

Types of Trigger Points

- Active = Can spontaneously trigger local pain or referred symptoms to more distal locations
  - "The TP" that reproduces the patients symptoms
  - Further they can cause muscle weakness, joint restrictions and autonomic phenomenon. Most needling courses only focus on local TP's
- Latent = Just a sore TP that may or may not refer symptoms and may or may not be linked to the patients problems
  - Do not usually spontaneously trigger local or referred symptoms but may alter muscle activation patterns and contribute to joint and movement dysfunctions
- Associated or Satellite = TP's that develop in the referral zone and often in the same myotome
- > Primary = A TP that is linked to main symptoms
- Secondary = a TP that develops due to the Primary TP, e.g. Trap refers pain over Temporalis and is Primary, while temporalis becomes involved and develops its own TP's which may or may not refer symptoms

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# Defining Terms...

- Trigger Point (TrP) = a hyperirritable spot within a taut band of skeletal muscle that is painful on:
  - ➤ Compression
  - > Stretch
  - > Overload or
  - > Contraction of the tissue which typically responds with referred symptoms perceived distal from the location
- Active TrP's = Those TrP's that reproduce famillar symptoms for the patient – may occur at rest and be the symptom generator
- Latent TrP's = Those TrP's that are clinically quiescent with respect to spontaneous symptoms, they may reproduce symptoms only when palpated
- https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4458928/

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# Defining Terms...

- BOTH Active and Latent TrP's have the ability to alter muscle movement patterns, function and reproduce pain and/or autonomic symptoms such as: altered sensation
- Myofascial Pain Syndrome (MPS) = is a common source of musculoskeletal pain and it is characterised by the presence of TrP's
- MPS = affects muscles and their connective tissue attachments (tendons/fascia)
- MPS are often under Diagnosed and affects numerous people suffering from pain





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### Travell and Simons TP Model



- The TP is painful on compression and can give rise to characteristic referred pain, tenderness and autonomic phenomena, either spontaneously or on digital compression
- Each muscle known referral pattern, well documented (Travell and Simons red books Vol 1 and 2)

Active

Primary

Latent

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Associated

Salellile

Secondary

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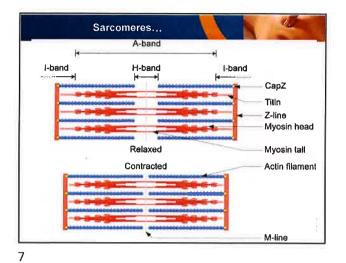
## Myofascial Pain Syndromes (MPS)

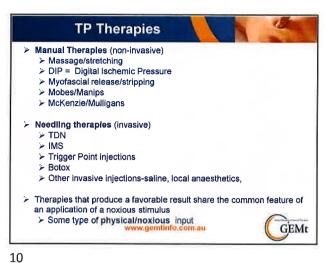


- > It is also a disease likely underpinning many orthopaedic conditions
- An individual contraction knot appears as a segment of a muscle fibre with contracted sarcomeres and an increased diameter
- > We can palpate this under physical examination





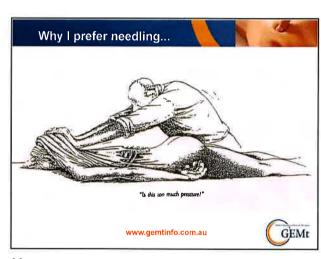




Clinical Features of Myofascial TP's

Point tenderness on a taut muscle band
Local Twitch Response (LTR)
Referred pain/symptoms
Reproduction of usual pain/symptoms
Restricted range of motion
Weakness without atrophy
Autonomic symptoms

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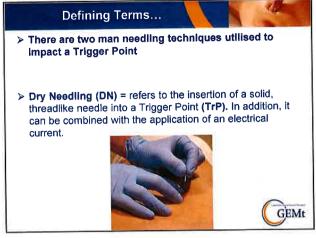


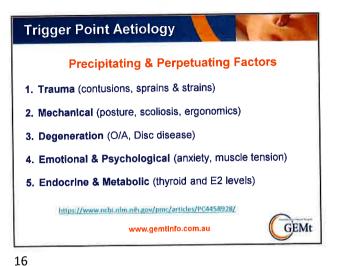
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Precipitating & Perpetuating Factors

6. Respiratory/Hypocapnia (shallow/rapid breathing, vasocon)

7. Infection (Viral or parasitic)

8. Nutritional (Vitamin/Mineral deficiencies)

9. Nerve root compression (Nerve root sensitization)

10. Peripheral Sensitization (nociceptive irritation)

https://www.ncbi.nlm.nih.gov/pmc/articles/PC4458928/

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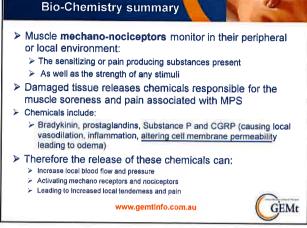
# 1. Energy Crisis Theory Postulates that increased demands on a muscle, such as: Increased trauma (macro or repeated micro) Increased neural input, leads to Increased Ca release from the sarcolemma and prolonged shortening of the sarcomeres, can lead to: Prolonged muscle shortening Compromised circulation Reduced O2 supply

> Rendering cells unable to produce enough ATP to initiate the active process of relaxation

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Shah and Gilliams

This bio-chemical cascade leading to altered pH and influencing both local and central processing

Painful MTrPs activate muscle nociceptors that, upon sustained noxious stimulation, initiate motor and sensory changes in the peripheral and central nervous systems

This process is called sensitization, which can be both peripheral and central

Sensitization of peripheral and central afferents is responsible for the transition from normal to aberrant pain perception

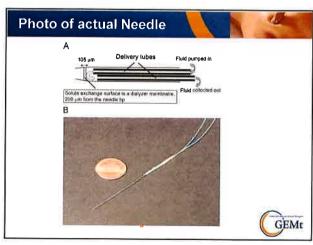
The authors developed a microdialysis technique to quantitatively measure the biochemical milieu of skeletal muscle

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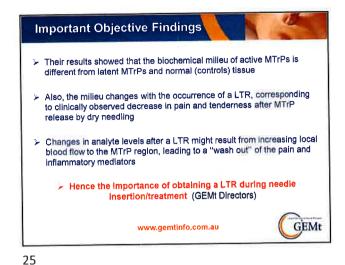
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2. Motor End Plate Hypothesis

The 'Motor End Plate' is where the motor nerve synapses with a muscle cell (neuro-muscular junction)

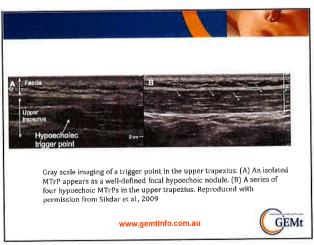
May co-exist with the previous

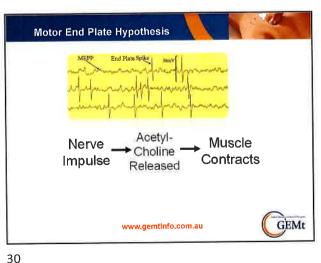
Needle EMG studies have found that each TP contains minute loci that produce characteristic electrical activity or SEA (Spontaneous Electrical Activity)

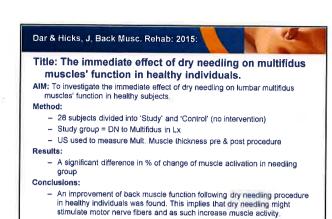
These loci are predominately found at the motor end plate zone

It is postulated that an increased rate of ACh is released from the nerve terminal resulting in increased action potentials (increased End Plate Noise) along the muscle cell membrane – resulting in muscle shortening

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Contraindications to Dry Needling

Scalp areas of infants
Following axillary or inguinal node dissection-avoid any breakage of skin (arm) due to risk of Lymphedema (Dr Beth Baughman DuPree)
Over rib cage or over a rib (Level 2)
High dose anti-coagulants
Immunosuppressed patients-Ca
Blood donors-plasma only 4 months
Nipples, umbilicus and external genitalia

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Complications Post Needling

> Pneumothorax
> Muscle Soreness
> Fatigue:
> Vasovagal Reaction/Fainting
> Bruising
> Needle insertion pain
> Metal allergies
> A Stuck needle
> A Broken needle
> A Forgotten needle
> Neurovascular bundles - knowledge of Anatomy is paramount

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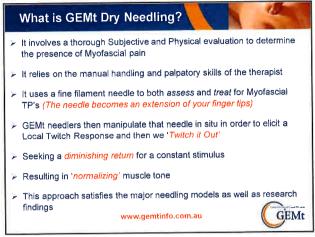
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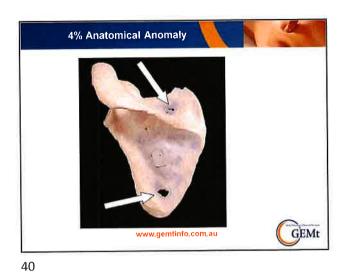
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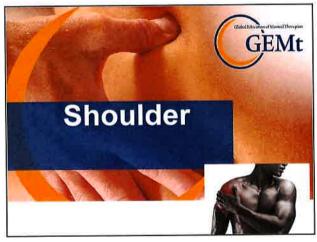


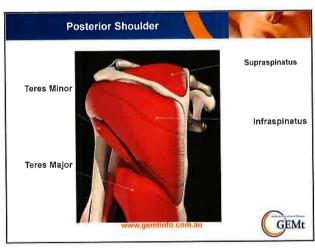
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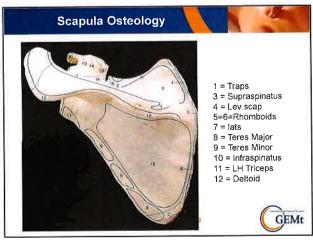


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