Although over the years the considerable breakthroughs made in our understanding of diabetic neuropathy has come from both neurologists and diabetologists, in recent times (in the UK) it is diabetologists that have assumed the main clinical role in diagnosing and managing the most common neuropathy in the western world. It may therefore be that referral of patients with diabetic neuropathy from the diabetic clinic only occurs if there is a neurologist with a particular interest or if help is required regarding a specific issue such as an atypical clinical picture or pain control. This creates a risk that those training in neurology end up with a somewhat skewed view of the spectrum of diabetic neuropathies. The other situation where neurologists come across diabetic neuropathy is when the patient is found to have diabetes in the course of investigating a peripheral neuropathy.

**DIAGNOSIS OF DIABETES MELLITUS**

Diagnosis of diabetes mellitus (DM) is made on a combination of typical symptoms—weight loss, thirst, weakness and fatigue—with a persistently raised blood glucose (table 1). Glycosuria and raised HbA1 values alone are not used to make the diagnosis of diabetes mellitus.

**CLASSIFICATION OF THE DIABETIC NEUROPATHIES**

Based on a modification of the classification proposed by PK Thomas, a number of distinct syndromes are identifiable (table 2).

**Hyperglycaemic neuropathy**

Tingling paraesthesia, pain or hyperaesthesia in the feet have long been described in patients with newly diagnosed DM or those with very poor glycaemic control, this being the phenomenon of hyperglycaemic neuropathy. The symptoms, and slowing of nerve conduction, are rapidly reversed by improving glucose control.

**Diabetic symmetric distal polyneuropathy with autonomic neuropathy**

This is the most common diabetic neuropathy and it is characterised by a length related distal distribution of sensory and motor symptoms and signs. As autonomic involvement occurs in many patients with diabetic symmetric distal polyneuropathy (DSDP), and forms an important part of the clinical complex, it is best that both are considered together.

It is highly likely that by the time DSDP is diagnosed, the patient with either type 1 or type 2 diabetes will have had a prolonged period (sometimes over years) of abnormal glucose metabolism. This is particularly the case in those with type 2 DM who tend to be “discovered” to be diabetic on presentation with symptoms and signs of a neuropathy. It is also worth remembering that it is usual to find evidence of either retinopathy or nephropathy in any case of DSDP—whether newly diagnosed or long established.

Some patients with DSDP have no symptoms, but the most common complaint is of tingling, buzzing or prickling sensations affecting the feet, which may also feel tight or hot or cold. The symptoms are often, but not exclusively, symmetric in distribution. The patient may complain of numbness or “as if my feet are wrapped up in cotton wool”. The pains in the feet are often worse at night—although this phenomenon is not unique to DSDP.

**Clinical signs**

The cardinal sign is absent ankle reflexes (table 3). Without this, it is difficult to make a diagnosis of DSDP. Loss of knee reflexes occurs in about two thirds of cases, but loss of any upper limb reflexes occurs in only a quarter of patients with DSDP. Muscle weakness is usually mild and confined to the feet, mainly in the distribution of the common peroneal nerve and more obviously affecting extensor hallucis longus and extensor digitorum brevis compared with dorsiflexion and eversion. Proximal leg weakness can be seen, but this, together with the presence of significant upper limb weakness, should make one suspicious of an alternative diagnosis and only after appropriate investigations are negative can a diagnosis of DSDP be made in these cases.
Sensory disturbance is very common. Vibration sense at the toes is most frequently affected. Pin prick, temperature, and light touch sensations are lost in a sock or stocking distribution, and if there is upper limb sensory loss in a glove distribution, the level of impairment in the legs has to have reached mid thigh. If not, look for another explanation for the distal neuropathy. Based on the relative loss of sensory modalities, one can arbitrarily divide the neuropathy into “large fibre type” (predominant loss of vibration, light touch, and joint position senses) and “small fibre type” (predominant loss of pain and temperature), but these subgroups are uncommon and represent both ends of the continuum of DSDP.

In more severe cases, sensory loss can extend to involve the trunk, affecting initially the anterior chest/abdominal wall in a “breastplate” distribution, and may extend laterally around the trunk.

Clinically significant symptomatic autonomic neuropathy is relatively uncommon, but specific autonomic function tests are said to show an abnormality in 97% of patients with DSDP. If there is a prominent autonomic neuropathy but only a very mild DSDP or no distal polyneuropathy in a patient with diabetes, think of another cause for autonomic disturbance.

What should be considered in a differential diagnosis? Although diabetes is a common condition, there is the possibility that there may be another cause for the distal neuropathy. Good history taking (alcohol, family history of neuropathy, drug history, etc) and a few basic blood tests (Table 4) should be enough to secure the diagnosis of DSDP.

What atypical features might suggest an alternative or additional neuropathy?

1. Severe autonomic neuropathy with mild DSDP: One has to consider amyloid neuropathy which can produce a small fibre neuropathy with spontaneous pains, but where autonomic features are prominent. The clinical features of familial and non-familial amyloidosis will be very similar.
2. Rapidly progressive motor component: When there is prominent weakness, the possibility of a superimposed chronic inflammatory demyelinating polyneuropathy (CIDP) has to be considered. There is some evidence that CIDP may be more common in patients with diabetes than in the general population. Nerve conduction studies may be helpful. CSF protein will be elevated in both CIDP and DSDP, but the presence of oligoclonal bands would point to CIDP.

Painful DSDP variants

Although it is not clear whether either of the two syndromes outlined below are distinct entities or simply part of the spectrum of painful DSDP, it is of value to be aware of their existence.

Insulin neuritis

A severe painful sensory neuropathy can be seen in relation to tightening of glucose control. This can be seen when trying to improve glucose control in a patient already established on insulin, or in patients being started on insulin for the first time. The pain is often difficult to control, and as with other painful diabetic neuropathy syndromes, is worse at night. Clinical signs are not uncommonly minimal or even absent. Nerve conduction studies may be normal.

Over a period of up to 12 months, recovery is usual.

Acute painful neuropathy with severe weight loss

Originally termed “diabetic neuropathic cachexia”, this uncommon condition occurs predominantly in male patients with poorly controlled type 1 diabetes. Females who try to control their diabetes by not eating and become anorexic also develop a similar painful neuropathy. Pronounced weight loss is the key feature. The distal lower limb pain is severe, with burning and tightness of the feet. Depression is not uncommon.

The weight loss is dramatic, and when body weight stabilises—usually after a period of improved glucose control with insulin—the pain begins to abate. It can take up to 12 months for improvement to occur and a further 1–2 years for recovery of body weight back to normal and for the pain to subside completely.

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**Table 1** Biochemical diagnosis of diabetes mellitus

<table>
<thead>
<tr>
<th>Test</th>
<th>Normal Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Random plasma glucose</td>
<td>$\geq 11.1$ mmol/l, or</td>
</tr>
<tr>
<td>Fasting plasma glucose</td>
<td>$\geq 7.8$ mmol/l, or</td>
</tr>
<tr>
<td>2 hour plasma glucose</td>
<td>$\geq 11.1$ mmol/l</td>
</tr>
</tbody>
</table>

*Glucose tolerance test using 75 g glucose load.

**Table 2** Classification of the diabetic neuropathies

- Generalised neuropathy
  - Hyperglycaemic neuropathy
  - Symmetric distal polyneuropathy with/without autonomic neuropathy
  - Acute painful sensory neuropathy variants
- Focal and multifocal neuropathy
  - Cranial neuropathies
  - Focal limb neuropathies
  - Thoracolumbar radiculoneuropathy
  - Lumbar sacral radiculoplexus neuropathy (Bruns-Garland syndrome)

**Table 3** Clinical assessment in diabetic symmetric distal polyneuropathy (DSDP)

- Look for any deformity, callus or foot ulcer, infection or fissure
- Absent ankle reflexes (mandatory)
- Test all sensory modalities starting at the toes and fingertips (in many diabetic screening clinics, vibration sensation and neuromaturation sensitivity are the only modalities tested and this seems to be an effective part of a neuropathy screen)
- Weakness of small foot muscles (extensor hallucis longus and extensor digitorum brevis)
- Check resting pulse and blood pressure lying and standing
- Check peripheral pulses
- Look for retinopathy
- Check urine for protein

**Table 4** Blood tests in a patient with DSDP

- B12 and folate values
- Serum protein electrophoresis
- ANA/ENA/rheumatoid factor
- Thyroid function
- FBC/ESR, U&Es and LFTs

ANA, antinuclear antibody; ENA, extractable nuclear antigen; ESR, erythrocyte sedimentation rate; FBC, full blood count; IFT, liver function test; U&E, urea and electrolytes

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*## NEUROLOGY IN PRACTICE##

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with DM who has developed symptoms and signs of a distal polyneuropathy associated with retinopathy and/or nephropathy. In a patient with diabetes, being male and tall. It is usually, if not invariably, the most important aspect of treatment. This has been clearly demonstrated in type 1 DM where tight glycaemic control reduces the risk of developing DSDP by 69% at five years. The same has not yet been shown for type 2 DM, but one might suspect similar outcomes. Once established, DSDP is irreversible and slowly progressive. At this stage, strict glucose control provides no clinically significant improvement from the patient’s perspective, despite modest improvement in vibration threshold and nerve conduction studies.

From the research studies that identified metabolic abnormalities, a variety of potential treatments have been investigated—for example, aldose reductase inhibitors, myo-inositol supplementation, α-lipoic acid, and administration of nerve growth factor—but none have had sufficient impact on DSDP to be approved as a specific long term treatment. Pancreatic transplantation is of real benefit only in those who have advanced renal failure and could undergo a combined kidney and pancreas transplant, despite studies that have shown that pancreatic transplantation can halt the progression of DSDP.

The treatment of pain associated with DSDP has received considerable attention over the last decade, but this still poses one of the most difficult aspects of management, as there is no single effective treatment. The list of approved drugs is increasing but often based on evidence from short term trials—for example, intravenous α-lipoic acid, which has been granted a licence for use in painful DSDP in Germany, but not in the UK. With such a variety of drugs available the best way to approach the treatment of pain in DSDP is to have a sequence of preferred choice drugs (often dictated by personal experience), and where possible to stick to monotherapy (table 7). Although the use of opiates is often frowned upon, they do have a role where antidepressants and anti-convulsants have failed or provided only modest pain control. Tramadol (up to 400 mg/day) is the best opiate to start with, and in certain circumstances morphine may be required.

**FOCAL AND MULTIFOCAL NEUROPATHIES**

This group of neuropathies should not be considered in isolation as they often co-exist with DSDP. Their aetiologies are likely to differ, however, with the focal and multifocal neuropathies having a prominent vascular component or a compressive aetiology as opposed to the predominant metabolic component in DSDP.
Diabetic truncal radiculoneuropathy

This is a phenomenon seen in both type 1 and type 2 diabetics, usually in middle or old age. The cardinal features are outlined in table 8. The diagnosis is made on clinical grounds, but evidence of denervation can be found in abdominal wall or intercostal or paraspinal muscles in the region where the pain and/or sensory loss has occurred. The abrupt onset and spontaneous recovery suggests a vascular aetiology, but an “inflammatory” aetiology cannot be excluded as it can coexist with diabetic lumbosacral radiculoplexus neuropathy (see below).

Management involves controlling the pain; topical capsaicin is useful if the affected area is small and well localised. Full recovery over a period of months is usual.

Diabetic lumbosacral radiculoplexus neuropathy (Bruns-Garland syndrome)

Previously termed diabetic amyotrophy, this clinical entity has had a variety of other names. The most recent attempt to define the condition is somewhat of a mouthful for everyday clinical use, and it may be simpler to use the eponym “Bruns-Garland syndrome” (after Bruns who initially described the syndrome in 1890, and Garland who rediscovered it and coined the term “amyotrophy”). It is most common in older patients with type 2 DM and is rarely encountered in those with type 1 DM.

The clinical features are outlined in table 9. The evolution of symptoms can be quite variable, and may progress to generalised lower limb paresis (“diabetic paraplegia”). A clinically indistinguishable syndrome occurs in patients without diabetes.

The main differential diagnosis to consider is that of an infiltrative pelvic malignancy, particularly when there is profound weight loss and unilateral weakness. A similar picture can be seen as a complication of radiotherapy. If there is no pain associated with progressive asymmetric leg weakness, the diagnosis is more likely to be CIDP.

Cranial neuropathies

The sixth nerve palsy associated with diabetes occurs in older patients and does not pose a great diagnostic dilemma. Onset is abrupt and usually painless. Full recovery occurs in the vast majority over 3–5 months. Detailed investigations are rarely required.

A third nerve palsy is not as common as a sixth nerve palsy, and again tends to be seen in patients over the age of 50 years, most of whom have evidence of a DSDP. In 50% of cases the onset is associated with retro-orbital pain, which may persist for several days. The key feature is that the palsy spares the pupil, possibly because the pupillomotor fibres of the IIIrd nerve are located on the outer layers of the fascicle, and ischaemia tends to affect the centre of the fascicle.

Although it is only rarely that a pupil sparing IIIrd nerve palsy arises from a posterior communicating artery aneurysm, as this possibility exists, imaging (magnetic resonance/computed tomographic angiography) is recommended. In diabetic IIIrd nerve palsy imaging is usually normal, although it has been postulated that small mesencephalic infarcts may be an alternative explanation. No specific treatment is required other than prismatic help with diplopia as the ptosis recovers. Complete resolution normally occurs over 3–6 months.

Limb mononeuropathies

Diabetic nerves seem to be more susceptible to compression injury, although exactly why this is the case is not well understood. The treatment of entrapment neuropathy (carpal tunnel syndrome, ulnar neuropathy or common peroneal neuropathy) should follow the same management guidelines as in a non-diabetic patient (see Fuller, p ii20). If the nerve compression is symptomatically troublesome, and especially if there is associated muscle weakness, decompression is needed. The results of decompressive surgery are not as good as in non-diabetic patients.

Table 7 Treatment of painful DSDP

<table>
<thead>
<tr>
<th>A. For localised pain (for example, dorsum of foot)</th>
<th>Capsaicin cream (0.075%) applied four times/day (maximum 6–8 weeks)</th>
</tr>
</thead>
<tbody>
<tr>
<td>B. For generalised distal pain</td>
<td>Gabapentin 600–3600, Carbamazepine 100–800, Amitriptyline 10–150, Nortriptyline 10–150, Imipramine 10–150, Desipramine 10–150</td>
</tr>
<tr>
<td>C. For those whose pain is resistant to the above drugs consider:</td>
<td>Mexiletine, SSRI (citalopram or paroxetine), Other anticonvulsants (lamotrigine, topiramate), Opioids: tramadol, pethidine/morphine, Spinal cord stimulator</td>
</tr>
<tr>
<td>D. Alternative</td>
<td>Opsite dressing (applied to painful areas), Acupuncture, Pain psychologist/counsellor</td>
</tr>
</tbody>
</table>

Table 8 Diabetic truncal radiculoneuropathy: clinical features

- Pain over a focal area on the chest and/or abdomen. Usually unilateral, the pain often burning in quality, and in a variety of distributions reflecting nerve root or intercostal nerve trunk involvement
- Focal contact hyperesthesiaeae in the same area as the pain
- Focal anterior abdominal wall weakness may be evident
- Associated weight loss in some cases may be profound
- Good prognosis with spontaneous recovery over several months

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- Focal contact hyperesthesiaeae in the same area as the pain
- Focal anterior abdominal wall weakness may be evident
- Associated weight loss in some cases may be profound
- Good prognosis with spontaneous recovery over several months
Suggested investigations are outlined (table 9). Neurophysiology is helpful, but CSF examination and nerve biopsy should only be considered if an alternative diagnosis is considered. Examination of nerve biopsies has shown evidence of microvasculitis and endoneurial inflammatory infiltration but these findings do not influence management. It is uncertain whether these inflammatory changes are primary or secondary phenomena to possible ischaemic injury.

There does appear to be a rarer brachial radiculoplexus neuropathy, reported in association with the Bruns-Garland syndrome in diabetic patients.

Treatment is centred initially around pain control (table 8). This can be difficult and often requires opiates. Physiotherapy and orthotic assessments are helpful in selected, often more severely affected cases. Although the suspected pathogenic mechanism is inflammatory change in the nerve or epineurial vessels, it is not established whether immunosuppression (steroids or intravenous immunoglobulin) has any role to play in the treatment of a syndrome, which eventually spontaneously recovers, albeit that this may be incomplete in a number of patients.

**Table 9  Diabetic lumbosacral radiculoplexus neuropathy (Bruns-Garland syndrome): clinical features and investigations**

<table>
<thead>
<tr>
<th>Clinical features</th>
<th>Investigations</th>
</tr>
</thead>
<tbody>
<tr>
<td>Males more frequently affected than females</td>
<td>Denervation changes in paraspinal, proximal and distal leg muscles</td>
</tr>
<tr>
<td>Pain: Severe, affecting lower back, buttocks or anterior thighs, burning and aching in quality; worse at night</td>
<td>There is often an associated DSDP</td>
</tr>
<tr>
<td>Weakness: Follows pain within a matter of a few days to several weeks and usually unilateral at onset. Later may be bilateral but asymmetric. Mainly proximal, but not uncommon for distal muscles to be involved</td>
<td>Lower limb f wave latencies may be more prolonged than usually seen in DSDP</td>
</tr>
<tr>
<td>Weight loss: May be dramatic (&gt;10–20 kg)</td>
<td>Only helps if the clinical picture is unusual</td>
</tr>
<tr>
<td>Prognosis is reasonable: recovery is heralded by stabilisation of body weight and resolution of pain. Muscle strength improves slowly over many months, but a number of patients never regain normal lower limb strength</td>
<td>Rarely helps management</td>
</tr>
</tbody>
</table>

Microvasculitis and endoneurial mononuclear cell infiltration found in the intermediate cutaneous nerve of the thigh and also in the sural nerve

**COMMENT**

The diabetic neuropathies are common and remain a major source of morbidity. Optimal treatment at this time requires good control of blood sugar, managing symptoms, and fastidious attention to foot care. The disappointing lack of benefit of other treatments may be related to the fact that most were trialled in patients with established neuropathy. It is probable that future pharmacological treatments for the more common DSDP will have to be directed at early neuropathy. This may provide the impetus to rekindle the neurologist’s interest and involvement in the diagnosis and management of diabetic neuropathy.

**KEY REFERENCES**