



**Do Cannabinoids have a
role in cancer pain?**

A short history of cannabis

Culpepper

“The decoction of the root eases the pains of the gout, the hard humours of knots in the joints, the pains and shrinking of the sinews, and the pains of the hips.”

- | **8000 BC** Hemp cultivation.
- ▣ **2800 BC** China *Pen ts'ao Ching*.
- | **2000 BC** India *Atharva Veda*.
- | **4 AD** Obstetric analgesia.
- | **60+ AD** Dioscorides and Galen.
- | **1653** Culpepper.
- ▣ **1842** O'Shaughnessy.
- | **1860s** Queen Victoria.

The Science

Cannabinoid Receptors

CB₁

Neurons

- Inhibit adenylate cyclase.
- Activate K⁺ channels.
- Inactivate Ca⁺⁺ channels.

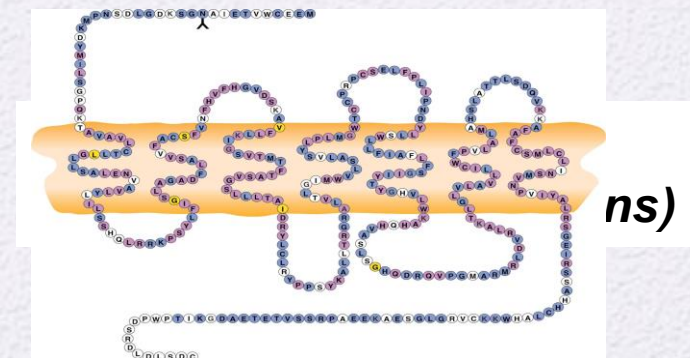
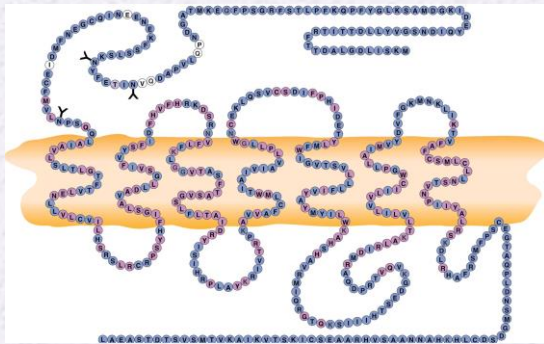
CB₃

GPR55

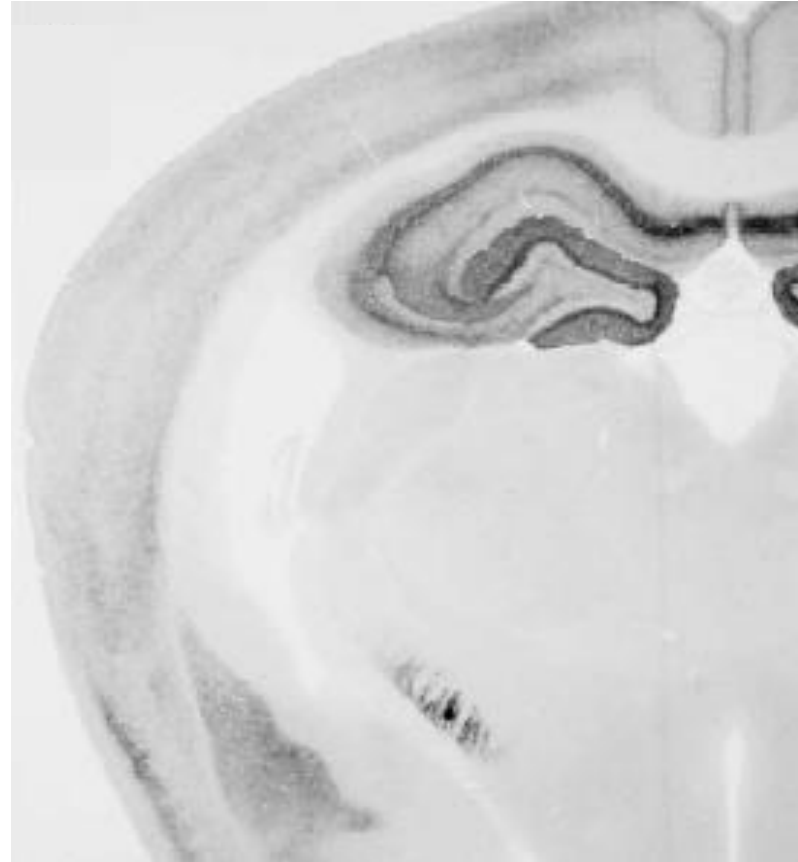
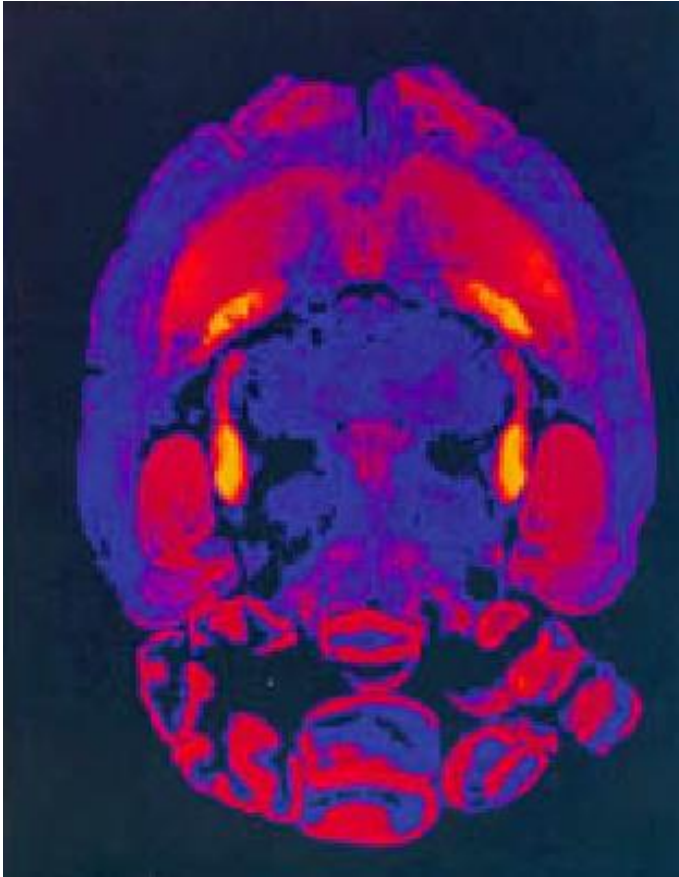
CB₂

Immune cells

- Inhibit adenylate cyclase.

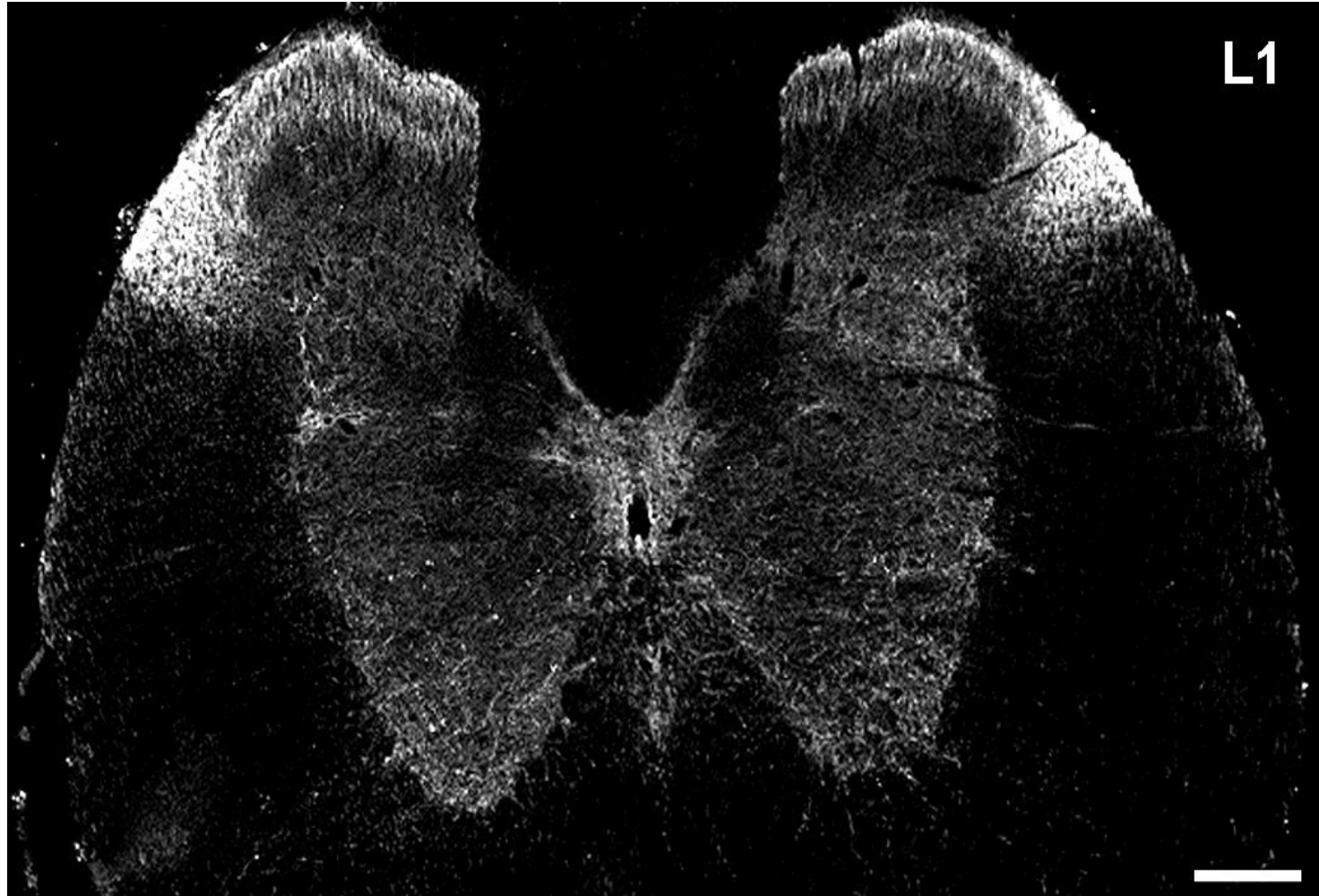


Brain



Herkenham et al 1991; Egertová et al 2003

Spinal Cord



Reported effects of cannabis and $\Delta^9\text{THC}$

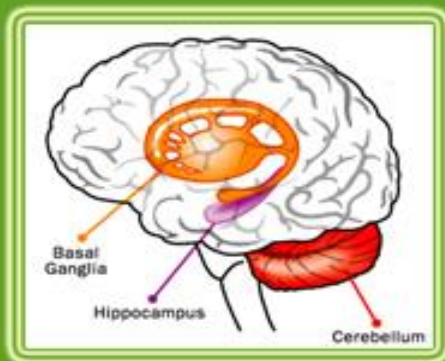
- **CNS:**

- Euphoria/"high".
- Heightened sensory perception.
- Impaired cognition and psychomotor performance.
- Distortion of space & time sense.
- Memory impairment.
- Fragmentation of thoughts.
- Antinociception and analgesia.
- Anti-emesis.
- Increased appetite.
- Altered thermoregulation.

- Schizophrenia-like syndrome.



CANNABINOIDS IN MEDICAL CANNABIS



ENDOCANNABINOIDS

Anandamide(AEA)

PHYTOCANNABINOIDS

THC, CBD, CBN, etc

SYNTHETIC CANNABINOIDS

THC Only(Marinol)

ENDOCANNABINOID RECEPTORS

(Brain Receptors)

CB1, CB2, etc

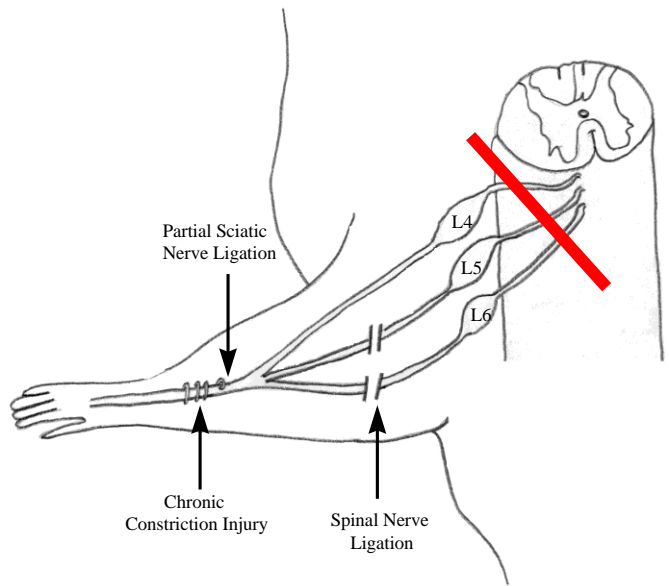
The Endocannabinoid System(ECS) is involved in regulating a variety of psychological processes including appetite, pain and pleasure sensation, immune system, mood and memory



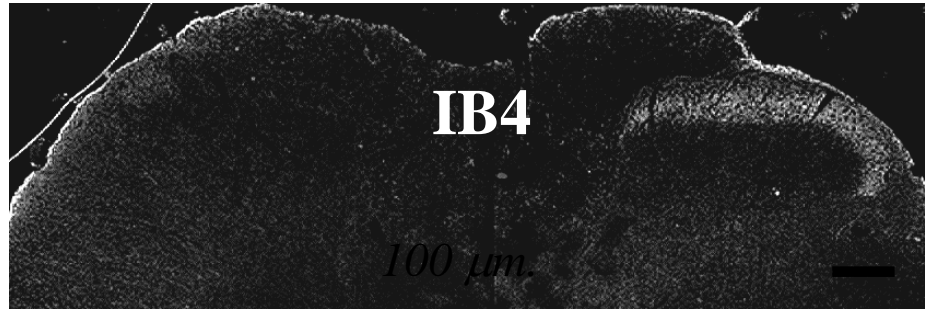
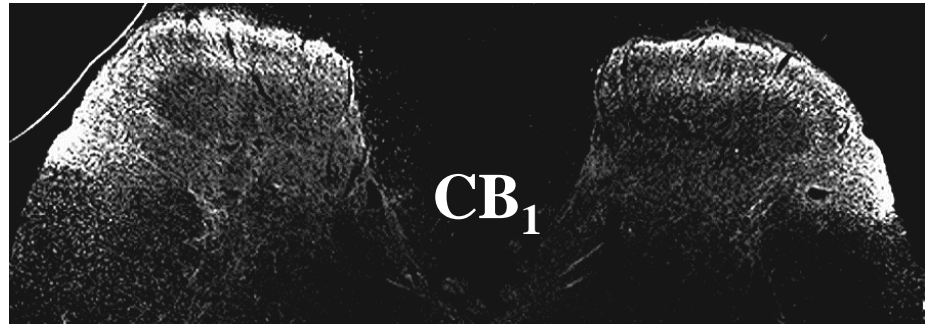
Pertinent pre-clinical
data

Cannabinoid Analgesia in Pain Models

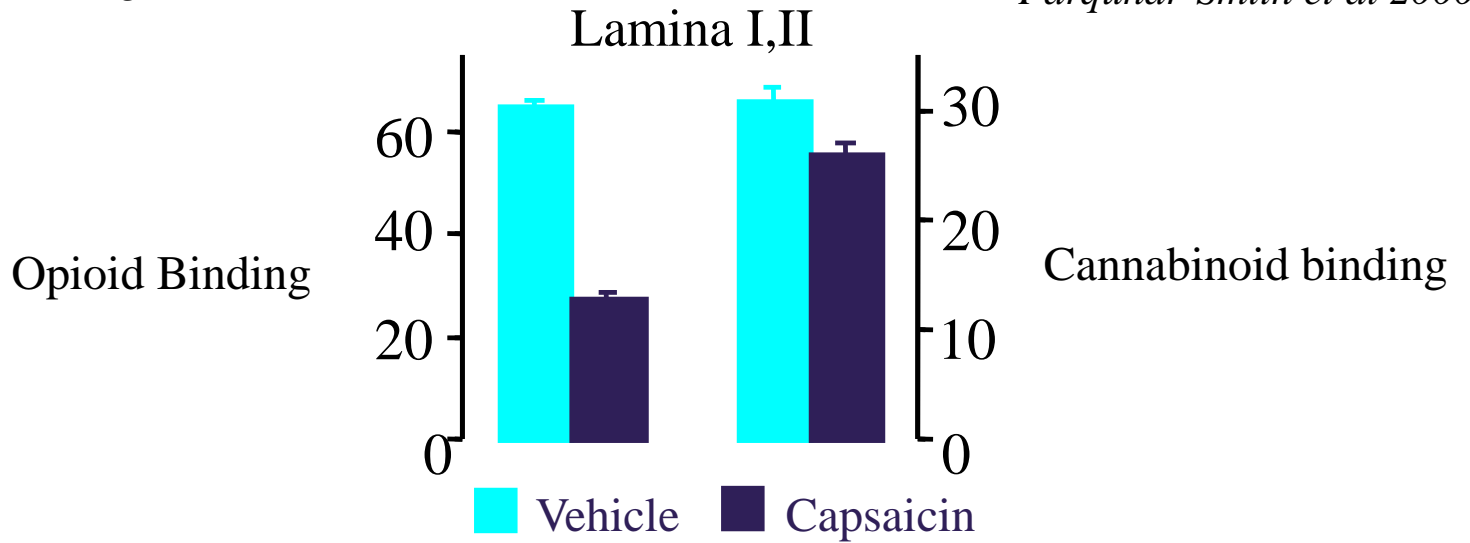
- **Visceral inflammatory pain and referred hyperalgesia**
(*Jaggar et al 1998a,b, Farquhar-Smith et al 2001, 2002*)
- **Formalin** (*Tsou et al 1996, Jaggar et al 1998a, Calignano et al 1998, Hanus et al 1999, Schreiber et al 2012*)
- **Carrageenan** (*Richardson et al 1998*)
- **Cystitis** (*Wang et al 2013*)
- **Collagen-induced arthritis** (*Malfait et al 2000*)
- **Complete Freund's Adjuvant** (*Martin et al 1999, Smith et al 1999*)
- **Chemotherapy induced neuropathic pain** (*Burgos et al 2012, Guindon et al 2013, Khasabova et al 2012*)
- **Tumour induced bone pain** (*Kehl et al 2003, Wang et al 2012, Khasabova et al 2011*)



Bridges et al 2001



Farquhar-Smith et al 2000



Hohmann & Herkenham 1998

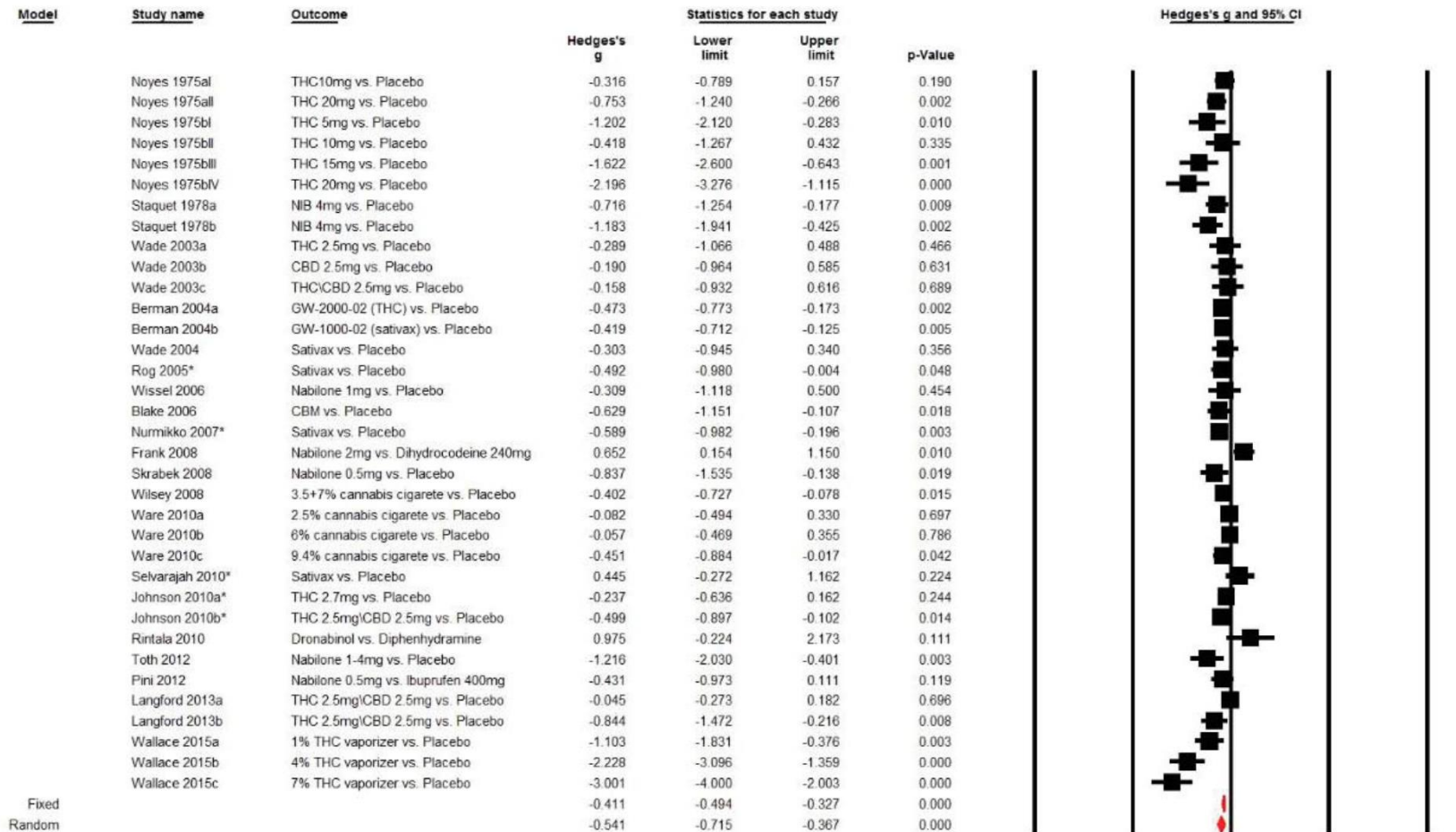
Evidence in humans

Meta-analysis of clinical evidence (17 years ago)

- All clinical trials published prior to 1999.
- 9 trials included
 - 5 cancer pain (119 patients analysed)
 - 2 chronic non-malignant pain (2 patients)
 - 2 acute pain (72 patients)
- THC (5-20 mg p.o.) ~ equi-analgesic to 50 –120 mg codeine
- Dose related and dose limiting CNS adverse events common.

More recent meta-analysis

- Meng et al 2017 Anesthesia Analgesia – chronic neuropathic pain
 - 11 RCT n=1219 NRS 0.65 –weak recommendation
- Whiting et al 2015 JAMA – Medical use
 - Portnoy (2012), Johnson (2010)
- Aviram et al 2017 Pain Physician -



x* = Parallel design.

Fixed 0.411
Random 0.541

Favours Cannabis Favours Placebo

Fig. 2. Meta-analysis- without acute postoperative pain.

All pain (without acute post-operative pain)

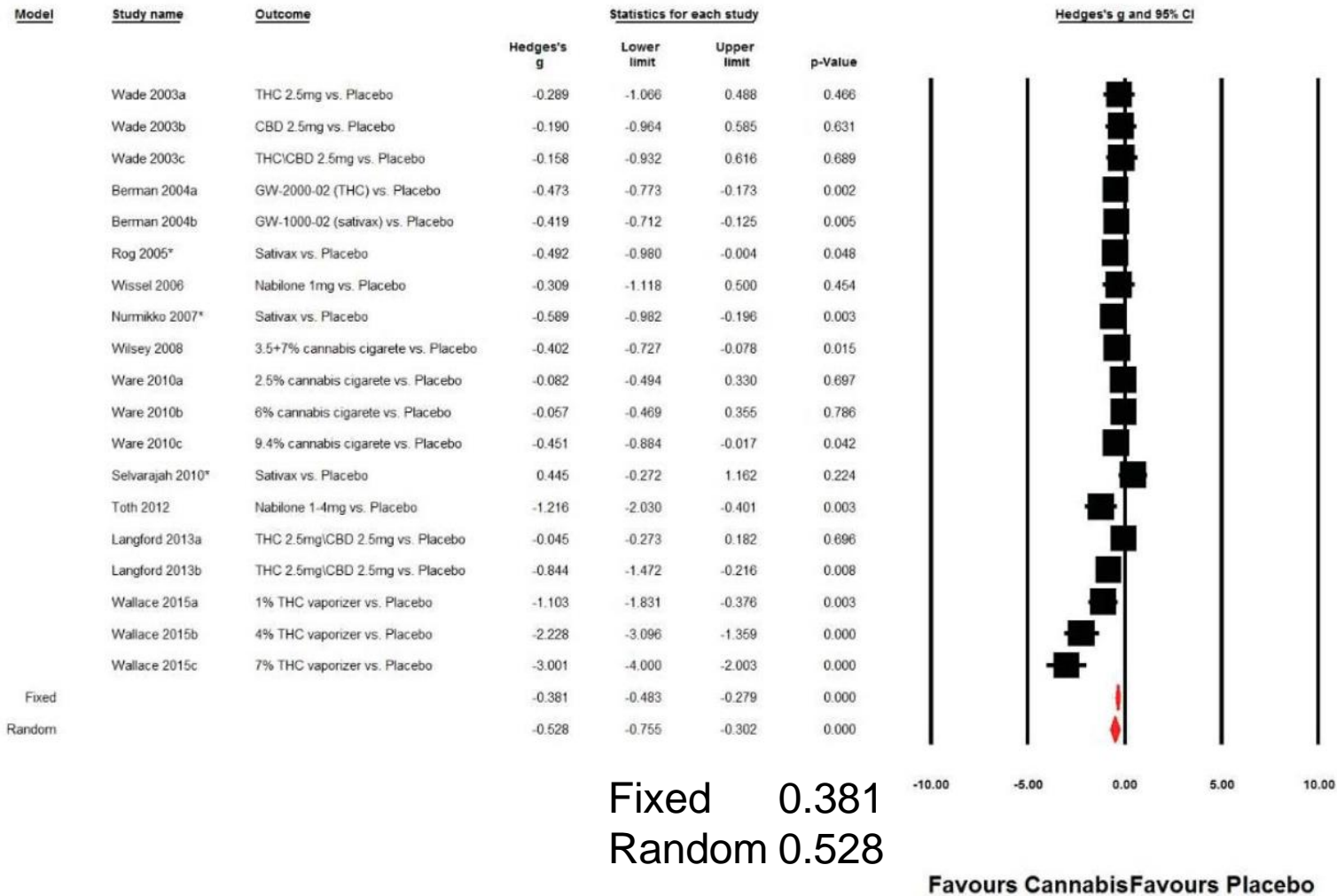
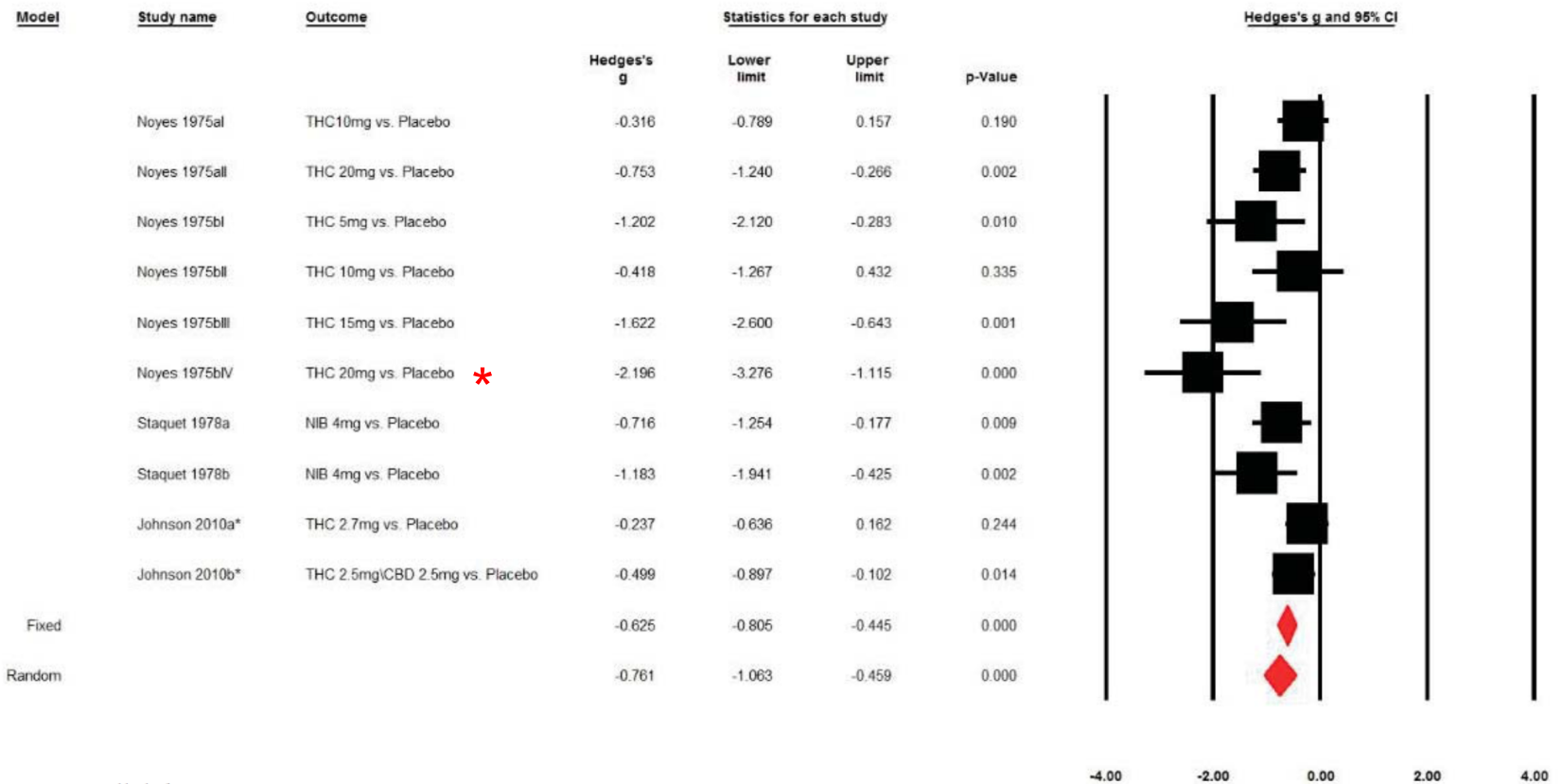


Fig. 6. Meta-analysis- cannabinoids effects on chronic NP.

Chronic neuropathic pain

Yes, but what about
cancer pain?



*= Parallel design

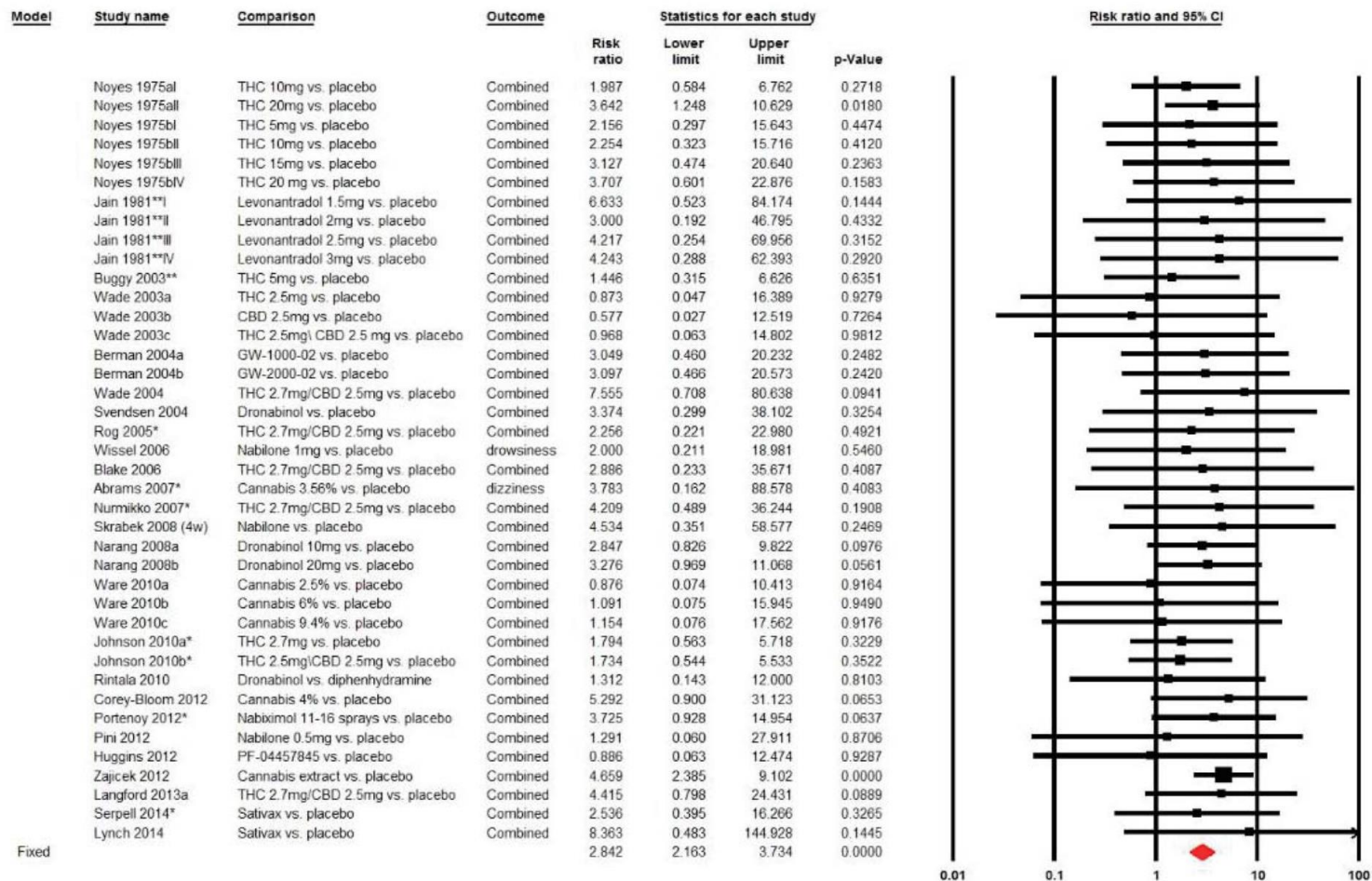
Fixed 0.625
Random 0.761

Favours Cannabis Favours Placebo

Fig. 7. Meta-analysis- cannabinoids effects on cancer pain.

Cancer pain

* “Alarming adverse reactions were also observed at this dose”



Fixed 2.842

Placebo Cannabis

*= Parallel design; **= Postoperative pain.

Fig. 10. Meta-analysis for CNS-related AEs.

CNS-related adverse effects



Cancer Pain

- Patients with uncontrolled cancer pain (NRS 5)
- N=117, (Sativex 60, THC 58, placebo 59)
- NRS -1.37 (Vs. -0.69) (THC NS)
- 30% reduction from baseline in 43% (c.f. 23%)
- No change opioid dose or breakthrough
- N&V increased (85% AE, c.f. 75% placebo)

Cancer Pain

- N=263 (360 randomised), Nabiximols (a.k.a. Sativex)
- 5 weeks low, medium (up to 10 sprays/day) and high dose spray
- 1° outcome: 30% responder, no difference
- But low/medium more 'analgesed' (average daily pain)
- No differences side effects

Cancer pain (not included in meta-analyses)

- N=397
- Self titration Sativex over 2 weeks
- 10.7% improvement (compared to 4.5% control) in NRS – not significant
- Subgroup analysis for QOL
- From US

Cancer pain (not included in meta-analyses)

- 2) PGIC better by 0.27 at week 5
- N=116 (64/52) withdrew in Sativex group
- Median improvement from baseline over 3.7 weeks (N=56 placebo)
7.2% (c.f. 9.5% placebo)
- 68% adverse effects (64% placebo)
- Sub group US <65y 11.2% vs. 4.8%

Pharmacotherapeutic considerations for use of cannabinoids to relieve pain in patients with malignant diseases

This article was published in the following Dove Press journal:
Journal of Pain Research

Results: Fifteen of the 18 trials demonstrated a significant analgesic effect of cannabinoids as compared to placebo. The most commonly reported adverse effects were generally well tolerated, mild to moderate. The main side effects were drowsiness, nausea, vomiting and dry mouth. There is evidence that cannabinoids are safe and modestly effective in neuropathic pain and also for relieving pain in patients with malignant diseases. The proportion of “responders” (patients who at the end of 2 weeks of treatment reported $\geq 30\%$ reduction in pain intensity on a scale of 0–10, which is considered to be clinically important) was 43% in comparison with placebo (21%).

The other problem

Long Term CNS Adverse Effects

- Historical cohort study 1969-70 Swedish conscripts
- n=50,087 (97% of Swedish 18-20 yr. olds)
- Questionnaire of drug use
- National register of psychiatric admissions 1970-1996
- Dose dependent increase risk of developing schizophrenia with cannabis use (30% increased risk of schizophrenia [OR: 6.7(2.1-21.7)])

The bottom line

‘May be beneficial’
(but potential of side effects)

Transdermal CB2 agonist

A close-up photograph showing a person's arm with a transdermal patch being applied. The patch is a small, rectangular, light-colored adhesive. A hand is visible in the upper left corner, holding the patch. The background is a blurred, warm-toned surface.

Chemically favourable

No central side effects

Anti-inflammatory with no NSAID problems



HEALTH BENEFITS OF

CANNABIDIOL

- 
- Antibacterial •
 - Neuron-Protective •
 - Inhibits cancer cell growth •
 - Promotes bone growth •
 - Reduces Blood sugar levels •
 - Reduces inflammation •
 - Reduces risk of artery blockage •
 - Reduces vomiting and nausea •
 - Reduces small intestine contractions •
 - Increases function in immune system •
 - Suppresses muscle spasms •
 - Relieves anxiety •
 - Relieves pain •
 - Tranquilizing •
 - Treats psoriasis •
 - Vasorelaxant •

Evidence base





Anxiolytic Effect During Public Speaking

**CBD significantly reduces
subjective anxiety during
and after public speaking**

**Correct dose is critical to
maximizing effectiveness**

60 men and women, aged 18 to 35

Double-blind study

**5 groups: Placebo, CBD 100mg, CBD 300mg,
CBD 900mg, clonazepam**

Researchers

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