

In this issue, Meditech extends its congratulations to Dr. Marie-Josée Robichaud and Dr. Pierre Levesque in Moncton, New Brunswick on the establishment of their new Laser Therapy Clinic, now in the final process of construction. What they have achieved is extraordinary and serves as an example to those who care and “those who dare”.

In 2009, we hosted four one-day Advanced Training Seminars on the management of back problems and wound healing. These have been highly successful and will continue throughout 2010. We plan to have some webinars beginning in February. The subjects to be covered include excerpts from our advanced training sessions, the

introduction of a new referral kit and other matters of interest.

For 2010 and the years to come, our efforts will continue to focus on extending the horizons of Laser Medicine.



Example of a new BioFlex Laser Therapy Clinic under construction in Moncton, NB. Proprietors Drs. Marie-Josée Robichaud and Pierre Levesque

## NEUROPATHIES AND THE APPLICATION OF LASER THERAPY

### DEFINITION

*Peripheral neuropathy is defined as damage to the peripheral nervous system resulting in a syndrome of sensory loss, muscle weakness and atrophy, along with vasomotor symptoms, alone or in any combination*<sup>1</sup>.

The basic peripheral nervous system components consist of a cell body located in either the anterior horn (motor) or the dorsal root ganglia (sensory) of the spinal cord and a long extension (axon) covered in a chain-like series of cells known as Schwann cells; these produce myelinated nerve fibers<sup>2</sup> (see Figure 1). The thin microscopic individual nerve fibers are surrounded by connective tissue referred to as the endoneurium. Bundled nerve fibers (fascicles) are then surrounded by a strong connective tissue known as perineurium. Finally, loose connective tissue called epineurium surrounds groups of bundles or fascicles. Thirty-one paired spinal and ten cranial nerves form the basis of the more numerous peripheral nerve trunks and their terminal branches.

Peripheral nerves are highways for the conduction of nerve impulses to and from the spinal cord. The larger the diameter of the nerve axon (i.e. width of the highway), the faster the impulses can be conveyed. For example, the sensations of light touch and vibration move quite rapidly along very large axons, while pain and temperature sensation move more slowly along smaller axons in their course towards the spinal cord;

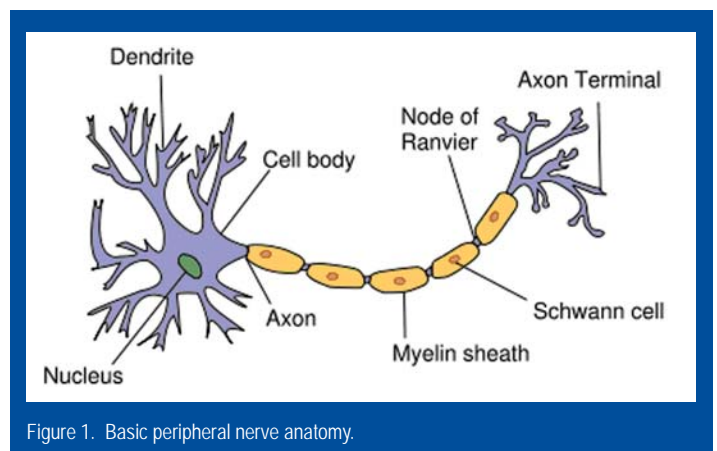


Figure 1. Basic peripheral nerve anatomy.

motor nerves have large diameter axons that rapidly convey impulses away from the spinal cord to the skeletal muscles.

### CLASSIFICATION

According to the National Institute of Neurological Disorders and Stroke, there exist over one hundred different types of identified peripheral neuropathies. The causes include toxic and metabolic factors, infectious diseases, mechanical compression, inflammation, ischemia and paraneoplastic lesions.

In developed countries, diabetes and alcoholism are the most frequent causes of peripheral neuropathy<sup>3</sup>. The common incidence of these conditions is often found to be time consuming from the diagnostic perspective, can be therapeutically frustrating and economically burdensome.

Generally speaking, the various types of neuropathy can be classified into three basic patterns of distribution:

- single peripheral nerve (focal mononeuropathy)
- multiple single peripheral nerves (multifocal mononeuropathy)
- peripheral nerves simultaneously (symmetrical polyneuropathy)

In addition to the three basic categories listed above, more specialized sub-patterns of peripheral neuropathy can include both an affected nerve root (radiculopathy) or the nerve plexus (plexopathy).

When classifying peripheral neuropathies, the time, course and presenting fiber deficit (sensory, motor or mixed) along with the pattern of distribution are important factors that must be considered. For example, peripheral neuropathies may present as an acute (<3 wks) purely motor neuropathy (e.g. Guillain-Barre Syndrome) or as chronic (>8-12 wks) symmetrical purely sensory polyneuropathy (e.g. Diabetes Mellitus). In addition, there are mixed sensory and motor neuropathies, which may or may not follow a specific dermatomal pattern (e.g. Carpal Tunnel, Disc Herniation etc.). In the above listed examples, both large and small fiber diameter axons are equally affected. In rare cases small fiber neuropathy presents exclusively, creating distal impairment of pain and temperature sensations only.

### CLINICAL MANIFESTATION AND PATHOPHYSIOLOGY

Clinical signs and symptoms of peripheral neuropathy depend primarily on the etiological factors involved. In this review, compressive neuropathies will be the primary focus. Compression of a normal healthy nerve results in paraesthesias and muscle weakness, generally without pain. On the other hand, compression of an inflamed nerve can cause pain, in addition to the objective neurological findings<sup>4</sup>. Compressive neuropathies frequently occur as the nerve passes through tight pathways or tunnels, formed by the encasing tissue borders. Many recognized examples of compression include the median nerve at the wrist, the ulnar nerve at the wrist or elbow but more commonly spinal nerve roots traversing the intervertebral foramen. Nerve roots are more susceptible to compression, compared to peripheral nerves, due to their less developed vascular network and lack of epineurium (outer connective tissue covering)<sup>5</sup>.

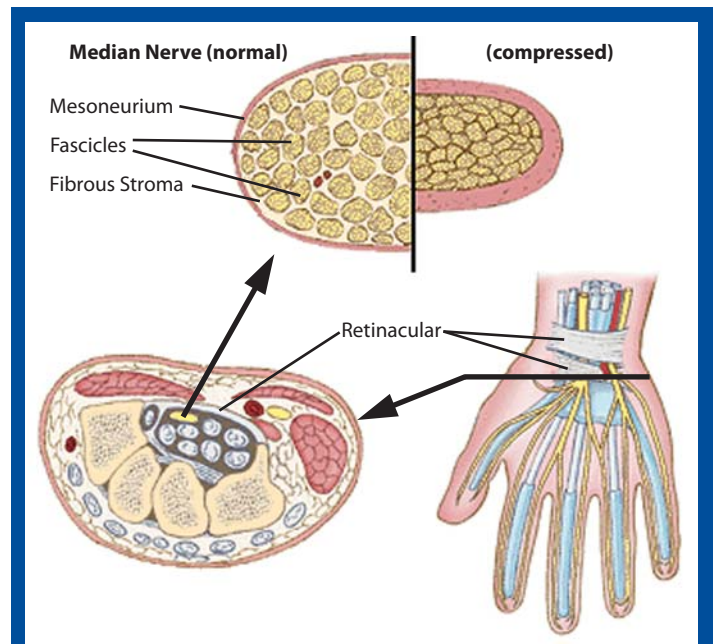


Figure 2. Pathology of Carpal Tunnel Syndrome

Compressive neuropathies involve nerve dysfunction secondary to localized interference of microvascular function and structural changes in the nerve or adjacent tissues<sup>6</sup>. As nerve tissue is compressed, pressure gradients develop, forcing tissues into areas of lower pressure. Early symptoms are primarily due to transient changes to microcirculation associated with edema and often result in morphological changes including segmental demyelination (chronic impairment of microcirculation)<sup>6</sup>. If nerve compression persists, distal axonal degeneration of nerve fibers occurs, manifesting as persistent paraesthesias (numbness/tingling) muscle weakness and atrophy. Finally, nerve compression appears to be both rate and pressure dependent, with acute duration resulting in only brief vascular interruption, while chronic sustained pressure results in both microvascular edema and permanent structural damage<sup>6</sup>.

### FOCAL MONONEUROPATHY (CARPAL TUNNEL)

Carpal tunnel is primarily a compressive neuropathy. Patients may initially present with intermittent numbness and tingling in the digits, occurring primarily at night. The pain and tingling may radiate proximally and numbness generally affects the digits according to the median nerve distribution<sup>7</sup>. If compression persists or progresses, more severe symptoms may occur. These include paraesthesias and numbness which can result in muscle weakness and atrophy (see Figure 2).

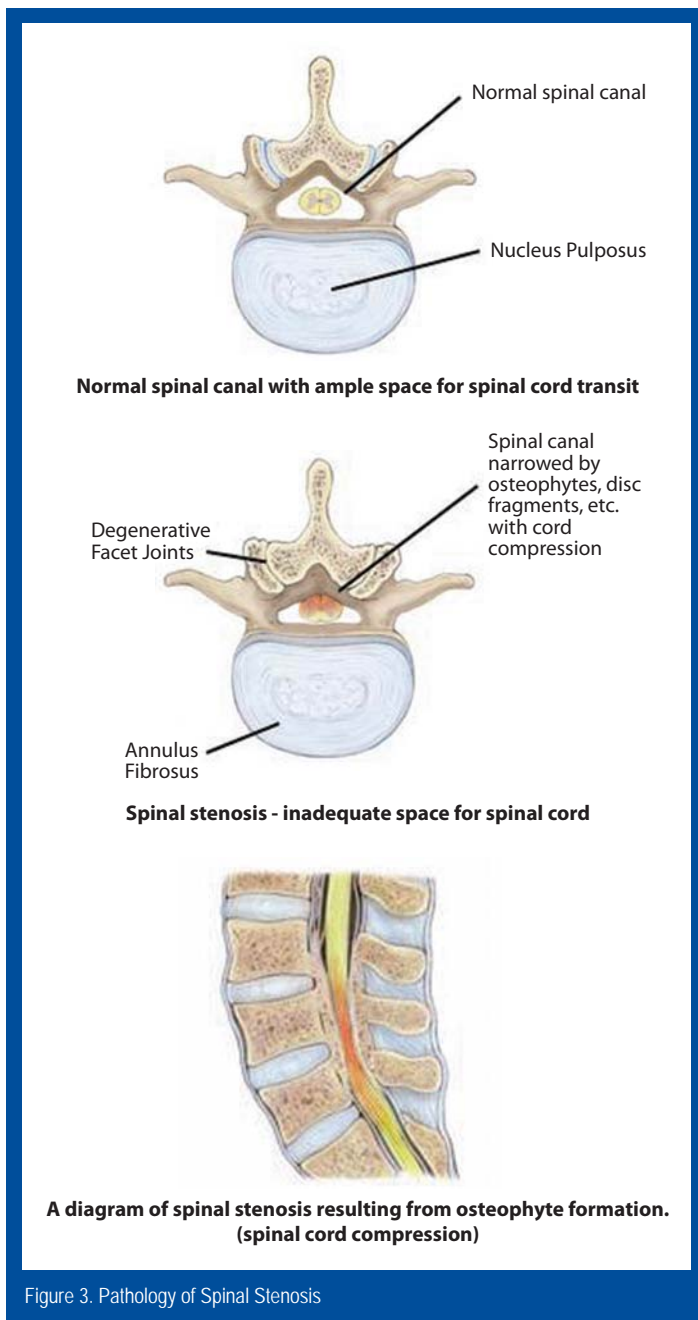


Figure 3. Pathology of Spinal Stenosis

### ***RADICULOPATHY (NERVE ROOT COMPRESSION)***

Radiculopathy is generally caused by compression of the nerve root(s), as demonstrated in degenerative disc disease, disc herniation, spinal stenosis etc.; this can result in paraesthesias and pain along the dermatomal distribution of the affected nerve root (see Figure 3). For example, a lumbar disc herniation affecting the L5 nerve root will cause pain over the lateral thigh and dorsum of the foot with potential loss of sensation, specifically in the interspace between the first two toes and eventually a decrease in the power of dorsi-flexion of the foot. Compression of a deep branch of the peroneal nerve

(mononeuropathy) at the inferior aspect of the extensor retinaculum causes a somewhat similar loss of sensation as an L5 radiculopathy but no loss in the power of dorsi-flexion of the foot.

Lumbar spinal stenosis (LSS) often causes the syndrome of activity related thigh and leg pain (uni- or bilateral) associated with low back pain. Bilateral radicular symptoms, such as numbness and weakness in the legs and feet occur with activity (i.e. walking) and are often relieved by rest (i.e. sitting, leaning forward). The compressive symptoms of LSS cause neural ischemia and neurogenic claudication.

### **TREATMENT**

Non-conservative treatment (surgical decompression) to remove the mechanical extraneural pressure has been shown to only be moderately successful in severe cases. For example, a recent systematic review found surgery for spinal stenosis to be associated with only short-term benefit compared to nonsurgical therapy, with benefits diminishing over long-term follow-up in some trials<sup>8</sup>. It has been demonstrated conclusively that neurological symptoms can persist even after the mechanical pressure has been removed<sup>9</sup>. Further, documented damage to nerves follows a dose dependent response sufficient to cause axonal degeneration, that may not be reversed by simply relieving the mechanical pressure. As compressive neuropathies span a continuum from the early phase of impaired blood flow and inflammation to eventual axonal degeneration, a treatment method targeting both the inflammatory process and regeneration is desirable from a result based perspective.

Low intensity laser therapy (LILT) has demonstrated the ability to stimulate the healing process and reduce the pain associated with peripheral nerve damage. At the cellular level, LILT has been shown to improve Schwann cell proliferation and reduce scar tissue formation in addition to reversing the process of progressive nerve degeneration post injury. Furthermore, a positive influence on axonal growth and re-myelination has been noted<sup>10</sup>.

A double-blind randomized study found an almost 70% vs. 18.2% improvement, in positive somatosensory evoked

responses and better quality (larger diameter axons) of the nerve regeneration process, with low intensity laser therapy after complete surgical transection and direct anastomosis of the sciatic nerve<sup>11</sup>. Clinically, a study by Iijima et al. (1991) using low intensity laser therapy, reduced pain levels by 45% in 18 patients with severe post herpetic neuralgia. A clinical double-blind, placebo-controlled randomized study compared the effectiveness of LILT on patients who had been suffering (6 months to several years) from incomplete peripheral nerve and brachial plexus injuries. This study revealed that LILT significantly improved motor function in addition to recruitment of voluntary muscle activity in the partially paralyzed limbs of brachial plexus injuries, compared to the placebo group, which failed to show any improvement.

More specifically, a randomized controlled trial carried out at General Motors Company found that carpal tunnel patients treated with low intensity laser had better functional recovery and a higher back to work percentage (72% active laser vs. 41% sham)<sup>12</sup>. Similarly, a more recent study showed that low intensity laser therapy improved both the sensory and distal motor latencies of the median nerve in carpal tunnel patients, compared to controls<sup>13</sup>. By safely and effectively healing damaged nerve tissue, restoring function along with the alleviation of pain, low intensity laser has been found to be a key therapeutic component in the treatment of peripheral nerve injuries.

#### ***TREATMENT OF FOCAL MONONEUROPATHY***

At Meditech focal mononeuropathy patients are initially assessed both clinically and utilizing Neurometrix™ nerve conduction testing. Patients with established neuropathy diagnoses are treated with low intensity laser therapy locally over the area of defined pathology and in some instances segmentally over the relevant nerve roots. For example, carpal tunnel patients are subjected to direct stimulation locally at the wrist to repair damaged nerve tissue, while segmental stimulation of nerve roots (cervical spine) stimulates axonal transport and circulation.

#### ***TREATMENT OF RADICULOPATHY***

Similarly, lumbar disc herniation and spinal stenosis patients are assessed initially and repeatedly as treatment progresses. Initial treatment of lumbar spinal stenosis for

example, may involve low intensity laser therapy treatment of the lumbar nerve roots only. Irradiation of the spinal nerve root(s) will promote neural and soft-tissue healing in addition to providing an analgesic effect. Treatment plans are outlined involving early symptom management, followed by progressive nerve repair and regeneration. Early pain relief is best obtained following four to five treatments in succession and subsequent appropriately spaced treatments until the objective of complete neurological recovery is reached.

#### **CONCLUSIONS**

- Neuropathies encompass an extensive number of medical conditions which may be simple or complex in nature.
- The key factor in the successful management of these lesions often requires individually designed therapy programmes, delivered both locally and over the dermatome innervating the area of involvement.
- Monitoring of the clinical response requires intense focus and serves as the basis for protocol adjustments.
- Laser Therapy is a non-invasive, non-toxic therapeutic approach which has consistently produced superior results in the treatment of these entities.
- The confirmation of nerve regeneration by Neurometrix and other studies, has provided evidence of the return to normal function.
- If on occasion a specific diagnosis cannot be established, this does not preclude the utilization of Laser Therapy.
- Despite the complex etiologies of peripheral neuropathies and sometimes difficulty in establishing the correct diagnosis, Laser Therapy has been found to be highly effective in the resolution of these conditions.
- With the application of Laser Therapy, there is nothing to be lost and a great deal to be gained.

**REFERENCES AVAILABLE UPON REQUEST**