Introduction to Medical Psychology Lecture 9: Chronic Pain

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https://youtu.be/yyCSYHy0JxM

Lecture video at above link.

Today: Chronic Pain

- Basics of pain perception (Nociception)
- Top-down control of pain perception
- Chronic low back pain
- Treatment strategies for chronic low back pain



How does pain work? Mechanoreceptors

Δ

Various receptor types are responsible for the perception of touch (marked with green arrow).

Free nerve endings in skin and internal organs respond to tissue damage, causing pain.



Nociceptors (neurons detecting painful stimulation)

Nociceptors located in skin, visceral organs, tendons, etc. can be grouped into four classes:

- Thermal (responds to temperatures > 43 deg Celsius or < 17 deg Celsius)
- Mechanical (responds to intense mechanical stimulation)
- Polymodal (responds to thermal, mechanical, or chemical stimulation)
- Silent nociceptor (responds to inflammation in visceral organs)

Nociception is mediated by activating membrane channels that open when stimulated (with a thermal, mechanical or other stimulus). When opening they let positively charged Na⁺ or Ca²⁺ ions into the neuron, triggering an action potential.



Nociceptive membrane channels

Several different membrane proteins are responsive to different types of stimulation. E.g., the <u>TRPV1</u> (transient receptor potential cation channel V1) channel responds to heat (> 43 degree Celsius), acid (protons), and capsaicin (a vanniloid found in chili plants). <u>TRPA1</u> is activated amongst other substances by AITC (mustard/wasabi) and cold temperature (< 17 degree Celsius).



Nociceptor pathway

Nociceptors have their cell bodies in the dorsal root ganglia (see cross section of spinal cord below). They enter the spinal cord through the dorsal root and connect to neurons within the substantia gelatinosa.



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Dermatomes

Somatosensory and pain nerves enter the spinal cord grouped along so-called dermatomes.



Pain Transduction

The pain pathway uses two types of nerve fibers:

- fast Aδ fibers
- slower C fibers

Speed depends on the diameter of the nerve fiber and its coating with myelin (electrical insulation).



Pain transduction

These differences in transmission speed lead to two pain stages:

1) A fast "first pain" phase mediated by A δ fibers.

2) A slower, longer lasting"second pain" phase mediatedby C fibers.



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Pain pathway

Nociceptors synapse with spinal cord neurons (1).

(3

Small dorsal

root axons

The nerve fibers cross the body midline at the level of the spinal cord segment.

The <u>spinothalamic</u> <u>tract</u> consist of fibers (2) from the spinal cord to thalamus ("sensory hub").

Neurons from thalamus project to somato-sensory cortex (3), insular and cingulatecortex.



Insula and Cingulate cortex



Hyperalgesia after pain

Intense pain can lead to changes in the sensitivity of synapses (long-term potentiation).

These implicit pain memory traces can lead to increased pain sensitivity and chronic pain.

Treating the acute pain with analgesic drugs might help to prevent this hyperalgesia effect.



Top-down control of pain



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In highly emotional or stressful situations (fight, sports tournament, etc.), pain perception can be reduced (analgesia).

This requires control of the pain pathway by cortical centers.



Top-down control of pain

The PAG (periaquaeductal gray) in midbrain and the Raphe nuclei are relay stations of cortical control of pain and can modulate pain-related neural activity in the spinal cord.

Electrical stimulation of these structures can reduce pain perception.



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Top-down control of pain



Opioids (from poppy plants) can reduce pain. Receptors for opioids can be found in the brain, but also in PAG, the Raphe nuclei and the dorsal horn of the spinal cord.

The nervous system produces endogenous opioids: endorphins.



Effect of attention on pain

Somatosensory cortex (SI, SII), anterior cingulate cortex (ACC), and insula are modulated by prefrontal areas (DLPFC: dorsolateral prefrontal cortex, VLPFC: ventrolateral prefrontal cortex), and in turn modulate ascending pain perception.

Attentional effects on pain perception (less pain when distracted) are assumed to be mediated by this network.



Effect of attention on pain

In this fMRI (functional magnetic resonance) study, 9 subjects either attended or did not attend ('think of s.th. else) a painful stimulus (hot plate).

Aversiveness of stimuli was decreased when pain was unattended. Brain activity in the PAG was higher when the pain was not attended, indicating top-down suppression of pain.





18

Lower back pain



Acute low back pain is very common, about 1/3 of the adult population is affected at least once a year. This can last a few days or weeks. 80% of population (US or Germany) will suffer from low back pain at least once in their lives.

If low back pain persists for longer than 3 months, we speak of chronic low back pain.

Causes for acute lower back pain / lumbago can be various, for example:

- inflammations
- vertebral disk hernia;
 this may lead to pressure
 on nerves, in particular
 the sciatic nerve



Source: Apothekenumschau

The symptoms during acute low back pain can include:

- Muscle ache
- Shooting/stabbing pain
- Pain that radiates down the leg
- Limited flexibility or range of motion



https://www.nlm.nih.gov/medlineplus/ency/imagepages/19503.htm

Only in ~20% of the cases (health information from Germany) can the cause be clearly identified. Nevertheless, the demand for imaging is surging (here for the US):



a Lumbar spine MR imaging, Medicare

Imaging is often not very helpful: even in healthy, asymptomatic patients, imaging reveals irrelevant but alarming findings in ~2/3 of the cases.

The problem with these "false alarms" is that aggressive treatment might follow:



c Lumbosacral injection rates, Medicare

Lumbosacral injections:

Epidural steroids (cortisone) against inflammation, sometimes in combination with local pain-killer injected into skin.

Facet injection: similar injection into facet joints.

http://www.spineinstitutenorthwest.com



Lumbosacral injections:

Epidural steroids (cortisone) against inflammation, sometimes in combination with local pain-killer injected into skin.

Facet injection: similar injection into facet joints.

Another form of aggressive treatment that might follow is lumbar fusion:

61.1 Fusions per 100,000 13.9

d Lumbar fusion rates, degenerative spine conditions

Lumbar fusion involves fusion of two vertebrae to prevent the painful motion.

The usefulness of lumbar injections seems to be limited, systematic reviews suggest that they do not reduce later surgery rates. Facet injections with corticosteroids seem to be no better than saline solution.

Fusion surgery is helpful for fractures and deformities, but seem to be of limited value for treating disk degeneration with back pain alone.



However, patients ask for a "quick fix" by injection or surgery.

But even if these measures are not helpful in all cases: you should go and see a doctor for your acute back pain, in particular when it was caused by a fall or trauma, or it gets worse over time, or it causes bowel/bladder problems, or it is accompanied by fever or weight loss.

Chronic low back pain has a social component, too: patients may become work disabled and may claim early retirement.



As shown in the graph, the proportion of disabled workers due to musculoskeletal conditions is increasing (US data).

From a learning theory perspective, patients may experience "rewards" when feeling pain:

- No need to go to work
- Early retirement
- Family might care more for patients
- "Back pain" as a physical impairment is more accepted than psychological ailments such as lack of motivation or depression.

Thus, instrumental conditioning (being rewarded for pain) can lead to intensified pain or more attentional focus on pain.



Multimodal treatment: Application of the biopsychosocial approach

- medical treatment of physical complaints
- strengthening of supporting muscles
- muscle relaxation
- pain diary
- cognitive-behavioral psychotherapy
- ergonomic workplace
- stress coping strategies

Kamper et al., BMJ, 2015: Cochrane systematic review and meta-analysis

What is already known on this topic

Multidisciplinary biopsychosocial rehabilitation programmes are widely used for people with chronic low back pain

Published reviews provide conflicting evidence regarding effectiveness of the programmes and do not quantify the size of the effects on key outcomes of pain, disability, and work absence

What this study adds

Based on the largest collection of trials and participants reviewed to date, this study provides robust estimates of the effects of multidisciplinary biopsychosocial rehabilitation programmes

Patients participating in these programmes are likely to gain small, long term benefits in improved pain and disability compared with usual care or physical treatments

They also have increased odds of being at work compared with patients receiving physical treatment

Patients participating in these programmes are likely to have a similar outcome to those receiving surgery but are less likely to experience adverse events

During acute pain, muscles are contracted and stiffened to protect tissue, skeleton. In chronic pain, this muscle tension can contribute to increased pain (by pressure) and decreased range of motion.

Muscle relaxation: via progressive muscle relaxation

or bio-feedback:

EMG (electromyography) can be shown to patients in real-time. Patients learn to control and relax muscle tension with this feedback.



Pain diary:

There may be relationships between situations and the experience of pain: Environment may be more caring when pain is expressed more.

(However, general social support has been shown to improve chronic pain; important is that social support should not be a consequence of the expression of pain).

Pain might occur when work is more stressful -> avoidance of work.

To understand these relationships and then address them, it is important to first clarify these relationships by logging the occurrence of pain.



In Japan, the Aichi Medical University (Nagoya) has a multidisciplinary pain center, founded in 2007, the first in Japan.



Von Korff et al. (Pain, 2005) investigated a population of ~5700 US citizens; within 1 year:

19% showed chronic spinal pain

of those:

69% showed another type of chronic pain 35% showed mental disorders in particular anxiety disorders mood disorders (depression)



Summary: Chronic Pain

Chronic pain

- Basics of pain perception (Nociception)
 - nociceptors, spinothalamic pathway, thalamus, somatosensory cortex, insula, anterior cingulate cortex
- Top-down control of pain perception
 PAG, opioid receptors, ventrolateral and dorsolateral prefrontal cortex, attentional modulation
- Chronic low back pain
 - psychological factors: e.g., attention, learning
- Treatment strategies for chronic low back pain

cognitive-behavioral psychotherapy, muscle relaxation, pain diary