

Riunione Annuale SIN Appulo-Lucana 2022

The Nicolaus Hotel

Via Cardinale Agostino Ciasca, 27 - Bari

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STRESS, DEPRESSIONE ED EMICRANIA: INFLUENZA DI GENERE

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Referral Center Rare Disease - Huntington Disease

Basic Medical Sciences, Neurosciences and Sense Organs Department

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Table 1 Pain conditions showing differential prevalence between men and women [9, 15–17]

<i>Female prevalence</i>	<i>Male prevalence</i>	<i>No sex prevalence</i>
Head and neck	Head and neck	Head and neck
Migraine with and without aura	Cluster headache	Acute tension-type headache
Chronic tension-type headache	Post-traumatic headache	Toothache due to dentinocervical defects
Chronic paroxysmal hemicrania	SUNCT syndrome (Short-lasting Unilateral Neuralgiform headache attacks with Conjunctival injection and Tearing)	Toothache due to pulpitis
Hemicrania continua	Lesions or diseases of the nervous system	Cracked tooth syndrome
Hypnic headache	Brachial plexus avulsion	Stylohyoid process syndrome
Primary stabbing headache	Postherpetic neuralgia	Lesions or diseases of the nervous system
Post-dural puncture headache	Meralgia paresthetica	Secondary trigeminal neuralgia
Cervicogenic headache	Lateral femoral cutaneous neuropathy	Neuralgia of nervus intermedius
Periapical periodontitis and abscess	Pain associated with Pancoast tumor	Painful ophthalmoplegia
Atypical odontalgia	Raeder paratrigeminal syndrome	Vagus nerve neuralgia
Burning mouth syndrome	Vascular & hematologic origin	Thoracic outlet syndrome
Carotidynia	Thromboangiitis obliterans	Brachial plexus tumors
Temporal arteritis	Coronary artery disease (before age 65 years)	Postherpetic neuralgia
Lesions or diseases of the nervous system	Erythromelalgia (over 50 years)	Maxillary sinusitis
Trigeminal neuralgia (tic douloureux)	Gastrointestinal tract & visceral organs	Dry socket
Carpal tunnel syndrome	Pancreatic disease	Pain in Parkinson's disease
Causalgia	Duodenal disease	Gastrointestinal tract & visceral organs
Reflex sympathetic dystrophy	Abdominal migraine	Esophageal motility disorders
Pain in multiple sclerosis	Myofascial and osteoarticular pains	Chronic gastric ulcer
Occipital neuralgia	Hemophilic arthropathy	Crohn's disease
Vascular & hematologic origin	Ankylosing spondylitis	Diverticular disease of the colon
Chronic venous insufficiency	Gout (after age 60 years)	Carcinoma of the colon
Acute intermittent porphyria	Osteoarthritis (before age 45 years)	Pain of vascular & hematologic origin
Raynaud's disease		Familial Mediterranean fever
Chilblains/Pernio		Hereditary coproporphria
Livedo reticularis (after age 40 years)		
Gastrointestinal tract & visceral organs		
Esophagitis		
Reflux esophagitis with peptic ulcer		
Gallbladder disease		
Post-cholecystectomy syndrome		
Irritable bowel syndrome		
Bladder pain syndrome		
Proctalgia fugax		
Myofascial and osteoarticular pain		
Piriformis syndrome		
Peroneal muscular atrophy		
Rheumatoid arthritis		
Fibromyalgia and chronic widespread pain		
Gout (before age 60 years)		
Slipping rib syndrome		
Twelfth rib syndrome		
Temporomandibular disorder pain		
Back pain (postmenopausal)		

Prevalenza di sindromi da dolore cronico nel sesso femminile? ..dipende

Pain Ther (2021) 10:287–314
<https://doi.org/10.1007/s40122-021-00244-1>

REVIEW

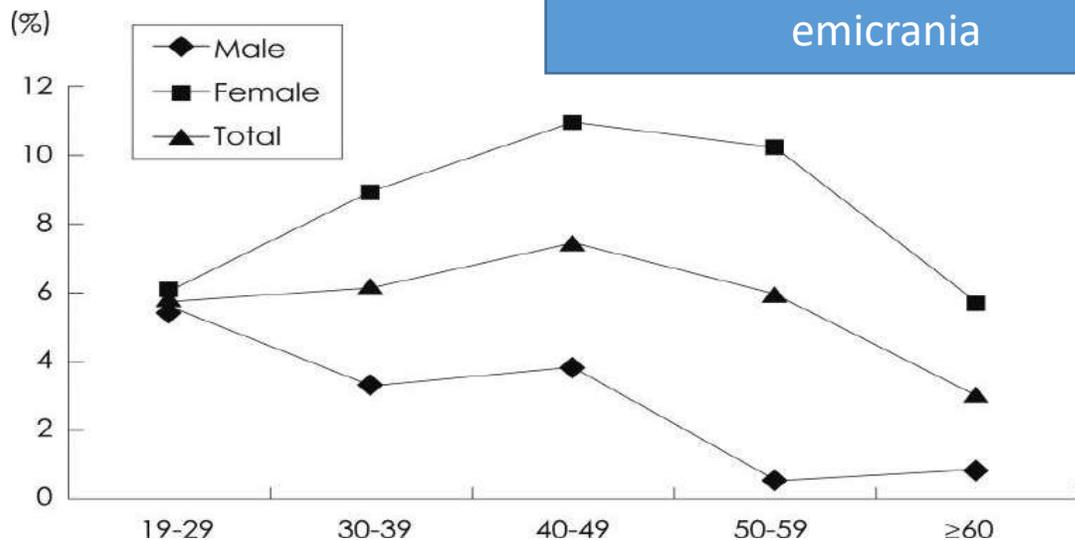
Pain in Women: A Perspective Review on a Relevant Clinical Issue that Deserves Prioritization

Roberto Casale  · Fabiola Atzeni · Laura Bazzichi · Giovanna Beretta ·

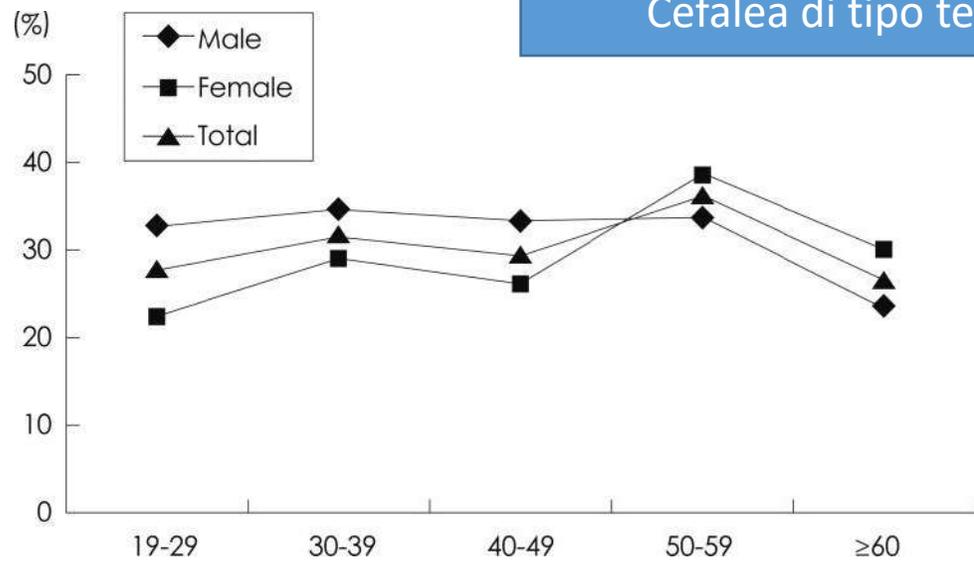
Elisabetta Costantini · Paola Sacerdote · Cristina Tassorelli



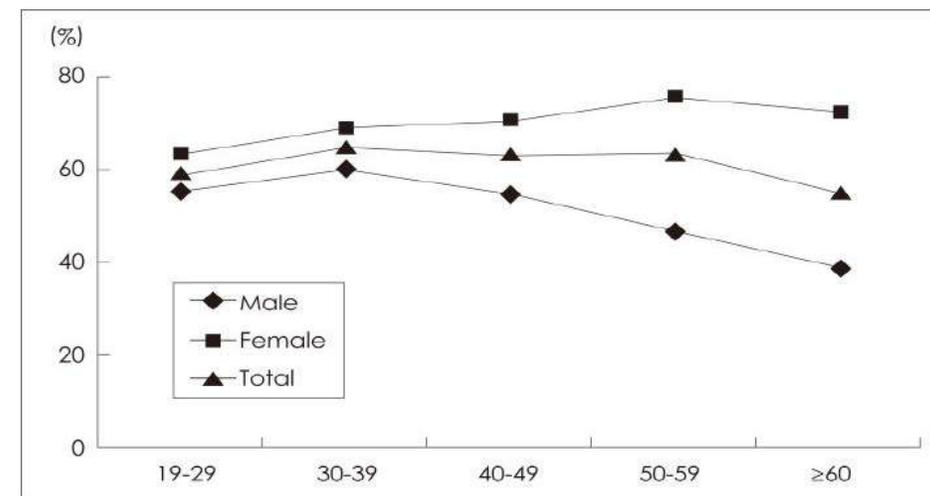
Prevalenza in 1 anno



Cefalea di tipo tensivo



Prevalenza di sindromi da dolore cronico nel sesso femminile? dipende...



Tutte le cefalee primarie

[J Clin Neurol. 2012 Sep; 8\(3\): 204–211.](#)

Nella neuropatia diabetica, la sensibilità al dolore è più elevata che nei pazienti di sesso maschile

Journal of the Neurological Sciences 388 (2018) 103–106



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Sex differences in neuropathic pain intensity in diabetes

Alon Abraham^{a,*}, Carolina Barnett^b, Hans D. Katzberg^b, Leif E. Lovblom^c, Bruce A. Perkins^c, Vera Brill^b



Diversa manifestazione fenotipica del dolore cronico nelle donne? Dipende....

Demographics, symptoms, sensory findings, and TCNS scores in 128 patients with type II diabetes and diabetic polyneuropathy (cohort #2).

	Males (n = 84)	Females (n = 44)	p-value
Age	63 ± 14	59 ± 14	0.13
DM duration (years)	13 ± 11	15 ± 10	0.61
HbA1c (%)	7.3 ± 1.8	7.9 ± 2.1	0.23
Symptoms			
Duration (years)	5	6	0.68
Pain (%)	46	60	0.14
NRS (0–10)	6.9 ± 2.5	8.5 ± 1.7	0.02
Numbness (%)	72	65	0.43
Tingling (%)	52	56	0.72
Upper limb (%)	41	58	0.08
Weakness (%)	41	34	0.48
Ataxia (%)	65	71	0.50
Weakness (%)	41	34	0.48
Sensory deficits			
Pinprick (%)	77	76	0.84
Temperature (%)	73	61	0.18
Light Touch (%)	53	57	0.69
Vibration (%)	76	70	0.46
Proprioception (%)	37	30	0.58
TCNS scores			
Symptoms	2.7 ± 1.3	3 ± 1.5	0.32
Sensory deficits	2.9 ± 1.7	2.8 ± 1.7	0.68
Reflexes	4 ± 2.7	4.42 ± 2.84	0.44
Total	9.74 ± 4.1	10.2 ± 4.5	0.62

DM – Diabetes Mellitus; NRS – numerical rating scale. Statistically significant p values (< 0.05) are bolded.



Sex Differences in Chronic Migraine: Focusing on Clinical Features, Pathophysiology, and Treatments

Chia-Kuang Tsai¹ · Chia-Lin Tsai¹ · Guan-Yu Lin¹ · Fu-Chi Yang¹ · Shuu-Jiun Wang^{2,3,4}

Diversa manifestazione fenotipica
del dolore cronico nelle donne?
Dipende....

Table 1 Clinical features of male and female patients with chronic migraine

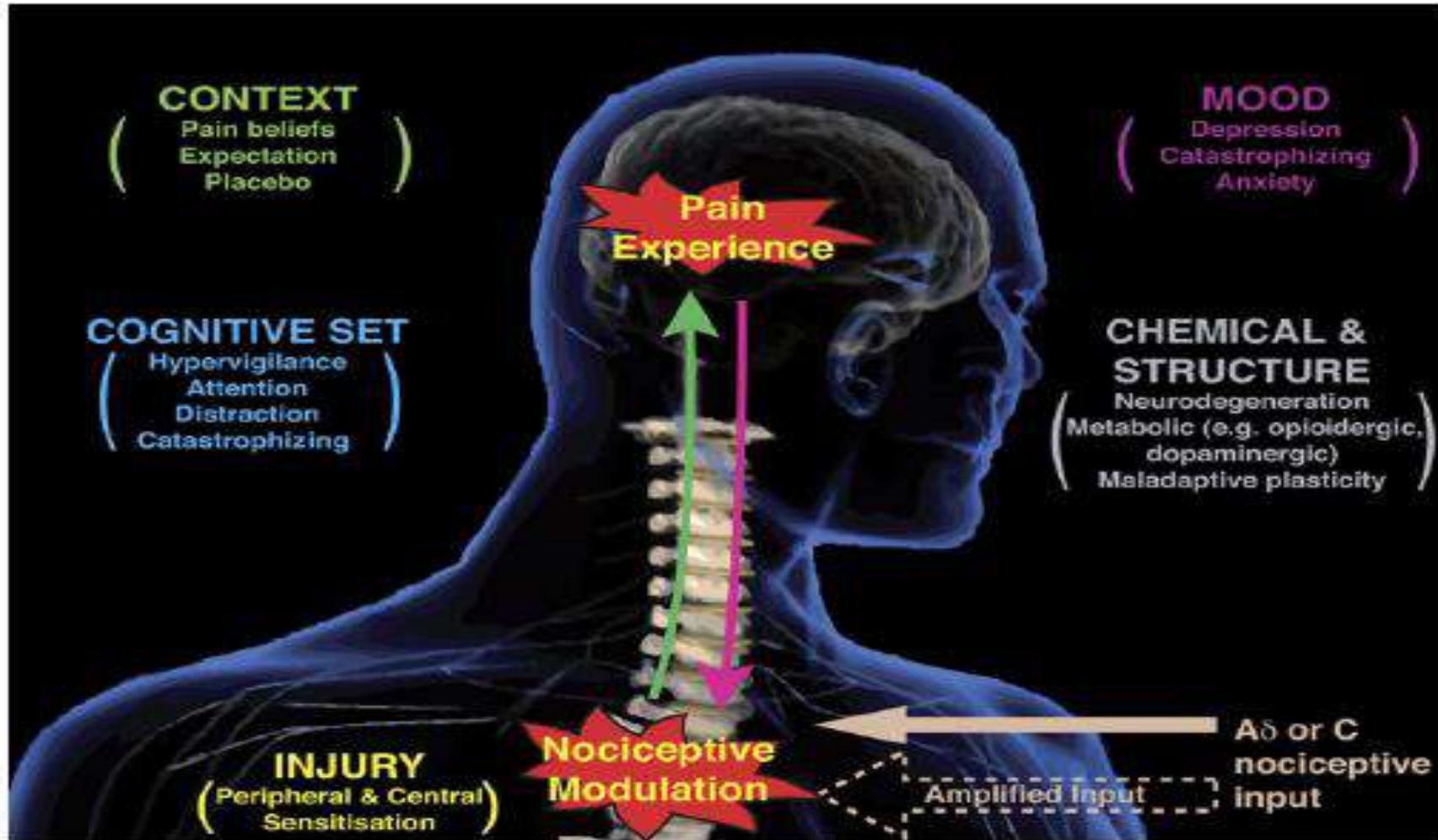
	Male	Female
Prevalence in all migraine population	5.9–6.5	6.9–9.6
Education		
At least graduate degree (%)	0.4	1.2
Less than 12 years (%)	1.2	4.6
Median headache frequency (days/month)	20.0–21.7	20.0
Median MIDAS scores	32–33	38–45
Rates of moderate/severe disability (%)	66.9–71	78.9–82.6
Lost 20 working days within 3 months due to migraine (%)	9.9	15.8

MIDAS Migraine Disability Assessment questionnaire

La manifestazione fenotipica dell'emicrania cronica è diversa nei 2 sessi, poichè le donne manifestano maggiore invalidità e perdita di giorni di lavoro

Neuron Review

Irene Tracey^{1,*} and Patrick W. Mantyh^{2,*}



Perché parlare di dolore di genere? E cosa intendiamo per dolore di genere?

The exploration of mechanisms of comorbidity between migraine and depression

Migraine and depression: bidirectional co-morbidities?

G. Bruti , [M. C. Magnotti](#) & [G. Iannetti](#)

Neurological Sciences **33**, 107–109 (2012) | [Cite this article](#)

546 Accesses | **18** Citations | [Metrics](#)

Abstract

Even if the bidirectional co-morbidity between migraine and depression has been supported by epidemiological and genetic studies, many aspects of this association have not been completely understood. This may be due to the heterogeneous character of migraine and depression as well as to their multifactorial pathogenesis. In this review, we have briefly reported the more recent findings published about the co-morbidity between migraine and depression by discussing the above reported issues and the relative clinical and therapeutic implications.

Meta-Analysis

> [Psychol Bull.](#) 2017 Aug;143(8):783-822. doi: 10.1037/bul0000102.

Epub 2017 Apr 27.

Gender differences in depression in representative national samples: Meta-analyses of diagnoses and symptoms

> [Psychiatry Res.](#) 2013 Dec 30;210(3):1301-3. doi: 10.1016/j.psychres.2013.09.027.

Epub 2013 Oct 2.

Gender differences in depression and anxiety: the role of age

[Carlo Faravelli](#)¹, [Maria Alessandra Scarpato](#), [Giovanni Castellini](#), [Carolina Lo Sauro](#)



Le differenze di genere nella percezione del dolore sono state un argomento di crescente interesse negli ultimi anni e conseguenti studi clinici hanno evidenziato che le donne dimostrano una maggiore sensibilità al dolore cronico. Soffrire nell'anima e nel corpo è fatto individuale, ma la cultura e gli ormoni portano a un soffrire diverso nell'uomo e nella donna.

- Gli ormoni sessuali modulano la neurotrasmissione agendo come “steroidi neuroattivi”.



LA TOLLERANZA E LA SOGLIA SONO *IN GENERE* RIDOTTE NEL SESSO FEMMINILE

Gli estrogeni e gli androgeni influenzano la percezione del dolore

Table 3
Studies Examining Sex Differences in Pressure, Electrical, and Ischemic Experimental Pain Models

Authors	Sample Size (M/F)	Stimulation Site	Method	Threshold*	Tolerance*	Ratings†
Pressure pain						
Ayesh et al ¹⁹	24/19	F	PA	F = M	M > F	—
Chesterton et al ⁶⁶	120/120	H	PA	M > F	—	—
Ellermeir and Westphal ¹¹¹	18/18	FNG	PA	—	—	F > M
Fillingim et al ¹²³	39/49	T, M, U	PA	M > F	—	—
Fillingim et al ¹³²	39/61	T, M, U	PA	M > F	—	—
Garcia et al ¹⁴¹	12/18	TP	PA	M > F	—	—
Komiyama and De Laat ²²⁷	16/16	M, H	PA	M > F	M > F	F = M
Komiyama et al ²²⁸	44/44	M, H, L	PA	M > F	M > F	F > M
Nie et al ²⁹⁵	12/12	H	CCPS	F = M	—	F = M
Electrical pain						
al' Absi et al ¹	59/40	FA	ES	M > F	M > F	—
Ashina et al ¹⁴	9/12	T, TM	ES	M > F	M > F	—
Ayesh et al ¹⁹	24/19	F	ES	F = M	—	—
Nyklicek et al ²⁹⁸	26/23	FA	ES	M > F	M > F	—
Ischemic pain						
Bragdon et al ⁴⁵	22/20	A	SETT	F = M	F = M	—
Edwards et al ¹⁰⁷	83/115	A	SETT	F = M	F = M	—
Fillingim and Maixner ¹²⁸	25/23	A	SETT	F = M	M > F	F = M
Fillingim et al ¹²³	39/49	A	SETT	F = M	F = M	—
Fillingim et al ¹³²	39/61	A	SETT	F = M	F = M	—
Girdler et al ¹⁶¹	40/37	A	SETT	M > F	M > F	F = M

Abbreviations for stimulation sites: A, arm; F, face; FA, forearm; FNG, finger; H, hand; L, Leg; M, masseter muscles; T, trapezius muscles; TM, temporal muscle; U, Ulna.

Abbreviations for methods: CCPS, computer controlled pressure stimulator; ES, electrical stimulation; PA, pressure algometry; SETT, submaximal effort tourniquet test.

* Lower levels of threshold and tolerance in females indicate greater pain sensitivity.

† Higher subjective ratings in females indicate greater pain sensitivity.



Sex differences in pain along the neuraxis

Peyton Presto^a, Mariacristina Mazzitelli^a, Riley Junell^a, Zach Griffin^a, Volker Neugebauer^{a,b,c,*}

Non c'è prevalenza di sesso nella resilienza al dolore.

Therefore, it is difficult to draw firm conclusions surrounding which sex is more strongly affected by or more resilient to pain, as the underlying mechanisms driving these dimensional differences are largely unknown.

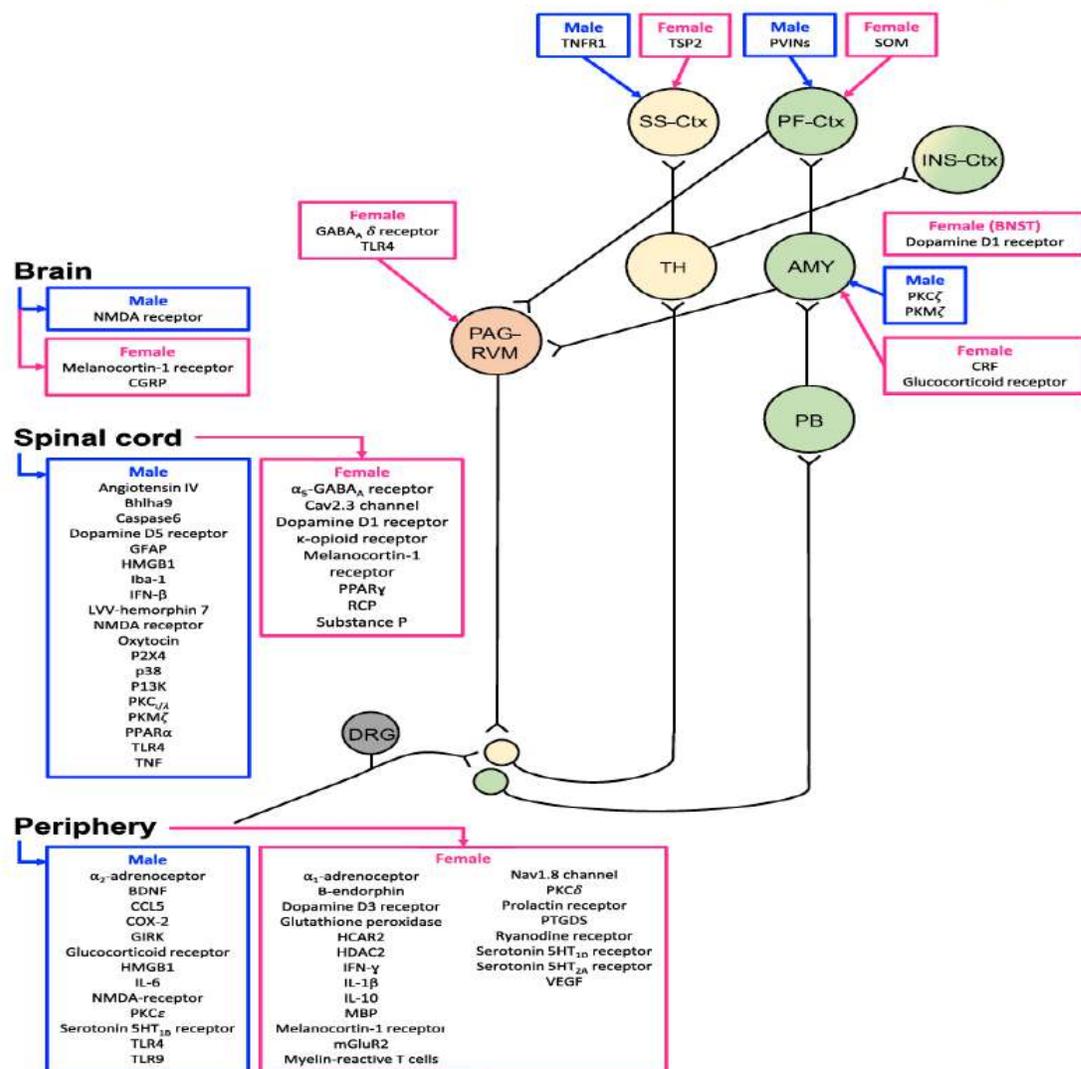
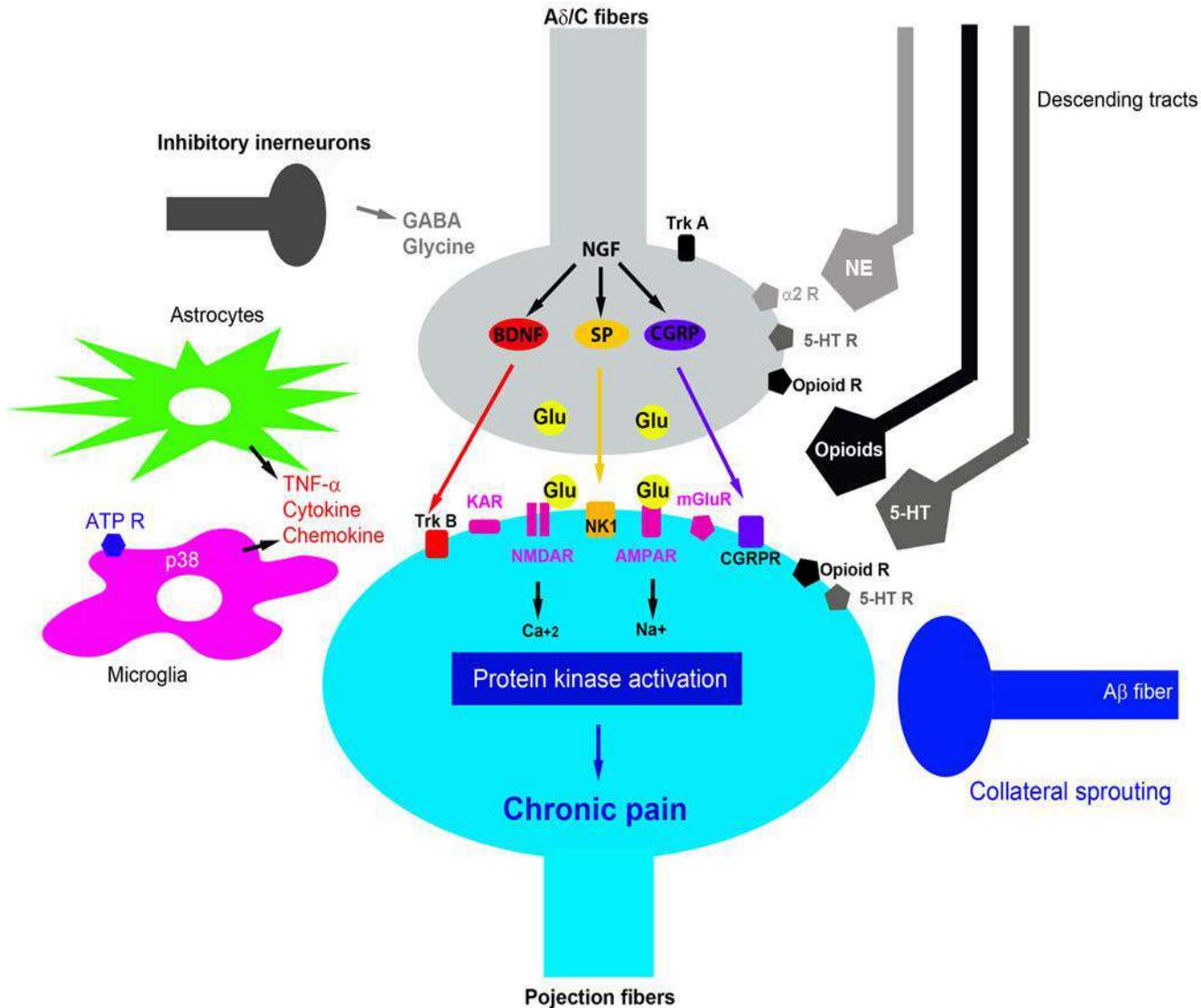


Fig. 1. Sex-predominant cellular and molecular factors within the pain system. First order nociceptive neurons are located in the dorsal root ganglia (DRG) and carry nociceptive information from the periphery into the CNS. These neurons synapse in the spinal cord dorsal horn (DH), and axons from second order neurons decussate in the anterior white commissure to ascend in the contralateral spinal cord. Pain-related information is conveyed through at least three main pathways: the spinothalamic tract (STT, shown in yellow), the spinoparabrachioamygdala tract (SPbA, shown in green), and the postsynaptic dorsal column (PSDC) pathway (not shown). Sensory information is conveyed through the STT to the ventral posterolateral and -medial (VPL and VPM) nuclei of the thalamus (TH) before synapsing in the somatosensory cortex (SS-Ctx) and posterior aspect of the insular cortex (INS-Ctx), allowing for the localization and intensity detection of pain. In the SPbA pathway, also known as the limbic pathway, nociceptive information relays in the parabrachial nucleus (PB) in the brainstem before reaching the amygdala (AMY). AMY integrates multimodal sensory and nociceptive information to alter central autonomic functions and emotional-affective states through associative processing and provides value-based information to prefrontal cortical regions (PF-Ctx) and anterior INS-Ctx. Various brain regions engage the descending pain modulatory system, including efferents from AMY and PF-Ctx that synapse onto the periaqueductal gray and rostral ventromedial medulla (PAG-RVM) system (shown in orange). These brainstem regions then modulate spinal nociceptive processing through descending monoaminergic projections. Cellular and molecular factors that have been reported to have sexually dimorphic effects are illustrated by pink (representing female-predominant effects) and blue (representing male-predominant effects) boxes. BNST, bed nucleus of stria terminalis.

Le modalità di attivazione del sistema nocicettivo sono diverse nei 2 sessi. Ancora meno chiare sono le differenze di genere nell'ambito dei meccanismi immunitari e dell'attivazione gliale



Sex differences in pain along the neuraxis

Peyton Presto^a, Mariacristina Mazzitelli^a, Riley Junell^a, Zach Griffin^a, Volker Neugebauer^{a,b,c,*}



Le donne si adatterebbero di meno al dolore fasico ripetuto, ma dimostrerebbero più abitudine al dolore tonico rispetto agli uomini



Topical review

Deconstructing sex differences in pain sensitivity



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^d Department of Surgery and Institute of Medical Science, University of Toronto, Toronto, ON, Canada

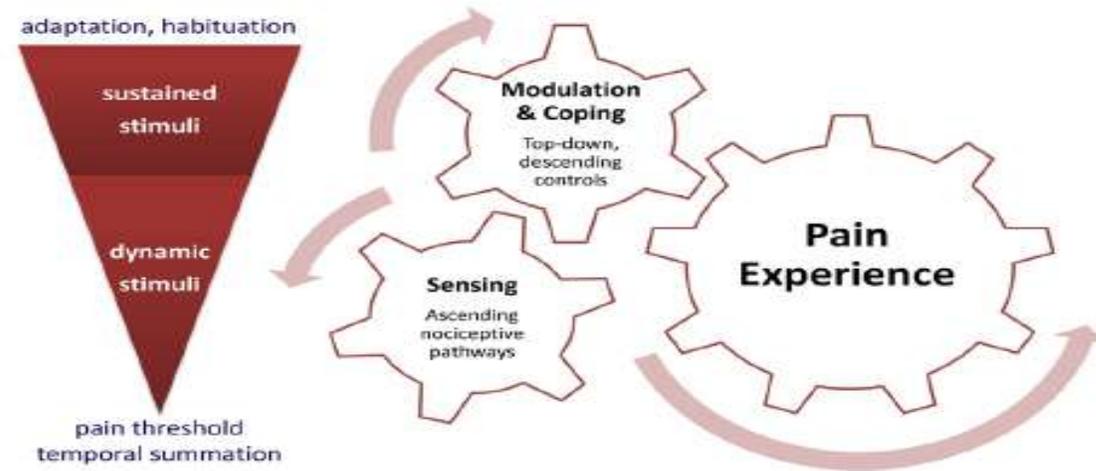


Fig. 2. Model for investigating sex differences in pain. We propose that the overall pain experience derives from “sensing” noxious stimuli by peripheral and ascending nociceptive pathways, which then induce activity in “modulation and coping” systems that include the descending modulation inhibitory pathway and top-down controls. Thus, pain sensitivity comprises elements from both the sensing and the modulation/coping systems. Existing evidence indicates that women are more responsive to dynamic stimuli where the temperature is vacillating or increasing, such as observed in temporal summation and threshold testing. In contrast, women can adapt and habituate more than men to sustained longer duration stimuli. We recommend that future studies of the concept of pain “sensitivity” and sex differences interpret their findings within this conceptual framework. This model indicates that tests that involve threshold and suprathreshold pain responses with brief or long stimuli, and with dynamic or static temperatures, may evoke different responses in men and women, and each variable may be critical for gaining insight into only one aspect of the overall pain sensitivity.

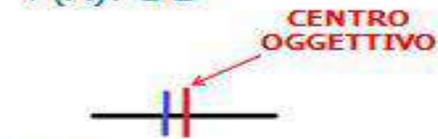
Article in Press

Oral contraceptive therapy modulates hemispheric asymmetry in spatial attention

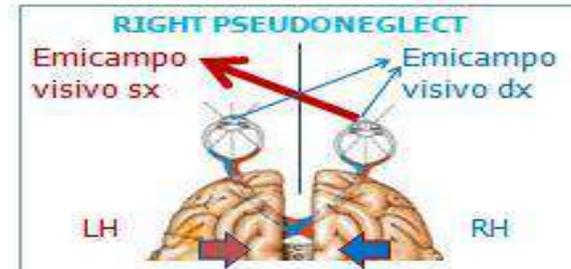
Ettore [Cicinelli](#), Marina De Tommaso, Antonio Cianci, Nicola Colacurci, Leonarda Rella, Luisa Loiudice, Maria Vittoria [Cicinelli](#), Paolo Livrea

DESTRIMANE

❖(A): OC



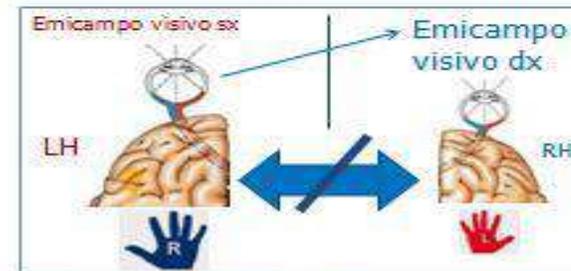
Tendenza a spostarsi a sx



❖(B): ~~OC~~



Tendenza a spostarsi a dx
HAND-USE EFFECT



✓ **MANCINE**: i risultati non sono statisticamente significativi.

MODIFICAZIONI ATTENTIVE
COMPLESSE

I risultati del presente studio confermano le interazioni tra ormoni steroidei e processi cognitivi, suggerendo una significativa influenza da parte dei contraccettivi orali.

Sex differences in the social brain and in social cognition

Alice Mado Proverbio 

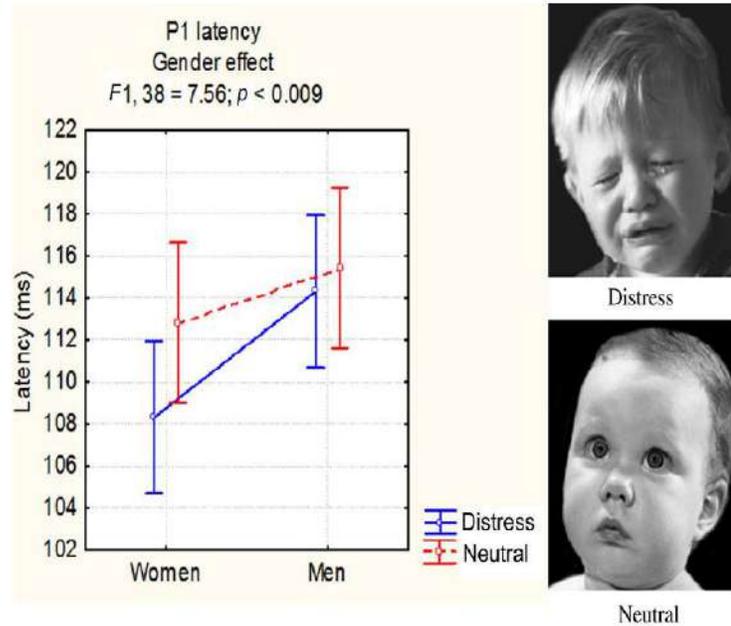
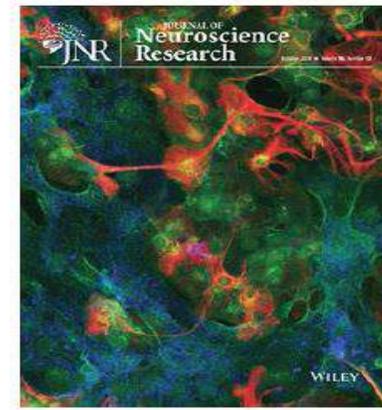


FIGURE 1 Mean latency of P1 component recorded at lateral occipital area (independent of hemispheric site) and analyzed according to viewer's sex and type of facial expression (taken and modified from Proverbio et al., 2006b study, protected by CC BY 2.0)

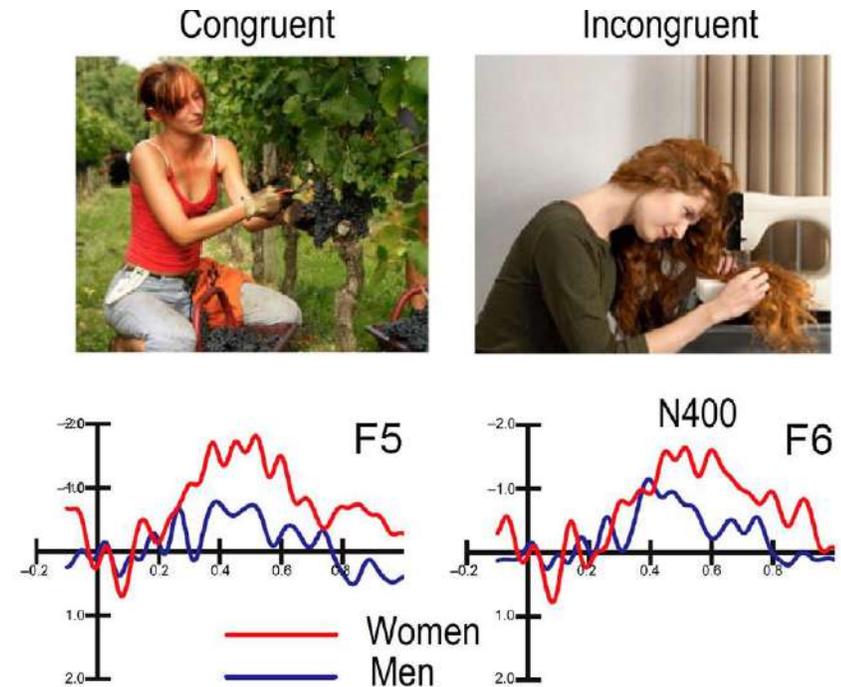


FIGURE 2 Event-related potential (ERP) difference waves obtained by subtracting ERPs to congruent from ERPs to incongruent actions separately for men and women, over anterior scalp sites. It can be appreciated a much larger N400 response to incongruent actions in women than men (original figure, relative to Proverbio et al., 2010, courtesy of the authors)

In summary, we have here outlined a series of sex differences in social behavioral and psychological processes that seem biologically associated with female chromosomal karyotype. They are: efficient and bilateral face processing, marked empathic attitude, sensitivity to facial mimicry and gestures, attachment to infants (induced by oxytocin), early interest for infants, curiosity about people and social information, emotional responsivity, lesser incidence of autistic, psychopathic, and sociopathic disorders.

Maggiore empatia per le
emozioni negative nelle
donne

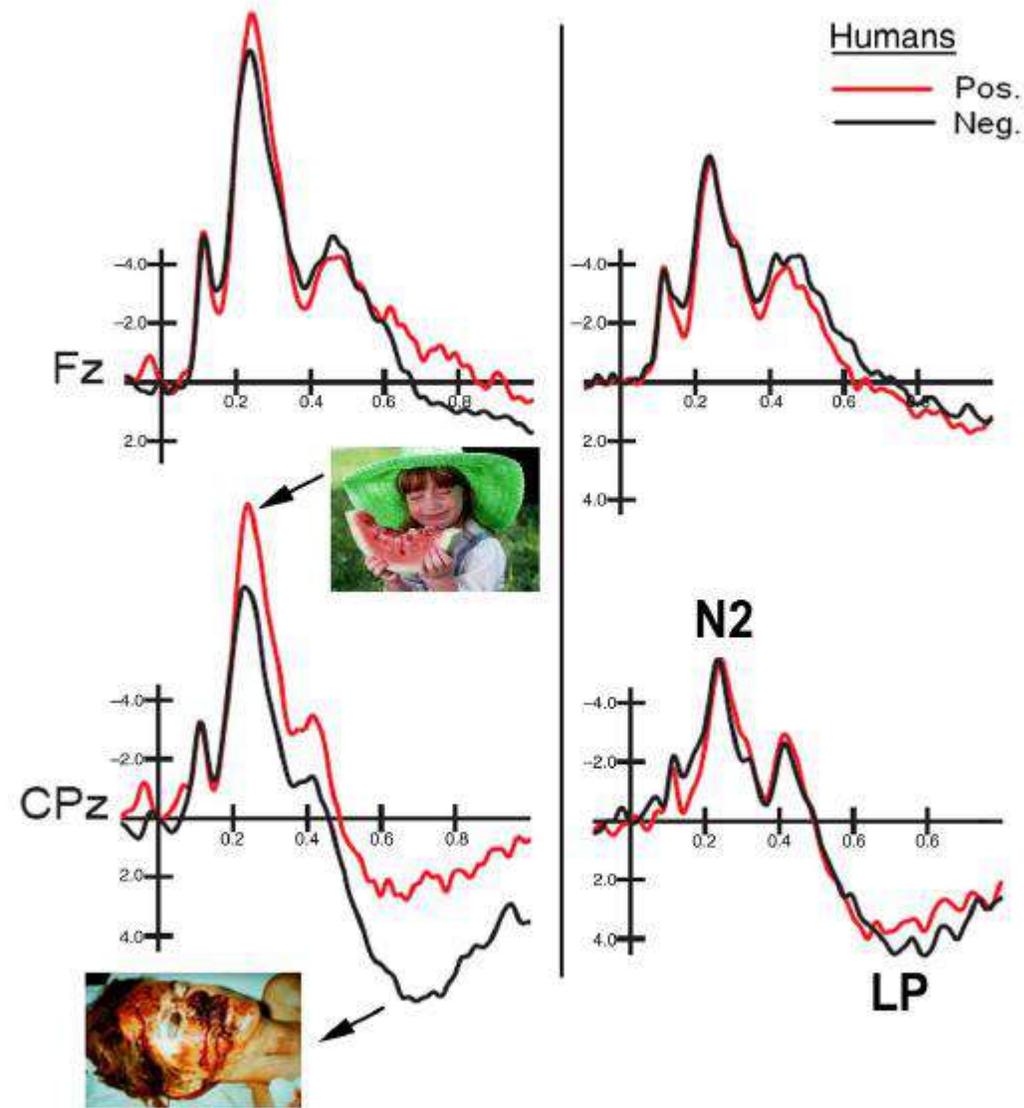
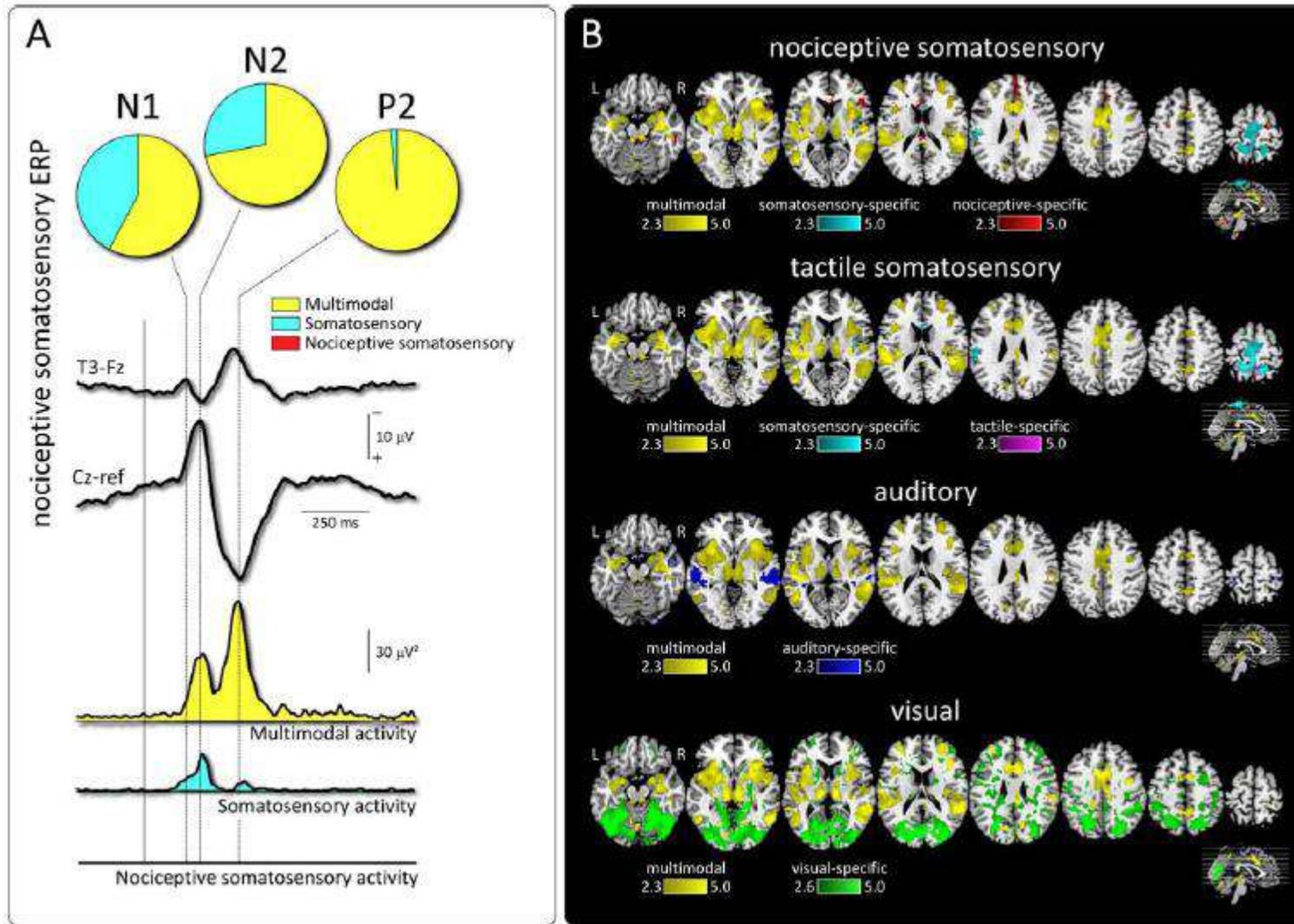


FIGURE 3 Event-related potentials (ERPs) recorded at midline frontal and centro/parietal sites as a function of stimulus valence and viewer's sex. It is visible a large effect of both emotional content (that can be appreciated by comparing ERPs to negative vs. positive unanimated scenes), and an effect of empathy for pain, especially in women (that can be appreciated by comparing ERPs to negative scenes vs. ERPs to pictures portraying humans). Original figure, relative to Proverbio et al. (2009)'s study, reproduced with permission of the authors and of Elsevier

I potenziali evocati da stimolo laser, soprattutto nella componente di vertice, sono un modello di risposta corticale ad uno stimolo “saliente”.
 Il dolore è lo stimolo “saliente” per eccellenza.

V. Legrain et al./Progress in Neurobiology 93 (2011) 111–124



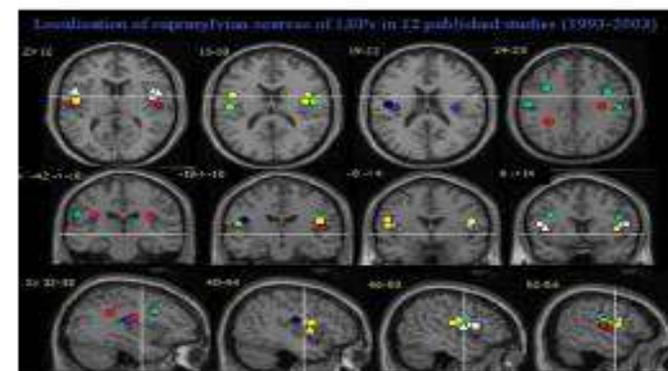
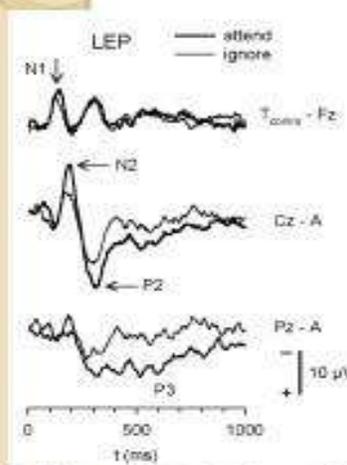
Age-related changes in laser-evoked potentials following trigeminal and hand stimulation in healthy subjects

M. de Tommaso, K. Ricci, A. Montemurno, E. Vecchio

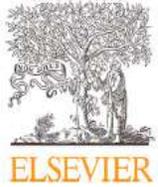
Table 6 Laser-evoked potential features by right hand in 166 females and 61 males and right supraorbital zone stimulation in 111 males and 69 females. The Student's *t*-test between sexes was not significant for any variable.

Sex		Face		Hand	
		Mean	SD	Mean	SD
Pain T (watt)	M	9.15	13.25	6.76	2.18
	F	9.01	12.91	7.10	3.30
VAS	M	50.66	40.10	18.52	12.98
	F	55.66	43.86	26.04	16.31
n1 (ms)	M	129.89	180.44	20.33	22.74
	F	130.19	181.46	19.11	21.04
n1 (uV)	M	5.91	6.12	4.22	6.17
	F	8.47	7.15	5.71	8.23
n2 (ms)	M	199.46	230.64	22.73	28.03
	F	203.07	219.10	27.19	29.59
p2 (ms)	M	321.27	348.75	30.44	34.10
	F	322.79	337.51	33.26	31.96
n2p2 (uV)	M	34.08	31.04	27.88	28.70
	F	28.12	25.11	15.19	17.85
Hab	M	0.66	0.76	0.50	0.47
	F	0.78	0.77	0.48	0.62

I potenziali evocati laser consentono lo studio psicofisiologico del dolore, poiché indicano il grado di attivazione delle aree corticali connesse agli aspetti cognitivi ed emotivi del dolore.



ASSENZA DI DIFFERENZE SIGNIFICATIVE DELLA SOGLIA DEL DOLORE E DEI POTENZIALI CORTICALI INDOTTI DA STIMOLO LASER



L'analisi delle singole risposte, rivela ridotta latenza del complesso di vertice di potenziali nocicettivi nel sesso femminile

Single-trial averaging improves the physiological interpretation of contact heat evoked potentials

Catherine R. Jutzeler^{a,b,c,1,*}, Lukas D. Linde^{d,e,f,1}, Jan Rosner^{c,g}, Michèle Hubli^c, Armin Curt^c, John L.K. Kramer^{d,e,f,*}



C.R. Jutzeler, L.D. Linde, J. Rosner et al.

NeuroImage 225 (2021) 117473

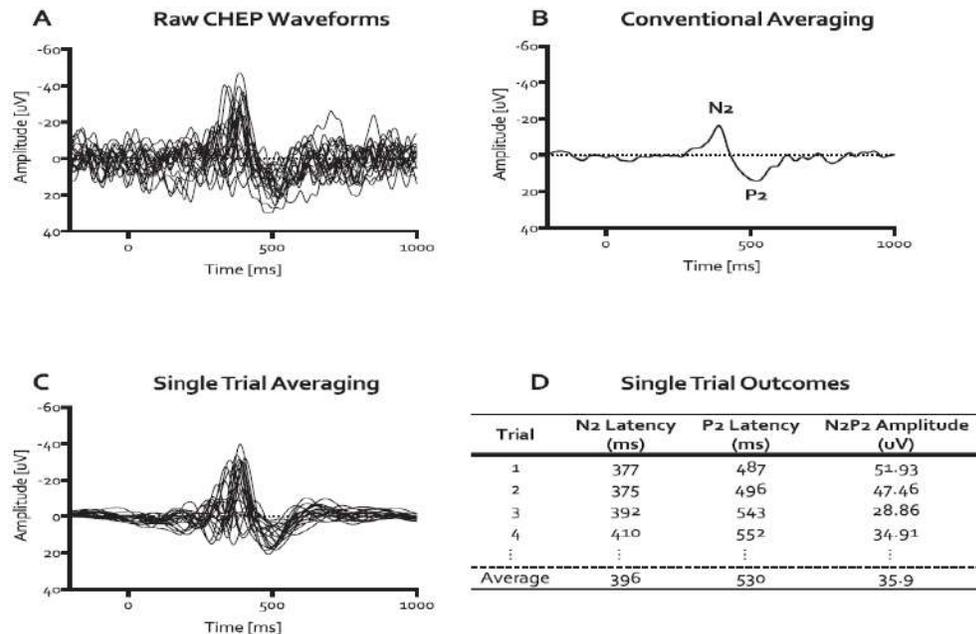
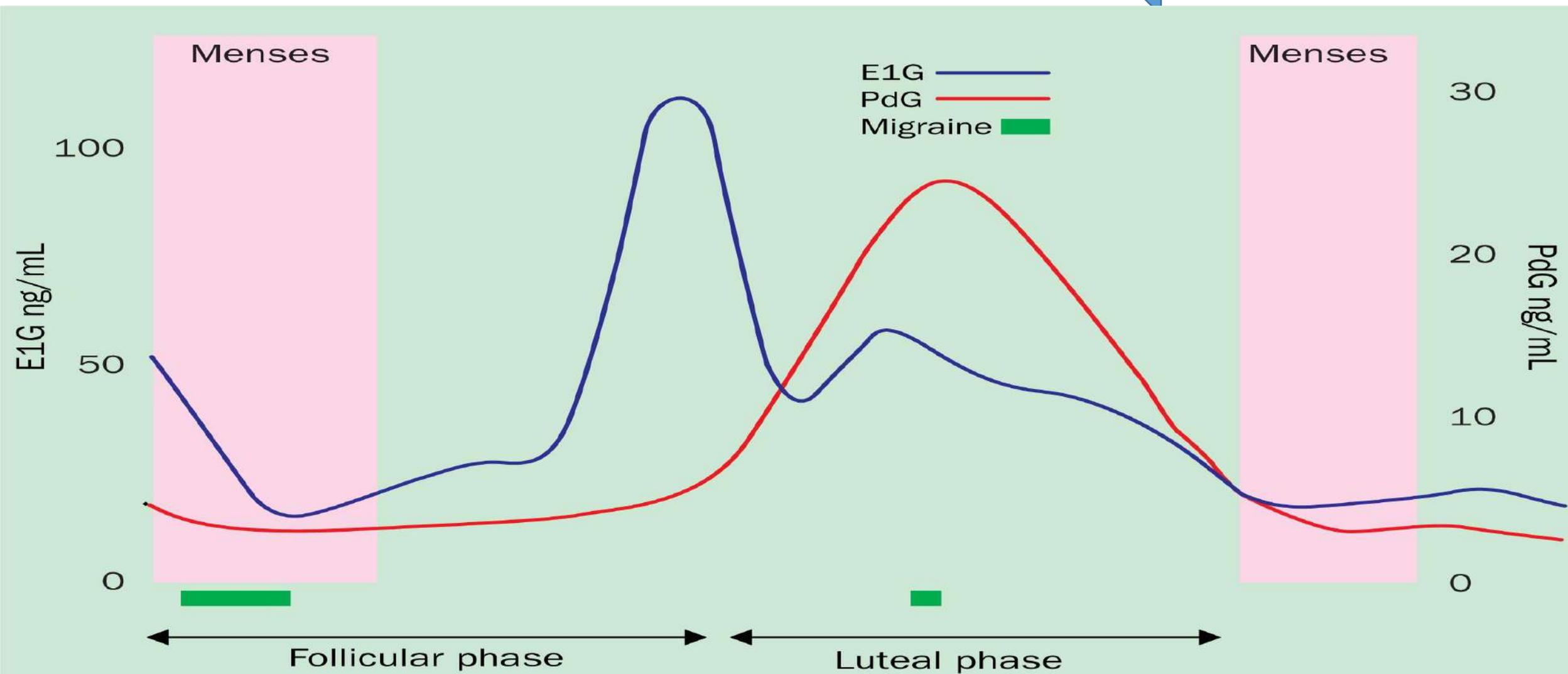


Fig. 1. Representative traces of conventional averaging and single trial averaging analysis methods. A) Individual contact heat evoked potential (CHEP) waveforms, filtered and re-referenced. B) Conventional averaging of individual CHEP waveforms, from which N2 and P2 outcome are derived. C) Individual CHEP waveforms following single trial analysis, via wavelet filtering and multiple linear regression with dispersion term (Hu et al., 2011, 2010). D) Averaged CHEP outcomes determined from single trial analysis.

Our analysis also revealed a significant main effect of sex on N2 ($F_{(1, 07.13)} = 7.06, p < 0.01$) and P2 latencies ($F_{(1, 85.6)} = 10.32, p < 0.01$), as well as a main effect of rating of perceived intensity on N2P2 amplitude ($F_{(1, 480.6)} = 31.7, p < 0.001$).

E' solo un problema di ormoni?

Le sindromi da dolore cronico sono influenzate dai mutamenti ormonali



Influenza degli estrogeni nella percezione del dolore

Misure psicofisiche in modelli animali

Estrogen-dependent changes in visceral afferent sensitivity

.....*

R. Sanoja, F. Cervero / *Autonomic Neuroscience: Basic and Clinical* 153 (2010) 84–89

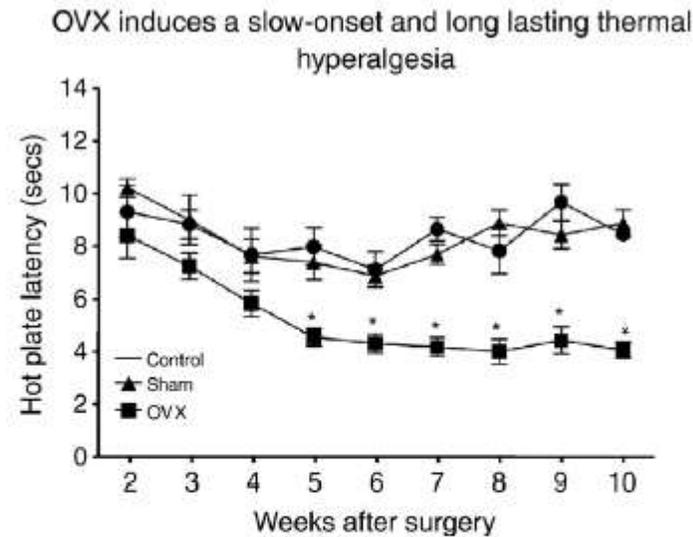


Fig. 2. Estrogen dependency of the hyperalgesic state induced by ovariectomy (OVX). Responses to mechanical stimulation (von Frey hairs) of the abdomen in ovariectomized mice to which slow release pellets containing 17 β -estradiol (OVX + 17 β -estradiol) or the vehicle only (OVX + placebo) had been implanted one week after surgery. The same animals were tested one week and five weeks after surgery. Note that estrogen-replacement after ovariectomy prevents the development of the hyperalgesic state. Significant differences (*, $p < 0.05$) were detected on week five after surgery between OVX + placebo animals and the other three groups. Data from Sanoja and Cervero (2005, 2008).

Nei modelli animali è riscontrabile una chiara relazione fra riduzione degli estrogeni e iperalgesia.

- Ma le donne sono più complesse.....

Il dolore sperimentale non si modifica in relazione ai cambiamenti ormonali

Misure psicofisiche della percezione del dolore correlate al ciclo mestruale

Menstrual cycle phase does not influence gender differences in experimental pain sensitivity

Rebecca R. Klatzkin^b, Beth Mechlin^b, Susan S. Girdler^{a,b,*}

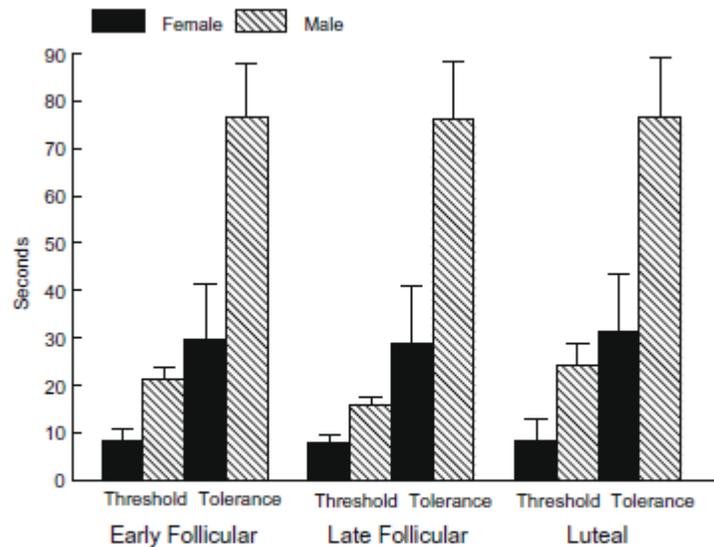


Fig. 1. Cold pressor pain threshold and tolerance as a function of gender and menstrual cycle phase.

- Le fasi del ciclo mestruale non condizionano le differenze di sesso relative alla soglia e alla tolleranza del dolore

Il dolore sperimentale non si modifica in relazione ai cambiamenti ormonali

ABITUAZIONE DEI LEPs NELL'EMICRANIA

Un pattern di ridotta abituazione è stato osservato durante la fase intercritica dell'emicrania. Esso potrebbe esser considerato un fenomeno corticale essenziale per la comparsa ed il ripetersi degli attacchi emicranici

Valeriani et al., 2003



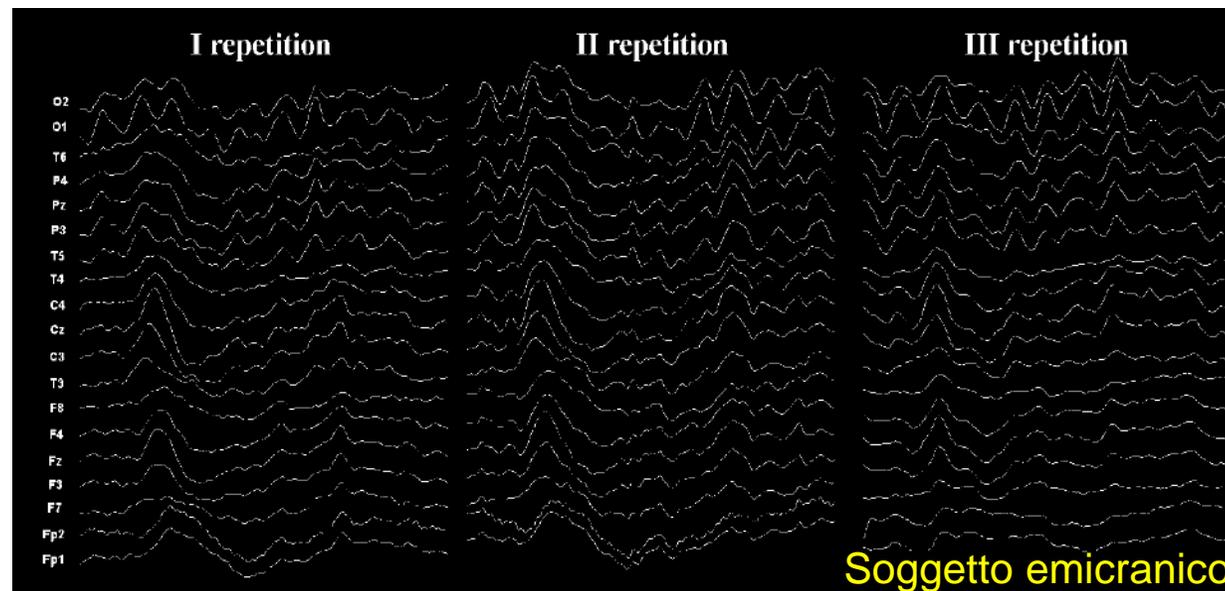
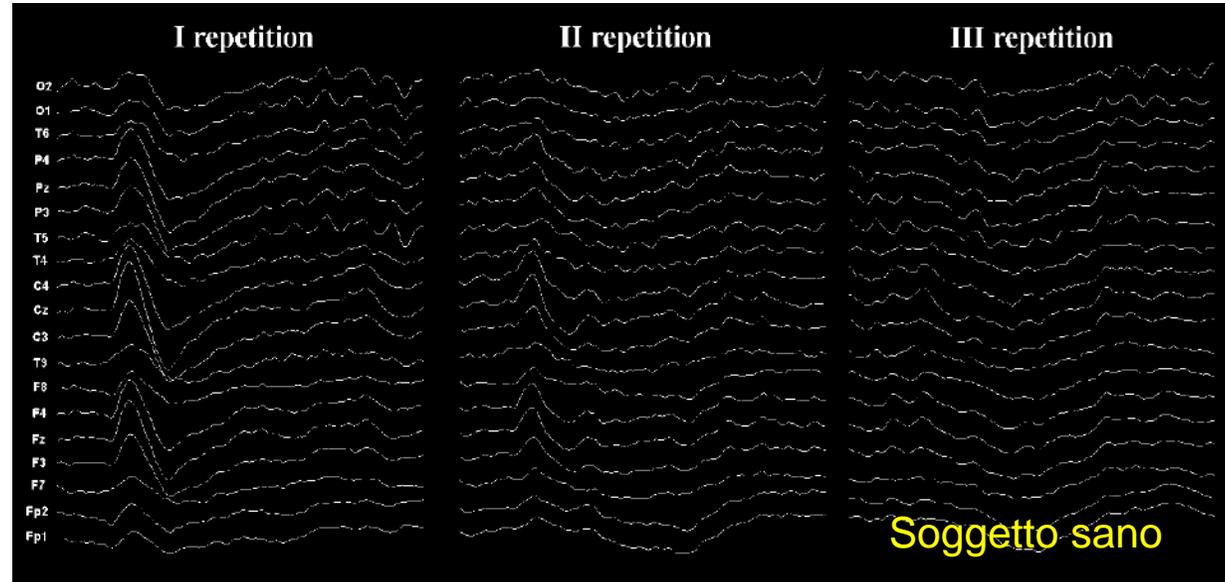
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PAIN

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Reduced habituation to experimental pain in migraine patients:
a CO₂ laser evoked potential study

M. Valeriani^{a,b,*}, M. De Tommaso^c, D. Restuccia^a, D. Le Pera^{a,d}, M. Guido^e, G.D. Iannetti^e,
G. Libro^e, A. Trini^e, G. Di Trapani^a, F. Pucca^e, P. Tonali^a, G. Cruccu^e



L'habituation del complesso di vertice si riduce sia nelle pazienti emicraniche che nelle donne non emicraniche

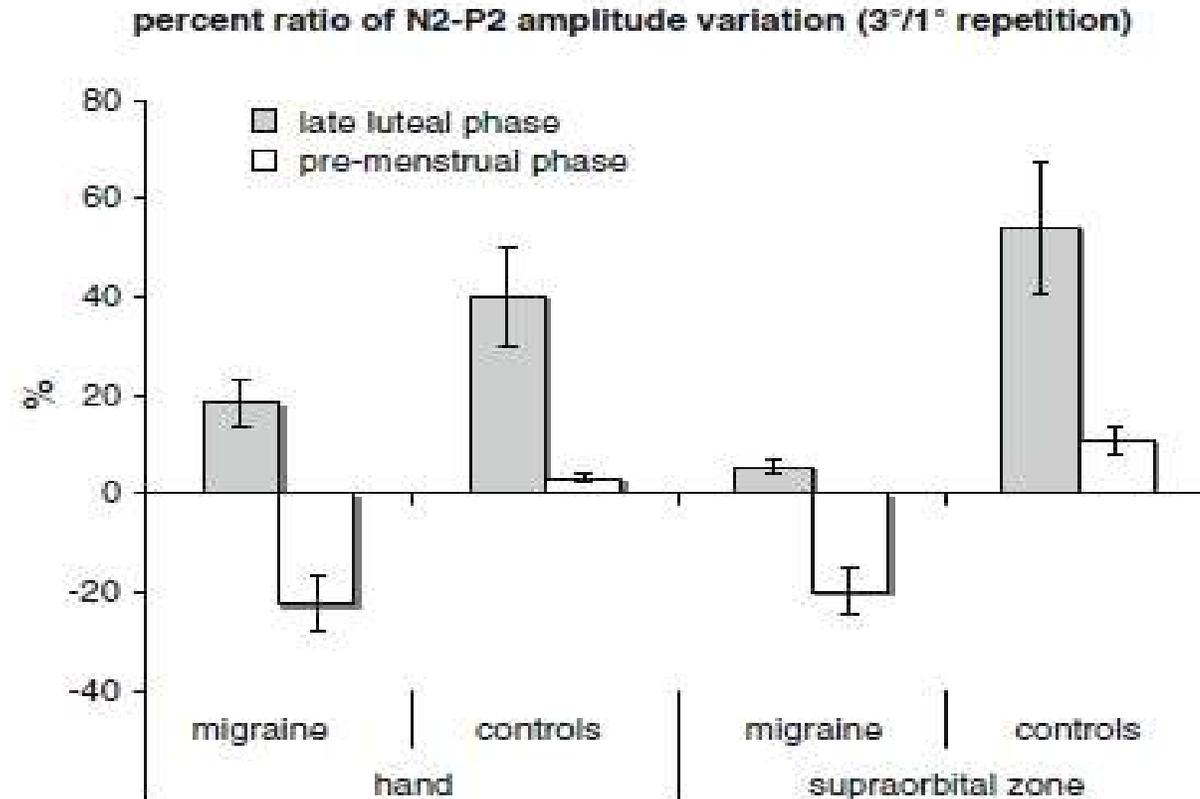


Fig. 4 Mean values and standard errors of percent ratio of amplitude variation, computed between the third and the first series of N2-P2 complexes, in migraine patients (no. 9) and controls (no. 10)

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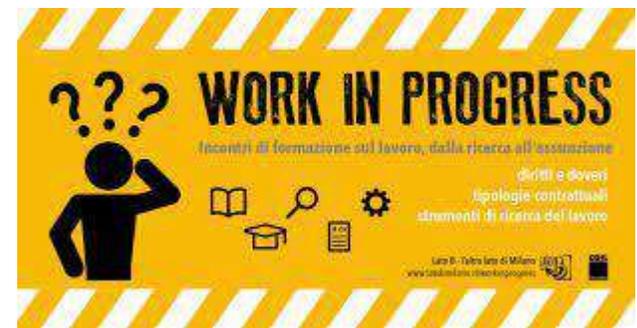
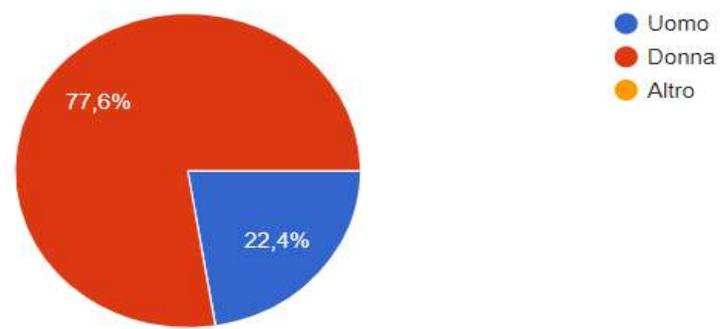
RESPONSABILI SCIENTIFICI:
PROF.SSA MARINA DE TOMMASO
PROF.SSA SARA INVITTO



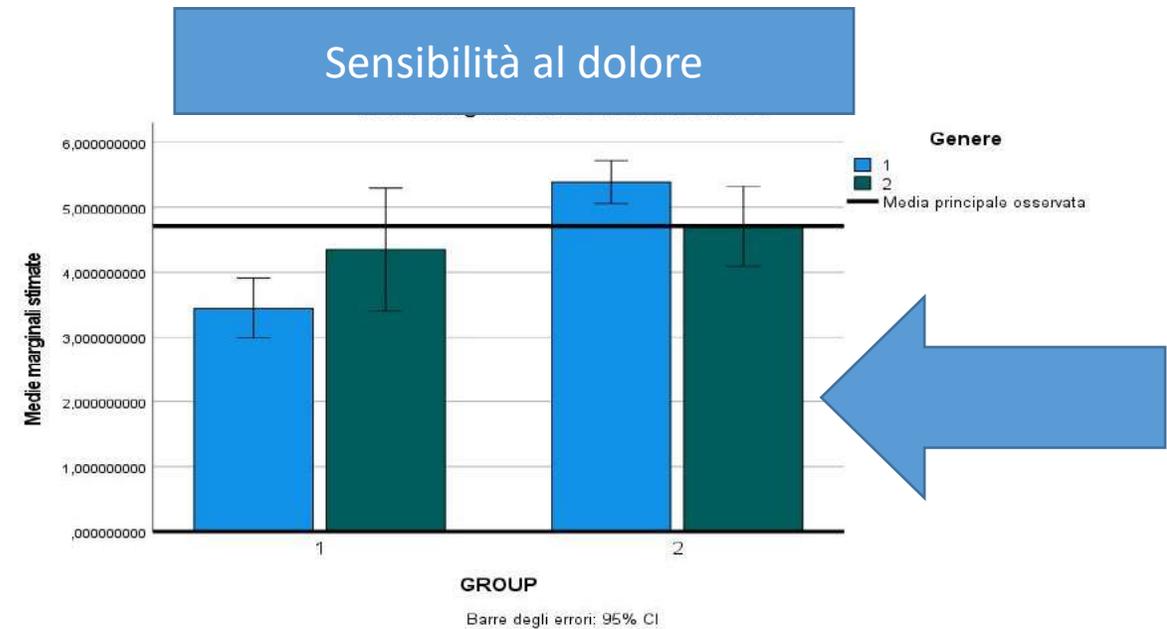
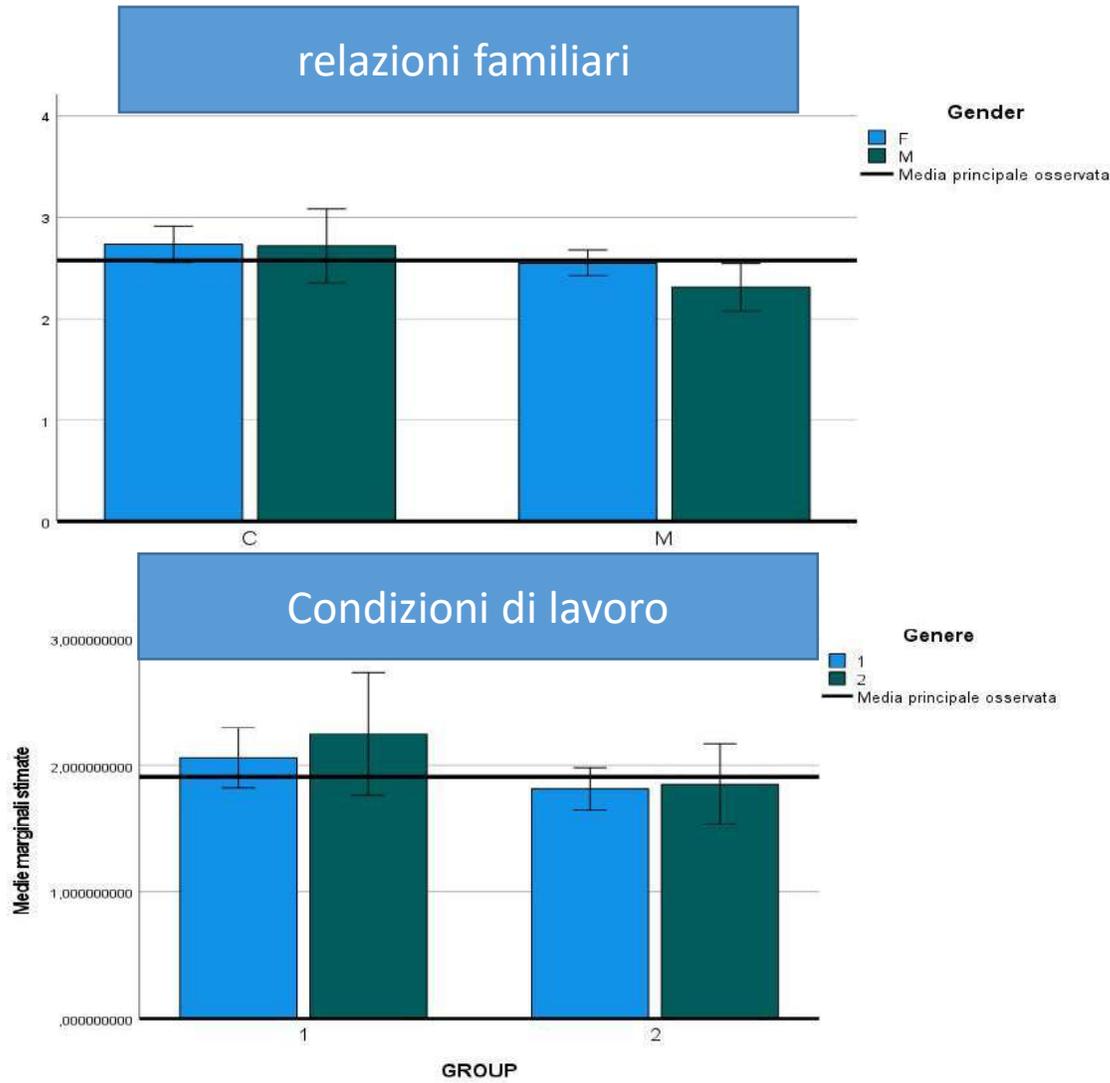
I RIUNIONE DEL GRUPPO SISC-INTERSOCIETARIO
"Donne Contro Il Dolore"
DOLORE E DIFFERENZE DI GENERE: IL MODELLO EMICRANICO

Lo Stress in Ambito Familiare e Lavorativo nell'Emicrania : Risultati di un'intervista online in pazienti afferenti a Centri Cefalee Survey -Test Somministrati

- BAI Beck Anxiety Inventory (Beck et al.)
- BPQ Body Perception Questionnaire (Porges)
- RSQ Romance Quality Scale
- CTS Conflict Scale
- ERQ Questionario Regolazione Emotiva (Versione Italiana Balzarotti et al.)
- PSS Scala Stress Percepito (S. Cohen)
- Stress Lavoro Correlato
- Percezione del Dolore Emicranico/Fibromialgico

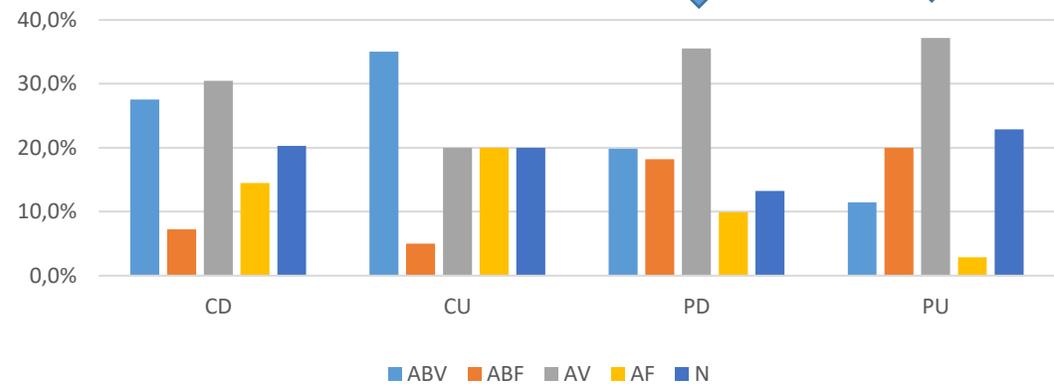


Non si evincono differenze di genere nel gruppo degli emicranici e dei controlli per le variabili considerate, ma solo differenze relative alla patologia. La percezione del dolore è più elevata nei pazienti emicranici, con lieve prevalenza femminile

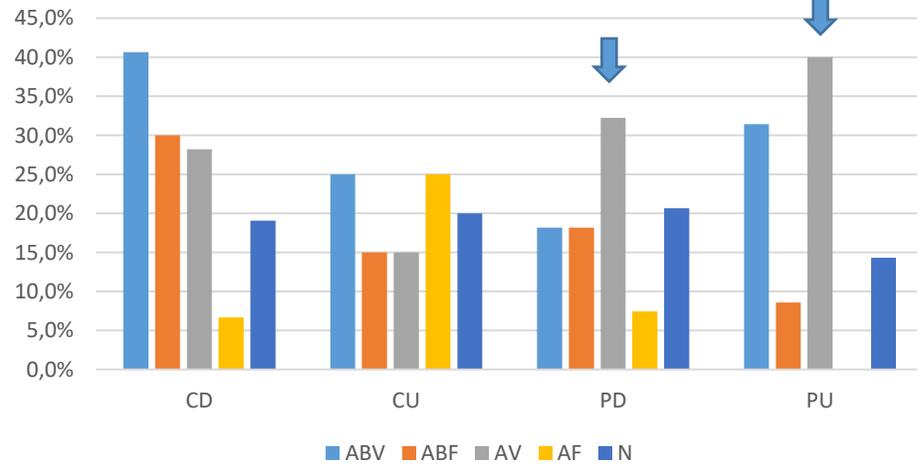


Nei rapporti familiari, l'emicrania fa la differenza

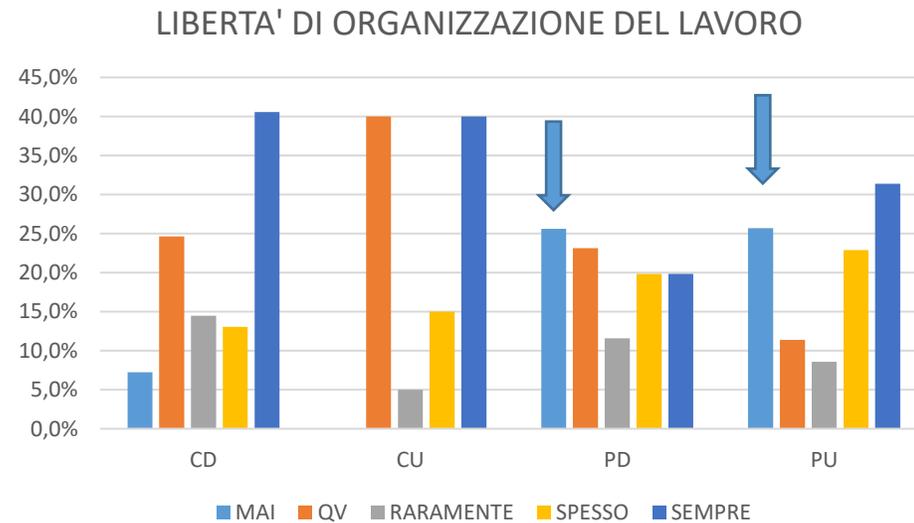
LITIGIOSITA' NELLA COPPIA



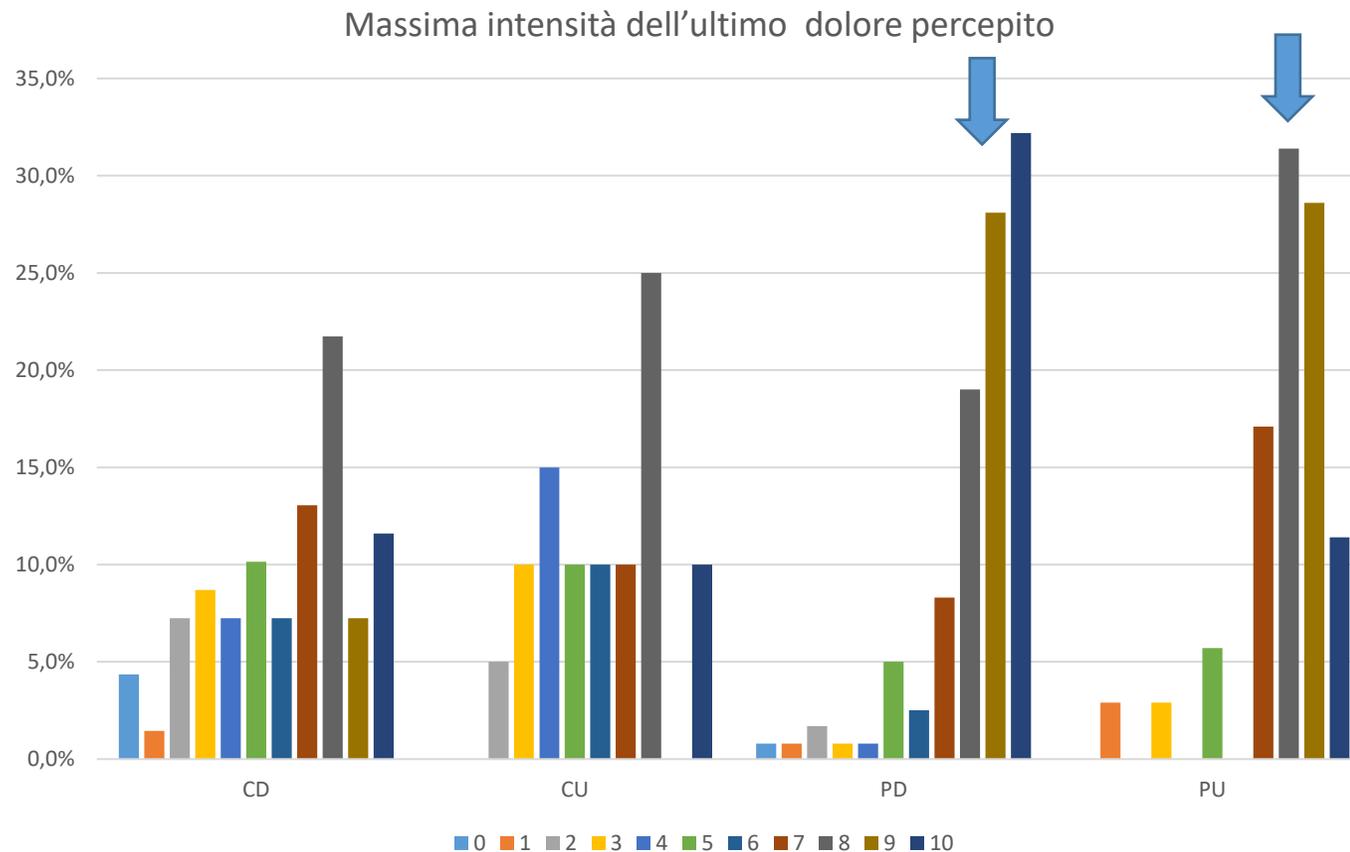
CONFLITTUALITA' DI COPPIA



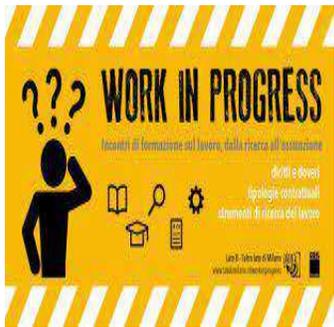
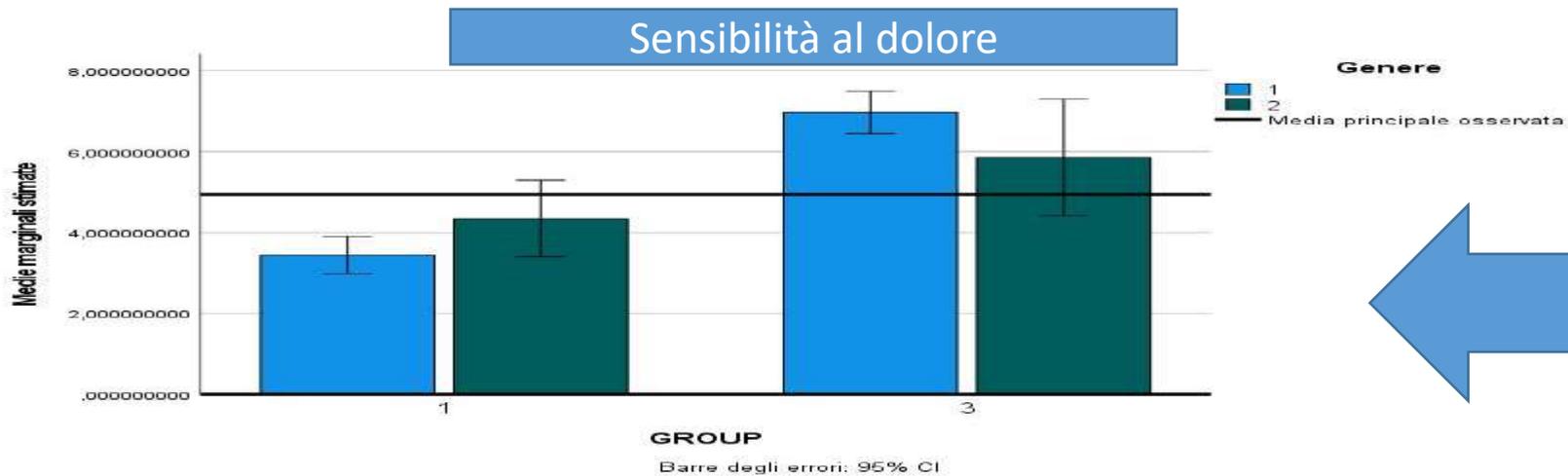
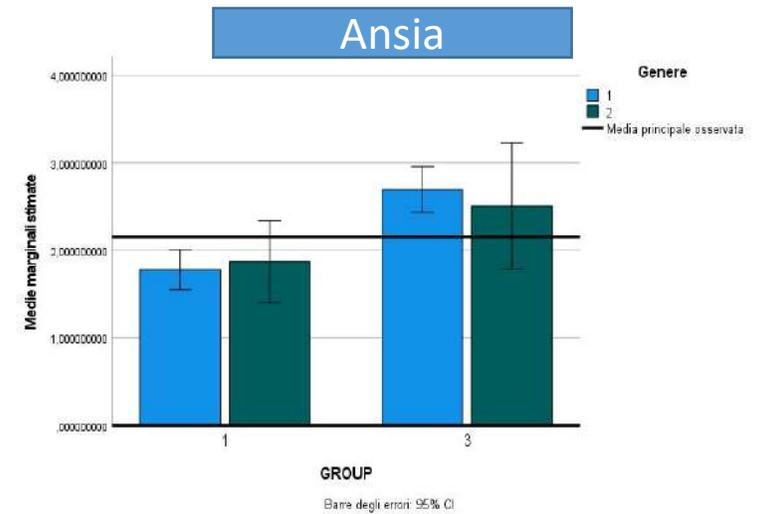
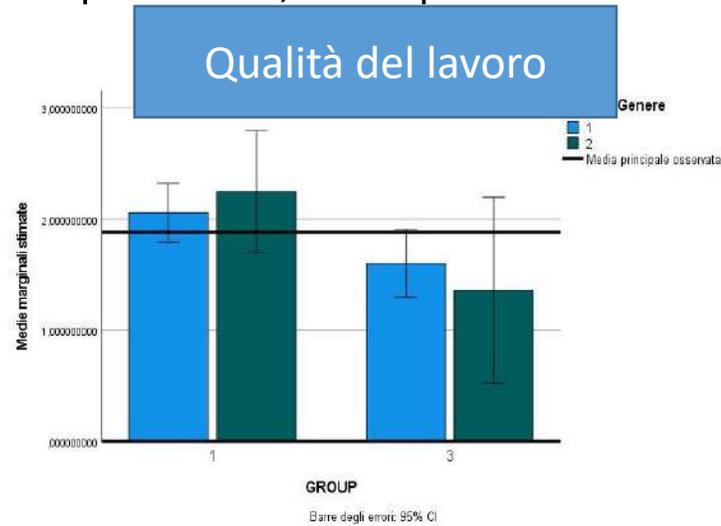
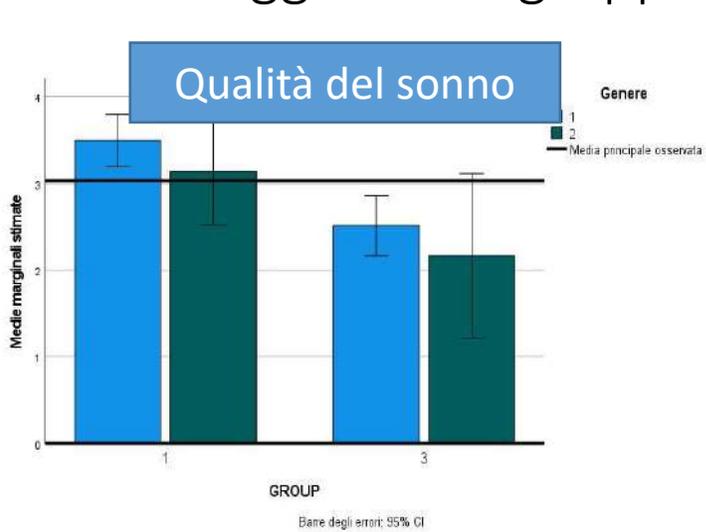
Nello stress da lavoro, l'emicrania fa la differenza



Nella sensibilità al dolore, l'emicrania fa la differenza
(ma le donne emicraniche sentono più dolore)



Nella fibromialgia, esempio di dolore cronico, gli indicatori di stress sono significativamente peggiori nel gruppo dei pazienti. La sensibilità al dolore è maggiore nel gruppo dei pazienti, con prevalenza femminile



Riflettiamo.....



- Sesso femminile  aumentata percezione del dolore indipendentemente dalla patologia e dalla specifica condizione ormonale
- La prevalenza di sesso nelle singole condizioni di dolore cronico sembra dipendere prevalentemente da variabili di tipo genetico e fattori ambientali comuni nei 2 sessi che condizionano comunque fenotipi simili
- Non c'è evidenza che il sesso femminile predica una evoluzione peggiore in tutte le condizioni di dolore cronico

La prevalenza di sesso nelle singole condizioni di dolore cronico sembra dipendere prevalentemente da variabili di tipo genetico e fattori ambientali comuni nei 2 sessi che condizionano comunque fenotipi similari .

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The Epidemiology of Migraine Headache in General Population of Tehran, Iran

Behnam Rabiee^a Atefeh Zeinoddini^a Ramin Kordi^a Masud Yunesian^b
Payam Mohammadnejad^a Mohammad A. Mansournia^a

The
1-year prevalence of migraine was 27.6% (36.7% in women and 21.6% in men) in Iranian general population,

[Iran Red Crescent Med J. 2016 October; 18\(10\):e40061.](#)

[doi: 10.5812/ircmj.40061.](#)

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Review Article

The Prevalence of Migraine in Iran: A Systematic Review and Meta-Analysis

Zeynab Farhadi,¹ Saeideh Alidoost,² Meysam Behzadifar,³ Roghayeh Mohammadibakhsh,⁴ Najmeh Khodadadi,⁵ Razieh Sepehrian,⁶ Rahim Sohrabi,⁷ Masood Taheri Mirghaed,⁸ Morteza Salemi,⁹ Hamid Ravaghi,⁶ and Masoud Behzadifar^{2,3,6,7}

Conclusions: The prevalence of migraine in Iran, which was estimated as 14%, was similar or even higher than that reported worldwide.



Un campione estratto dalla
UOC Neurofisiopatologia Universitaria
...esempio di prevalenza di genere....

