MRI of Neuropathic Pain: Spectrum of Imaging Findings

Poster No.: C-2152
Congress: ECR 2013
Type: Educational Exhibit
Authors: E. Blanco Pérez¹, F. Mata Escolano¹, E. Cascón Sánchez¹, A. González-Cruz Soler¹, J. C. quiles teodoro¹, V. MARTINEZ SANJUAN², ¹Valencia/ES, ²PICASSENT-VALENCIA/ES
Keywords: Metabolic disorders, Ischaemia / Infarction, Inflammation, Diagnostic procedure, MR, Neuroradiology peripheral nerve, Neuroradiology brain
DOI: 10.1594/ecr2013/C-2152

Any information contained in this pdf file is automatically generated from digital material submitted to EPOS by third parties in the form of scientific presentations. References to any names, marks, products, or services of third parties or hypertext links to third-party sites or information are provided solely as a convenience to you and do not in any way constitute or imply ECR's endorsement, sponsorship or recommendation of the third party, information, product or service. ECR is not responsible for the content of these pages and does not make any representations regarding the content or accuracy of material in this file. As per copyright regulations, any unauthorised use of the material or parts thereof as well as commercial reproduction or multiple distribution by any traditional or electronically based reproduction/publication method ist strictly prohibited. You agree to defend, indemnify, and hold ECR harmless from and against any and all claims, damages, costs, and expenses, including attorneys' fees, arising from or related to your use of these pages. Please note: Links to movies, ppt slide shows and any other multimedia files are not available in the pdf version of presentations. www.myESR.org
Learning objectives

To know the etiology of neuropathic pain (NP) and the utility of MRI for differential diagnosis. To provide spectrum of imaging features of these entities. To analyze the future directions of the role of new MRI techniques for characterization if the NP (functional MRI, Diffusion tensor MRI tractography).
Background

It was defined by IASP (International Association for the Study of Pain) in 1994 as a consequence of injury or disease affecting the somatosensory system. So NP refers to pain that originates from injury or disfunction of the nervous system, including diseases of the peripheral or central nervous system.

The patient with neuropathic pain may have concomitant non-neuropathic pain and it should be consider that NP may be the first manifestation of a systemic disease.

A wide variety of abnormalities can cause NP, i.e., metabolic disease, infection, ischemia, injury, entrapment, connective tissue disease, acquired immunodeficiency, malignancy, drugs, and toxins. Although NP may develop without any identifiable cause (intercostal neuralgia, idiopathic polyneuropathy, etc).

We can classify neuropathic pain syndromes in mononeuropathies (post-traumatic neuroma, nerve or radicular compression and idiopathic causes), polyneuropathies (ischemic, metabolic), deafferentation (postherpetic neuralgia, amputation neuromas and phantom limb syndrome) and Complex Regional Pain Syndrome. Central nervous system diseases includes thalamic stroke, Wernicke's encephalopathy, multiple sclerosis, HIV encephalopathy and granulomatous diseases.
Imaging findings OR Procedure details

Fields in 1991 proposed a classification of NP, into four groups.

- Mononeuropathy: post-traumatic neuromas, nerve root compression and idiopathic neuropathies as trigeminal neuralgia.

- Polyneuropathies: ischemic, metabolic (diabetic neuropathy), etc.

- Deafferentation pain: postherpetic neuralgia, amputation neuromas phantom limb pain syndrome and central pain syndrome.

- Complex Regional Pain Syndrome type1 (Reflex Sympathetic Dystrophy -RSD-) and type 2 (Causalgia).

Another classification of NP is based on the location of the lesion:

- Central Nervous System:
  - Ischemic etiology: Central poststroke pain due to thalamic infarct (image 1)
  - Metabolic cause: Wernicke (image 2)
  - Immunological diseases: Multiple Sclerosis (image 3), AIDS.
  - Granulomatosus diseases: sarcoidosis, tuberculosis.
  - Intrinsic and extrinsic compression: tumors, meningiomas.

- Cranial nerves:
  - Neoplastic disease: benign and malignant (image 5)
  - Compression neuropathy (image 6)

- Peripheral nervous system:
  - Nerve tumors: Morton's neuromas (image 7), schwannomas (image 8), traumatic neuromas, etc
  - Nerve root compression: herniated disc.
  - Plexopathies (image 9).
  - Systemic disease.
FUNCTIONAL IMAGING STUDIES

Application of new MRI techniques will play a very important role in the study of patients with neuropathic pain.

These imaging procedures include functional magnetic resonance (fMRI) performed with a blood oxygenation level-dependent (BOLD) sensitive T2*-weighted multislice gradient echo EPI sequence, to establish the extent of cortical reorganization from face to hand area in motor and somatosensory representational maps.

Generally, fMRI studies are based on the acquisition of images, while the patient is at rest and while performing a specific cognitive, emotional or sensitivo-motora task. In this procedure the resonance signal changes depend on the concentrations of oxygen in blood, fMRI is able to detect this change is due to a fundamental difference in the paramagnetic properties of oxyHb and deoxyHb.

When a specific region of the cortex increases its activity in response to a task, the extraction fraction of oxygen from the local capillaries leads to an initial drop in oxygenated haemoglobin (oxyHb) and an increase in local carbon dioxide (CO2) and deoxygenated haemoglobin (deoxyHb). Following a lag of 2 - 6 seconds, regional cerebral blood flow (RCBF) increased, delivering an surplus of oxygenated haemoglobin, washing away deoxyhemoglobin. Apparently, the RCBF increase could be up to 50% in response to the increase in neural activity however, the increase in oxygen consumption is much lower than the increase of the contribution through the blood. So in venous phase (venous capillaries, venules and veins) there is an elevated concentration of oxyhemoglobin (diamagnetic compound) with regard to deoxyhemoglobin (paramagnetic substance).

In addition this technique supposes an advantage because it do not require contrast administration for the study of brain activity.

With this technique Moseley (2006) demonstrated significant pain relief in patients with complex regional pain syndrome (CRPS). Other authors have evaluated the relationship between cortical reorganization, the various forms of pain in patients with phantom limb pain syndrome and the analgesic effect of mental imagery by fMRI; they demonstrated measured activation, in the phantom limb area and explained it by a change in the excitability of cortical neurons previously responsive to functions involving the hand or arm only.
Tractography is performed using DTI and computer post-processing to track the fiber bundles which exist in the brain and spinal cord and visualize them as two and three dimensional images. This technique is being studied to determine if neuropathic pain is associated with changes in regional brain anatomy and connectivity.
Fig. 1: Thalamic infarct

© - Valencia/ES
Metabolic cause: Wernicke

Fig. 2: Wernicke

© - Valencia/ES
Fig. 3: Neusosarcoidosis

© - Valencia/ES
Fig. 4: Vascular etiology

© - Valencia/ES
Fig. 5: Schwannoma

© - Valencia/ES
Fig. 6: Vascular etiology

© - Valencia/ES
Fig. 7: Morton’s neuroma

© - Valencia/ES
Fig. 8: Schwannoma

© - Valencia/ES
Fig. 9: Plexopathy

© - Valencia/ES
Conclusion

Multiple mechanisms can cause NP. MRI plays a key role in diagnosis and management of these structural and functional pathologies. The development of MRI methods will provide a new approach in the study of NP syndromes.
References

Patterns of neurovascular compression in patients with classic trigeminal neuralgia: A high-resolution MRI-based study.


Central poststroke pain: An abstruse outcome


Advances in brain imaging of neuropathic pain

CHEN Fu-yong, TAO Wei and LI Yong-jie Chinese Medical Journal 2008; 121(7):653-657

Brain imaging of neuropathic pain


Neuroimaging of pain: what does it tell us?

Karen D. Davis DOI:10.1097/SPC.0b013e3283458f96

Neuroimaging Studies of Chronic Pain. Do Hyung Kang, MD, June Hee Son, BA†, and Yong Chul Kim, MD*Korean J Pain 2010 September; Vol. 23, No. 3: 159-165