalgesia was present in six patients. Moderate sensory loss was observed in patient 21, whose illness began a week after exposure to a concentrate of malathion. He was treated with a chelating agent because of a questionable increase in urinary lead excretion. There was no change in his clinical state and no increase in lead excretion during therapy. Marked vibratory sense loss was observed in patient 1 at the peak of his illness. The distribution of weakness in the extremities did not follow any standard pattern. It was more prominent proximally in 11 patients (nos. 1, 3, 7, 9-11, 16-19, and 21); of equal prominence proximally and distally in six patients (nos. 4, 5, 8, 12, and 20); and more prominent distally in the remaining five patients.

THREATENING WEAKNESS—That is, in category I—were treated with aqueous adrenocorticotrophic hormone (ACTH) or prednisone in high dosage, in view of the opinion that steroid therapy is of benefit in idiopathic polyneuritis (Heller and DeJong, 1963). Patient 6 was not given steroids since he had been taking isonicotinic acid hydrazide (INH) for two years before the onset of his polyneuritis. The INH was prescribed because of a family history of tuberculosis and a personal history of skin test conversion.

Beginning in March 1968, a random, double-blind trial was started to compare the effect of ACTH (1,000 units intramuscularly over a seven to 10 day period) with that of a placebo. Patient 3 in category I was the first subject to enter this trial. He developed hyponatraemia after two days of therapy. The code was broken, it was found that he was receiving ACTH, and this was stopped. Hyponatraemia improved on restriction of fluids. Three weeks later he developed a persistent cardiac arrhythmia thought to be due to a myocarditis. The arrhythmia disappeared only after he was given high doses of prednisone (80 mg by mouth per day). Because of this experience, the trial has been limited since then to patients in categories II and III. Separate random groups for children and adults were established. The two children (patients 9 and 12) who received ACTH during the trial had a relatively brief illness, of two to eight months, respectively. The duration of illness in children (patients 10 and 15) and adults (patients 7 and 20) receiving placebo ranged from 3-5 to 18 months.

Plasma cortisol levels and/or urinary steroid output were followed in six patients (nos. 1, 7, 9, 10, 12, and 20). Cortisol levels rose proportionately higher than did steroid output in the one patient (no. 9) in whom both sorts of determinations were done. These studies correctly identified the patients receiving ACTH, by virtue of a rise in cortisol level or steroid output, in all but one instance (patient 12). Despite the fact that this patient received ACTH she showed no significant rise in 17-ketosteroid or 17-ketogenic steroid excretion in the urine. The apparent failure to respond to ACTH may be an error of measurement, since plasma cortisol levels (if obtained) might have shown a rise not mirrored by changes in urinary steroid excretion. Such findings would indicate rapid metabolism or inactivation of ACTH, and in this way could be related to the duration of her illness. Patient 12 was given INH during and after ACTH treatment, in view of a personal history of pulmonary tuberculosis four years before the onset of her polyneuritis. No clinical activation of the tuberculosis occurred.

STUDIES OF PERIPHERAL NERVE FUNCTION Studies of peripheral nerve function were obtained in all but
two patients (nos. 8 and 19). Selected data from remaining 20 patients are recorded in Table 2. Nine patients had a modal motor nerve conduction velocity (modal NCV) within normal range for our laboratory at the time of first study. Eight patients showed an increase in distal latency. Only two patients had a muscle action potential of normal amplitude. In two patients (nos. 4 and 5) a muscle action potential could not be recorded, although a nerve action potential was seen proximally with stimulation of mixed nerve distally. Children showed the clearest deviation from normal in all parameters. However, there was considerable overlap among clinical and age categories.

Six adults and six children were followed to clinical recovery with serial studies. Two patterns of nerve function were observed. In pattern one, modal NCV remained nearly within a normal range during the course of the illness with the major change noticed in the amplitude of the evoked potential (Fig. 1) or in distal latency. In pattern two a marked change was seen in all parameters (Fig. 2). Peak abnormalities were observed from 7-10 weeks before to 10 weeks after the point of maximal weakness was reached. Recovery of normal function paralleled clinical recovery, although it lagged behind return to normal strength. Deep tendon reflexes returned to normal at essentially the same time as peripheral nerve function. Again, children showed the most severe changes. Mean modal NCV was 17.3±3.6 (m/sec) in children and 40.4±11.8 m/sec in adults, at the time of slowest study. Mean modal NCV was slower in pattern two (18.5±9.3 m/sec) than in pattern one (43.9±10 m/sec), at this time, also.

Attempts to find a correlation between changes in peripheral nerve function and clinical course were productive in pattern two only. In this instance there is a strong correlation between the degree of slowing of modal NCV and the duration of the illness (Fig. 3). The slowest conduction velocities were associated with the shortest clinical courses. Among the 12 patients followed to clinical recovery with serial studies, all six children and one adult (patient 21) followed pattern two.

CEREBROSPINAL FLUID (CSF) FINDINGS Spinal puncture was performed 47 times on the 22 patients. All but five punctures were made at the peak of weakness ± one month. Total CSF protein ranged from 7 to 300 mg/100 ml.

Serial studies were done on six adults and eight children. No consistent pattern of change was observed. Three children showed a decrease in total protein before peak weakness. One child and three

### Table 2

<table>
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<tr>
<th>Category</th>
<th>No.</th>
<th>Time (days)</th>
<th>V₁†</th>
<th>V₉†</th>
<th>Distal latency (msec)</th>
<th>Map Amp (mV)</th>
<th>Time*</th>
<th>V₁†</th>
<th>V₉†</th>
<th>Distal latency (msec)</th>
<th>Map Amp (mV)</th>
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<td>III</td>
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<td>20</td>
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</tbody>
</table>

*From onset of symptoms to first study.
†In metres per second; = action potential looked for but not seen; normal range: V₁ = 38.1-64.3; V₉ = 48.7-61.4.
‡From peak of weakness to time of study or recovery.
adults showed an increase after peak weakness. Four children showed an increase followed by a decrease after peak weakness.

Quantitative measurement of CSF immunoglobulin G (IgG) content was obtained in seven children and four adults (Table 3). Paired determinations were done in four children and three adults. IgG amounted to less than 10% of total protein on six occasions; more than 20% on four occasions; and between 10 and 20% on seven occasions. No consistent relation to peak weakness was noticed on single studies. No consistent change in level with reference to peak weakness was observed.

Quantitative determination of serum immuno-
Again, no of nine globulins of nine patients generally showed a normal content of IgG, IgA and IgM for age (Table 3). Although the total number of patients in this study is small with respect to other surveys of idiopathic polyneuritis there are some conclusions which seem warranted. In the first place, children with idiopathic polyneuritis may have an illness as serious and serious as adults.

**LYMPHOCYTE TRANSFORMATION** The ability of an antigen to alter the rate of transformation of lymphocytes in cell culture was studied in 10 patients (Table 4). Serial studies were done on three patients. Four patients were receiving steroids at the time of study. All studies were obtained at or after the peak of weakness.

A significant increase—that is, $x - 2 \text{SD} > 100\%$—in lymphocyte transformation rate was noted in four patients. In patient 20, the increase occurred on exposure to specific brain protein, as well as on exposure to homologous peripheral nerve. One patient showed a decrease in enhanced transformation, while receiving ACTH. Two other patients, on ACTH at the time of study, showed no enhanced transformation. The use of prednisone did not appear to alter the marked increase in transformation noted in patient 3. Patient 7, who showed no significant change in transformation when homologous peripheral nerve was used as an antigen, exhibited a significant decrease in the rate of transformation when specific brain protein was used as the antigen.

**DISCUSSION**

**TABLE 3**

**IMMUNOGLOBULIN DETERMINATIONS**

<table>
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<tr>
<th>Category</th>
<th>No.</th>
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<td></td>
<td></td>
<td>TP†</td>
<td>IgG†</td>
<td>IgG/TP × 100</td>
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<td>Before</td>
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*In days, relative to peak weakness.
†In mg per 100 ml.
‡x = ACTH (prednisone for the second set of values on patient 3), being given at the time of study.
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by
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was
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15-8%
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evoked
by
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wrist.
This
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Cerra
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stimulation
of
mixed
nerve
in
median
and
ulnar
nerves
in
one
man,
whose
illness
lasted
55
days.

Such
potentials
were
found
first
at
50
days,
when
clinical
recovery
was
almost
complete. Distal
motor
latency
was
never
less
than
5
to
6
msec
in
median
nerve,
in
studies
repeated
until
day
50. Isch-Treussard,
Buchheit,
and
Isch
(1962)
obtained
three
measurements
of
peroneal
motor
nerve
conduction
velocity
in
two
adult
patients,
and
two
measure-
ments
in
another. Studies
were
made
from
one
to
nine
months
after
the
onset
of
the
illness. Looking
at
velocity
and
distal
latency
only,
there
was
a
lack
of
correlation
between
the
most
marked
slowing
and
the
most
severe
disease.
In
general,
the
velocity
returned
to
normal,
in
a
fashion
parallel
to
but
lagging
behind
clinical
recovery.
In
a
later
paper,
the
same
authors
commented
that
these
changes
might
be
present
from
the
start
of
the
illness,
or
might
take
several
weeks
to
appear
(Isch,
Isch-Treussard,
Buchheit,
Delgado,
and
Kircher,
1969).
All
patients
studied
by
them
did
show
slowing
at
some
time,
in
erve
conduction
velocity
and
distal
latency
in
a
parallel
fashion.
Humphrey
(1964)
did
not
observe
a
return
of
ulnar
and
peroneal
nerve
conduction
velocities
to
normal
until
all
deep
tendon
reflexes
had
recovered.
Bergamini,
Gandiglia,
and
Fra
(1966)
studied
nerve
conduction
velocities
in
proximal
and
distal
segments
of
ulnar
(motor
and
mixed)
and
pero-
neal
(motor)
nerves
in
five
patients. Studies
were
made
within
four
weeks
of
the
onset
of
the
illness,
and
serially
thereafter
(at
four,
12,
24,
and
48
weeks).
Two
patients
showed
no
significant
change
at
any
time,
two
patients
had
pronounced
slowing
in
the

| TABLE 4
<p>| LYMPHOCYTE TRANSFORMATION |
|------------------------------------------|---------------------|</p>
<table>
<thead>
<tr>
<th>Category</th>
<th>No.</th>
<th>Antigen</th>
<th>Transformation</th>
<th>Therapy</th>
</tr>
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<td>I</td>
<td></td>
<td>P.N. x</td>
<td>B.P. x</td>
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<tr>
<td>I</td>
<td></td>
<td>x</td>
<td></td>
<td>108±12</td>
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<tr>
<td>II</td>
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<td></td>
<td>x</td>
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<td>107±15</td>
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<td></td>
<td>130±21</td>
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</table>

*PN = homogenized extract of human sciatic nerve; BP = specific brain protein.
†Expressed as a per cent of four control cultures, ±2 SD.
axilla-to-elbow segment of ulnar nerve, and one patient had pronounced distal slowing in peroneal nerve only. Other segmental changes were observed. Slowing in the peroneal nerve was not as great as clinical signs would suggest. The severity of slowing paralleled rapidity of onset, rather than the degree of clinical involvement. Pronounced slowing (especially distally) was more common in the younger patients. The major alteration (when seen) was present by eight weeks, with recovery occurring before 16 weeks.

Twelve patients in this study were followed to clinical recovery with serial observation of peripheral nerve function. The parameters measured included modal motor nerve conduction velocity, distal motor latency, and amplitude of the evoked muscle action potential. When a marked slowing in nerve conduction velocity occurs with a change in both of the other parameters (Fig. 2), there is a strong correlation between the degree of slowing and the length of the illness (Fig. 3). The slowest conduction velocities were associated with the shortest clinical courses. From studies of guinea-pig sciatic nerve, Kaeser and Lambert (1962) concluded that a slight alteration in myelin structure could greatly impair nerve conduction velocity, while selective destruction of large motor neurones does not impair velocity in fibres that remain excitable. They observed a marked slowing of velocity (to 30% of normal) in diphtheric polyneuritis but little slowing with Wallerian degeneration or thallium poisoning. A gradation of change, from mild segmental demyelination to severe axonal loss, is seen in peripheral nerves of patients who die of idiopathic polyneuritis (Asbury et al., 1969). Pattern two may reflect this gradation; pattern one does not exclude it. Whatever the explanation for the correlation between slowing and clinical course, it is important to consider this in assessing the benefit of any therapy for idiopathic polyneuritis.

Steroid therapy is based on the thesis that idiopathic polyneuritis is an immune disorder (Heller and DeJong, 1963). Although this thesis does not hold true for every patient (Drachman, Paterson, Berlin, and Roguska, 1970), it is an attractive one not only because of the pathological similarity between experimental allergic neuritis and idiopathic polyneuritis, but also in view of the demonstration of various immune phenomena in patients with idiopathic polyneuritis (Asbury et al., 1969). Circulating antibodies to peripheral nerve have been demonstrated in patients with idiopathic polyneuritis, but appear to be non-specific and not correlated with the clinical state of the patient (Melnick, 1963; Osuntokun, Prineas, and Field, 1966). Since such antibodies would be expected to comprise only a minute fraction of serum IgG, it is not surprising that total immunoglobulin levels—in our patients and in the literature (Dencker, Swahn, and Ursing, 1964; Wiederholt and Mulder, 1965)—show no striking abnormalities. In vitro techniques show no difference between CSF protein in idiopathic polyneuritis and the proteins in serum and CSF from normal individuals (Ravn and Jensen, 1965).

The common denominator in pathological studies of idiopathic polyneuritis is an inflammatory demyelinative neuritis, marked by focal, perivascular, lymphocytic infiltrates affecting any level of the peripheral nervous system (Asbury et al., 1969). Several lines of study have been used to support this evidence that immunity in idiopathic polyneuritis is cell-mediated. Buffy coat cells from patients have been shown to destroy myelin in vitro (Armason, Winkler, and Hadler, 1969). Mononuclear cells thought to be synthesizing deoxyribonucleic acid are found in the blood of patients with idiopathic polyneuritis, as well as a number of other disorders (Cook and Dowling, 1968; Cook, Dowling, and Whitaker 1970; Whitaker, Hirano, Cook, and Dowling, 1970). Such cells are found more commonly when the polyneuritis is acute, and in larger number in patients with the greatest morbidity (Cook et al., 1970).

Enhanced lymphocyte transformation on exposure of cultures of lymphocytes from patients with idiopathic polyneuritis to homologous peripheral nerve is recorded in the literature (Knowles, Saunders, Currie, Walton, and Field, 1969; Behan, Lamarche, Feldman, and Sheramata, 1970) and in this series. This method is thought to be a valid test in vitro for the presence of cell-mediated immunity (Mills, 1966). In this series, however, it was not a phenomenon specific to peripheral nerve as an antigen (Table 4). It did not occur regularly. This may derive in part from the observation that enhanced transformation can be suppressed by steroids (Ono, Terayama, Takaku, and Nakoo, 1968). Less transformation while on ACTH was observed in one patient. Two other patients, who showed no enhanced transformation, were on ACTH at the time of study. Nevertheless, the two patients (Table 4; nos. 3 and 15) who showed the greatest amount of transformation had the longest clinical course (27 and 18 months, respectively). Unfortunately, the data are insufficient to draw a correlation between the degree of activity and the duration of disease. Thus we cannot say that the patients with the most severe disease have the most intense immune reaction, as others have claimed (Cook et al., 1970).

Plasma and urinary steroids were determined in the laboratories of Dr. C. Charlton Mabry and Dr. William
Idiopathic polyneuritis: serial studies of nerve and immune functions

W. Wintenritz. Serum and cerebrospinal fluid immunoglobulins were measured in the laboratory of Dr. Kenneth Gerson. Lymphocyte transformation was studied by Ira Fowler.

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Idiopathic polyneuritis: serial studies of nerve and immune functions
Michael P. McQuillen

*J Neurol Neurosurg Psychiatry* 1971 34: 607-615
doi: 10.1136/jnnp.34.5.607

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