THE DIAGNOSTIC VALUE OF THE EXAMINATION OF THE CEREBROSPINAL FLUID

By J. N. CUMINGS, M.D., M.R.C.P.

Lumbar puncture is in most diseases safe but somewhat uncomfortable to the patient, yet while examination of the cerebrospinal fluid (C.S.F.) frequently yields most valuable aid to clinical diagnosis, there are many occasions when such examinations are disappointing. It seems best to limit discussion to specific diseases in which assistance to the clinician may be given by the pathologist.

Normal Cerebrospinal Fluid

Before turning to the findings in various diseases it is necessary to be certain of the normal range and of the alterations that may occur naturally, for example as a result of increasing age. Table 1 gives the ranges that are commonly accepted for most of the substances examined.

The exact figure for chlorides may depend to some extent on the blood chloride level, but the sugar level does not vary greatly from one time of the day to another as raised blood sugar levels following a carbohydrate meal are hardly reflected at all in the fluid, and when they are there is a considerable time lag. Even so it is best to try to take C.S.F. at about the same time if one patient is being followed from day to day.

Age affects the findings particularly in relation to protein content. Subjects of 60 or over very frequently show figures of up to 55 or 60 mg./100 ml., whereas children under 10 frequently have only 15 mg./100 ml. in the fluid and there is a corresponding variation between these ages. Pro-longed immobility in bed also tends to raise the protein content. The other substances commonly examined show little variation due to these causes.

The C.S.F. in Disease

Syphilis—Acquired

Primary. A small proportion (10 to 20 per cent.) of patients with primary syphilis have been shown to have a positive Wassermann and a lympho-cytosis in the fluid, but following adequate treatment only a very few retain these abnormalities. Such patients should still be treated and the vast majority will finally show no abnormality.

Secondary. An increase of cells of up to 50 per cmm. composed of lymphocytes, a raised protein of up to 100 mg./100 ml. and a positive Wassermann are not uncommon. The Lange in these patients shows a mid-zone curve. Treatment usually removes these abnormalities. Occasionally, a spinal block from an arachnoiditis may be found with a fluid similar to that seen in a block from a spinal tumour, but showing in addition a pleocytosis and a positive Wassermann.

Tertiary. Although three common types of disease involving the central nervous system are seen—a meningo-vascular lesion, tabes and general paralysis of the insane—yet other varieties of affection are possible, but these latter do not usually show any absolute diagnostic features in the C.S.F.

Table 2 gives in brief the type of results that are usually found in each of these three conditions with the chief distinguishing features such as type of cell and frequency of positivity of the Wassermann, together with the relative degree of positivity in blood and C.S.F. indicated respectively by the figures in brackets and the number of + signs.

Not every patient shows such clear-cut results at the first fluid examination. The Lange in particular does not always conform to these findings, and paretic curves such as 55 43 32 100 can be seen in meningo-vascular disease. Should such findings persist despite treatment, general paralysis of the insane must be seriously considered, as such curves will change to mid-zone or luetic curves in treated cases of tabes and meningo-vascular syphilis, but not in general paralysis of the insane. The Wassermann reaction in the fluid is much more easily influenced in meningo-vascular disease than it is in the blood; whereas in tabes and general paralysis of the insane prolonged treatment lowers the Wassermann levels in both blood and fluid much more slowly. In general paralysis of the insane an abnormal Lange curve and positive blood Wassermann may be found after prolonged treatment, even though the fluid Wassermann has become negative.

The Wassermann reaction is only positive in the
C.S.F. in syphilitic conditions, but a negative test does not exclude syphilis, and a fluid showing all the other findings, even with a negative Wassermann, only rarely occurs in other diseases. The exceptions to this that have been seen are in meningeal spread of a secondary carcinoma, in tuberculosis, rarely in disseminated sclerosis and in sarcoidosis.

**Syphilis—Congenital**

Some patients show no fluid changes. Cases of general paralysis of the insane usually show a slight pleocytosis, slight protein rise, a positive Nonne Apelt test and a Lange of paretic type with a positive Wassermann. These findings are often difficult to alter by treatment or at the best only show slow improvement.

**Poliomyelitis**

It is generally agreed that the fluid differs at varying stages of the disease and the usual findings are given in Table 3.

The particular features to note are that the cells decrease and the protein increases during paralysis. Should the protein be sufficiently raised (e.g. over 150 mg./100 ml.) then a fine coagulum may form, such as is seen in tuberculous meningitis. The differential diagnosis from this latter disease is that in poliomyelitis the sugar and chlorides are normal at all stages. Polymorphs do not occur in the pleocytosis of encephalitis, but are seen in about a quarter of the cases of poliomyelitis until towards the end of the first week of paralysis, after which time only lymphocytes are present. It is not easy by an examination of the C.S.F. to distinguish some cases of poliomyelitis from some forms of benign lymphocytic meningitis. Here there may be a slight reduction in chlorides, a lymphocytic pleocytosis and a raised protein content in the fluid fairly early in the disease. Nevertheless, it is rare to see a patient with lymphocytic meningitis in such an early stage as one may in poliomyelitis, e.g. before paralysis, so that the clinical picture answers the problem except in those cases of poliomyelitis without paralytic signs.

**Encephalitis**

The C.S.F. during the active stage is frequently under raised pressure. The fluid is clear and colourless and there is never a coagulum. Cells are increased and are always lymphocytes, together with a few mononuclears in a small number of patients. Protein is only very slightly raised with usually a negative Nonne Apelt test. While the chloride level is normal, the glucose content may frequently be raised. The Lange is frequently abnormal and, while a luetic type is the more common, paretic curves do occur.

Patients with inclusion body encephalitis show a normal fluid apart from the Lange which is paretic in type.

In post encephalitic cases changes in the fluid are not seen, apart from slight protein increase and perhaps a very low luetic type Lange.

**Meningitis**

There are three main sub-groups to be discussed which, according to their aetiology, can be described as pyogenic, tuberculous or virus meningitis.

Although the first of these is not quite so commonly seen now as formerly, the findings in the C.S.F. can be recorded as in Table 4.

Exact figures are, of course, impossible to give as they depend on the length of infection and the severity of the condition. However, the important features are that in pyogenic conditions the polymorphs outnumber the lymphocytes in the cell count, whereas the reverse is usually, but not invariably, true in tuberculous meningitis. Sugar is absent or almost absent in pyogenic cases, but in tuberculous meningitis there is only a reduction in amount and no alteration is seen in virus meningitis.

The finding of the responsible organism is, naturally the most important single diagnostic feature. It should and can be found in nearly every case of tuberculous meningitis—by guinea pig inoculation if necessary—but in practice we have not failed to demonstrate it if sufficient fluid (e.g. 8 ml.) and time are used; even in one case a small child with only 12 cells one organism was seen after a prolonged search, and post-mortem revealed the diagnosis as correct.

It cannot be too frequently stated that the sugar content of the C.S.F. is the surest guide to prognosis. Given a rising sugar level, whatever the cell, protein or chloride estimations, the patient is improving; both this and the reverse have been seen many times during streptomycin treatment alone, as well as in cases shown me in Zurich under continuous P.A.S treatment together with streptomycin.

It must always be remembered that intrathecal streptomycin in tuberculous meningitis almost invariably gives rise to an increase in cells and protein, so that no prognostic significance can be placed on results of the examination of the fluid as regards these two components. It has, moreover, been recently shown that streptomycin itself in no way affects the sugar level of the fluid.

The differential diagnosis between virus meningitis and poliomyelitis is very difficult when no paralysis is present in the latter condition, as has already been mentioned.

Another difficulty that arises is in sarcoidosis.
DIAGNOSTIC VALUE OF EXAMINATION OF CEREBROSPINAL FLUID

### Table 1
**Normal Cerebrospinal Fluid**

<table>
<thead>
<tr>
<th></th>
<th>Lumbar Fluid</th>
<th>Cisternal Fluid</th>
<th>Ventricular Fluid</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Appearance</strong></td>
<td>Clear and colourless.</td>
<td>Clear and colourless.</td>
<td>Clear and colourless.</td>
</tr>
<tr>
<td><strong>Cells</strong></td>
<td>Up to 4 lymphocytes per c.mm.</td>
<td>1 to 2 lymphocytes per c.mm.</td>
<td>1 lymphocyte per c.mm.</td>
</tr>
<tr>
<td><strong>Protein</strong></td>
<td>Up to 30 mg./100 ml.</td>
<td>About 15 mg./100 ml.</td>
<td>Up to 10 mg./100 ml.</td>
</tr>
<tr>
<td><strong>Globulin</strong> (e.g. Nonne Apelt)</td>
<td>Mostly lymphocytes, few large mononuclears.</td>
<td>Lymphocytes, a few large mononuclears.</td>
<td>Luetic, i.e. mid-zone.</td>
</tr>
<tr>
<td><strong>Chlorides</strong></td>
<td>Usually less than 100 mg./100 ml.</td>
<td>Usually under 100 mg./100 ml.</td>
<td>Usually 150 mg./100 ml.</td>
</tr>
<tr>
<td><strong>Lange</strong></td>
<td>Variable but usually mid-zone.</td>
<td>++</td>
<td>++ (in 70 per cent.)</td>
</tr>
<tr>
<td><strong>Sugar</strong></td>
<td>No change.</td>
<td>++ (in 70 per cent.)</td>
<td>++ (in 100 per cent.)</td>
</tr>
<tr>
<td><strong>W.R.</strong></td>
<td>Negative.</td>
<td>++</td>
<td>++</td>
</tr>
</tbody>
</table>

### Table 2
**The C.S.F. in Syphilis**

<table>
<thead>
<tr>
<th>Pressure</th>
<th>Meningo-Vascular</th>
<th>Tabs</th>
<th>G.P.I.</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Appearance</strong></td>
<td>Clear and colourless.</td>
<td>Clear and colourless.</td>
<td>Clear and colourless.</td>
</tr>
<tr>
<td><strong>Cells</strong></td>
<td>20 to 150 per c.mm.</td>
<td>10 to 50 per c.mm.</td>
<td>10 to 50 per c.mm.</td>
</tr>
<tr>
<td><strong>Type of Cell</strong></td>
<td>Mostly lymphocytes, few large mononuclears.</td>
<td>Lymphocytes, a few large mononuclears.</td>
<td>Usually lymphocytes and large mononuclears.</td>
</tr>
<tr>
<td><strong>Protein</strong></td>
<td>Usually less than 100 mg./100 ml.</td>
<td>Usually under 100 mg./100 ml.</td>
<td>50 to 150 mg./100 ml.</td>
</tr>
<tr>
<td><strong>Nonne Apelt Test</strong></td>
<td>+</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td><strong>Lange</strong></td>
<td>Variable but usually mid-zone.</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td><strong>W.R. C.S.F. Blood</strong></td>
<td>+</td>
<td>+</td>
<td>+</td>
</tr>
</tbody>
</table>

### Table 3
**C.S.F. in Poliomyelitis**

<table>
<thead>
<tr>
<th>Appearance</th>
<th>Clear and colourless.</th>
<th>Clear and colourless.</th>
<th>Clear and colourless or with a coagulum.</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Cells</strong></td>
<td>Up to 50 (can be 2,000), per c.mm. Polymorphs, 25 per cent. Lymphocytes, 75 per cent.</td>
<td>5 to 10 per c.mm., usually lymphocytes.</td>
<td>4 per c.mm.</td>
</tr>
<tr>
<td><strong>Protein</strong></td>
<td>Normal or only slightly raised.</td>
<td>Raised → 100 mg./100 ml.</td>
<td>→ 300 mg./100 ml.</td>
</tr>
<tr>
<td><strong>Nonne Apelt Test</strong></td>
<td>Negative.</td>
<td>Positive.</td>
<td>Positive.</td>
</tr>
<tr>
<td><strong>Chlorides and Sugar</strong></td>
<td>Normal.</td>
<td>Normal.</td>
<td>Normal.</td>
</tr>
<tr>
<td><strong>Lange</strong></td>
<td>No change.</td>
<td>Normal.</td>
<td>Normal or slightly mid-zone.</td>
</tr>
</tbody>
</table>

### Table 4
**C.S.F. in Meningitis**

<table>
<thead>
<tr>
<th>Pressure</th>
<th>Pyogenic</th>
<th>T.B.</th>
<th>Virus</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Appearance</strong></td>
<td>Raised + +</td>
<td>Raised + +</td>
<td>Slight increase.</td>
</tr>
<tr>
<td><strong>Cells</strong></td>
<td>Turbid in varying degree. Hundreds or thousands. Polymorphs.</td>
<td>Clear or slightly opalescent. Tens or hundreds.</td>
<td>Usually clear.</td>
</tr>
<tr>
<td><strong>Type of Cell</strong></td>
<td></td>
<td></td>
<td>Tens, occasionally hundreds.</td>
</tr>
<tr>
<td><strong>Protein</strong></td>
<td>100+ mg./100 ml.</td>
<td>Usually more lymphocytes than polymorphs (2 or 3 to 1).</td>
<td>Lymphocytes.</td>
</tr>
<tr>
<td><strong>Globulin</strong></td>
<td>Positive.</td>
<td>Positive.</td>
<td>Up to 150 mg./100 ml.</td>
</tr>
<tr>
<td><strong>Chlorides</strong></td>
<td>Decreased to about 650 mg./100 ml.</td>
<td>Decreased to 600 to 650 mg./100 ml.</td>
<td>Positive.</td>
</tr>
<tr>
<td><strong>Sugar</strong></td>
<td>Absent.</td>
<td>Decreased to 20 to 35 mg./100 ml.</td>
<td>Slight decrease—680 mg./100 ml.</td>
</tr>
<tr>
<td><strong>Direct Smear</strong></td>
<td>Causal organism.</td>
<td>T.B.</td>
<td>Normal.</td>
</tr>
<tr>
<td><strong>Culture</strong></td>
<td>Causal organism.</td>
<td>T.B.</td>
<td>Nil.</td>
</tr>
</tbody>
</table>

**CEREBROSPINAL F.LUID IN C.S.F. OF MENINGITIS**

**BEFORE PARALYSIS**

- Clear and colourless.
- Up to 50 (can be 2,000), per c.mm. Polymorphs, 25 per cent. Lymphocytes, 75 per cent.
- Normal or only slightly raised.

**FIRST WEEK OF PARALYSIS**

- Raised → 100 mg./100 ml.
- Positive.
- Normal.
- 0011210000.

**SUBSEQUENT WEEKS**

- Clear and colourless or with a coagulum.
- 4 per c.mm.
- → 300 mg./100 ml.
- Positive.
- Normal.
- Normal or slightly mid-zone.

**DIAGNOSTIC VALUE OF EXAMINATION OF CEREBROSPINAL FLUID**

**G.P.I.**

- Clear and colourless.
- 10 to 50 per c.mm.
- Usually lymphocytes and large mononuclears.
- 50 to 150 mg./100 ml.

**PARETIC, I.E. 1ST STAGE.**

- Positive.
- Normal.
- Normal or slightly mid-zone.

**CEREBROSPINAL Fluid IN C.S.F. OF Tabes Meningo-Vascular Before Paralysis.**

- Clear and colourless.
- Up to 760 mg./100 ml.
- No change.
- Negative.

**Clear and colourless.**

- Up to 10 mg./100 ml.
- Negative.
- 700 to 760 mg./100 ml.
- No change.
- Negative.

**Clear and colourless.**

- Up to 10 mg./100 ml.
- Negative.
- 700 to 760 mg./100 ml.
- No change.
- Negative.
Two or three cases have been seen and all showed a fluid similar to that seen in tuberculous meningitis, except that the sugar was never decreased to such a low level and tubercle bacilli were never found. Post-mortem alone revealed the truth, for in one case a negative Kveim's test was also obtained.

No verified case of torulosis has been seen, but here the sugar is not greatly reduced and the organism should be found or cultured.

**Cerebral Abscess**

A really deep-seated abscess may give no more changes in the C.S.F. than may a similarly sited tumour. This type of abscess is rare and oedema is common around an abscess, hence it is usual to find a raised protein level. A moderate pleocytosis is about half as common as a raised protein and the cells are usually lymphocytes, although polymorphs are not infrequently seen in small numbers. When the abscess is near the ventricular surface a considerable increase in cells, especially in polymorphs, is present. Fluids from such cases also show a lowered sugar and chloride content, but the more chronic abscess with few cells gives rise to little depression in the levels of sugar, and chloride levels in such cases are frequently unaltered. It is uncommon to find organisms in the fluid.

**Cerebral Tumour**

The value of the examination of the C.S.F. is limited in tumours of the brain, and this is reflected in the fact that one-tenth of the number of lumbar fluids from such cases are now examined as compared with the years before the last war. However, when lumbar puncture is performed there is in many cases either a high normal or a raised C.S.F. pressure. Frequently pressure figures of 200 to 300 mm. are found. Examination of the fluid reveals a raised cell count in about 10 per cent. of cases, with a raised protein level in some 80 per cent. of cases. Very occasionally tumour cells can be found. Raised cell counts are found in tumours in close apposition to the ventricles or at the base of the brain. Such patients also show increased protein and while malignant gliomata are more commonly associated with higher figures than the more innocent cerebral tumours, this is not always true, the pontine glioma is a typical exception.

Neurofibromata of the eighth cranial nerve are almost invariably associated with a high protein content and not infrequently the C.S.F. is xanthochromic in appearance. Secondary carcinomatous deposits in the brain frequently show a raised protein and this is rather more common than in malignant gliomata as there is not uncommonly more oedema around the former than the latter.

The value of the examination of the fluid from the lateral ventricles is now well recognized. A raised protein content in the fluid from one side as compared with the other is an almost sure indication that the side of the tumour is the same as that of the raised protein level.

**Spinal Tumour**

Manometry is of vital importance here. The usual Queckenstedt test is adequate and an absence of any significant rise in the lumbar manometer on jugular compression is indicative of a spinal block above the level of the puncture. When the block is incomplete there may be a slow and partial rise followed by a similar slow fall. Frequently the actual pressure may be as low as 40 mm. The fluid is rarely yellow and as rarely it contains clots spontaneously. Cells are not usually increased, but in meningeal sarcomatosis, or widespread meningeal tumours there may be a slight pleocytosis. Protein is always raised in complete block and figures as high as 1,000 mg./100 ml. are to be seen. There is practically always some increase in protein in any spinal tumour whether there is a block or not.

The examination of the cisternal fluid is of value as the protein in this fluid will be normal in many cases of spinal tumour. However, as there may be an increased protein in the fluid just above a tumour, a cervical tumour may give rise to an increased cisternal fluid protein, but even so there is always a wide disparity between the protein levels in the two regions. Similarly, a raised protein level in the fluid from the lumbar region may be found above a herniated lumbar disc or a lower lumbar chondroma.

**Disseminated Sclerosis**

Characteristic changes can be found in the fluid in this condition. The acute phases give rise to a slight to moderate pleocytosis in which the predominating cell is the lymphocyte. In some 5 per cent. of cases polymorphs in small numbers are present. The protein content is only very slightly raised above the normal and, in fact, almost never exceeds 100 mg./100 ml. Globulin is always in excess and an abnormal Lange curve is very frequently seen. The paretic curve in association with a raised cell count, slight rise in protein and a negative Wassermann is seen but only occurs in some 15 per cent. of all cases, whereas the luetic type of curve is seen in some 30 to 40 per cent. of all cases, while some 40 per cent. of all cases show no appreciable change in the Lange.

The C.S.F. in neuromyelitis optica may show changes similar to those found in disseminated...
sclerosis, but on the whole cells and protein tend to be higher provided the lumbar puncture is performed in the acute stages.

There are many other diseases in which the C.S.F. is examined but in which there are few, if any, diagnostic features. The one obvious exception to this is in subarachnoid haemorrhage where a fluid intimately mixed with blood is found. Should the haemorrhage be three or more days old the supernatant fluid, after centrifugation, may be pale yellow and after one week the red cells diminish or may be entirely lost, but the yellow colouration may persist along with a slightly raised protein content.

The description given here of the results of the examination of the cerebrospinal fluid has of necessity an air of dogmatism about it, but this is inevitable unless statistical tables and long descriptions of each condition are given. Therefore, one must be prepared for slight variations from the absolutely typical findings recorded here, but on the whole these are relatively rare, so that the writer may perhaps be forgiven for the approach that has been used.

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