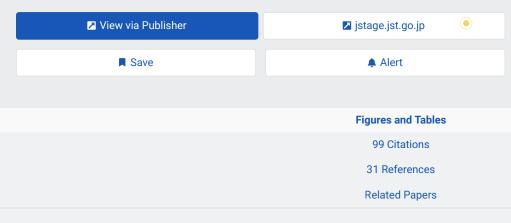
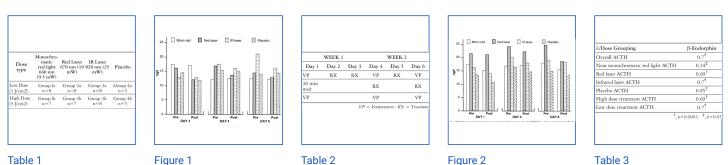
## PLASMA ACTH AND β-ENDORPHIN LEVELS IN RESPONSE TO LOW LEVEL LASER THERAPY (LLLT) FOR MYOFASCIAL **TRIGGER POINTS**

E. Laakso, T. Cramond, +1 author J. Galligan • Published 1994 • Medicine • Laser therapy

The mechanism by which laser phototherapy (Low Level Laser Therapy-LLLT) induces analgesia in the treatmet of chronic pain is not understood. To investigate a possible role for opioids in this treatment, a double-blind placebo-controlled study was designed to compare the effect of two dosages (1 j/cm and 5 j/cm) of an infrared (IR) lase r (820 nm), a visible red laser (670 nm) and A near-mo nochromatic light emitting dcvice (660 nm. 30 nm bandwidth) on Trigger points. Fifty-six consenting subjects with chronic pain conditions exhibiting myofascial trigger points in the neck and upper trunk region underwent six experimental sessions over a two week period. Blood samples were withdrawn before and after treatment on three of six appointments. Plasma was assayed for β-endorphin (radioimmunoassay, RIA) and adrenoeorti cotropic hormone (ACTH two site immunoradiometric assay, IRMA) to assess opioid response. ACTH was shown to have a cumulative response to treatment with 1 J/cm infrared laser (p < 0.001) and 5 J/cm red laser (p < 0.05) responding significantly.  $\beta$ -endorphin was noted to be significantly elevated between days one and four (p <0,05) in subjects who received IR(5 j/cm) laser. Results indicated that the analgesic response to phototherapy may be mediated through hormonal/opioid mechanisms, and that responses to LLLT are dose and wavelength dependent A mechanism by which peripheral stimulation using LLLT may clicit activity in the central pathways is proposed. Collapse



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Figure 1

Table 2

Figure 2

Table 3

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