

Sommeil et Psychiatrie

Pr. Pierre Alexis GEOFFROY

↓ *Speaker honorarium:*







Biocodex, Bioprojet, Idorsia, Janssen-Cilag, Isis Medical, Jazz pharmaceuticals, NordicPharma, Pharmanovia, ProBTP.

↓ *Fees for consulting:*

Apneal, Arrow, Biocodex, Di&Care, Idorsia, Jazz pharmaceuticals, Myndblue, Petit Bambou, Pileje.

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Apneal, Di&Care, Idorsia, Janssen-Cilag, Myndblue.

Pierre Alexis Geoffroy¹ , Romain Roue², Sophie B. Sebille² , Julia Maruani³ , Estelle Taupinard⁴, Michel Lejoyeux³ , Anne Perozziello²  and Sibylle Mauries³ 

Population & Data Source



13,913 adult psychiatric inpatients



2021-2023



Paris Psychiatry Hospital Group- Health Data Warehouse



81% had chronic sleep disorders (CSD)

Main Findings

- ↑ Hospitalizations per patients
- ↑ Longer stay duration
- ↑ Seclusion
- ↑ Physical restraint
- ↑ Resistance to drugs
- ↑ Depression, Bipolar disorders, Anxiety disorders, SUD, etc

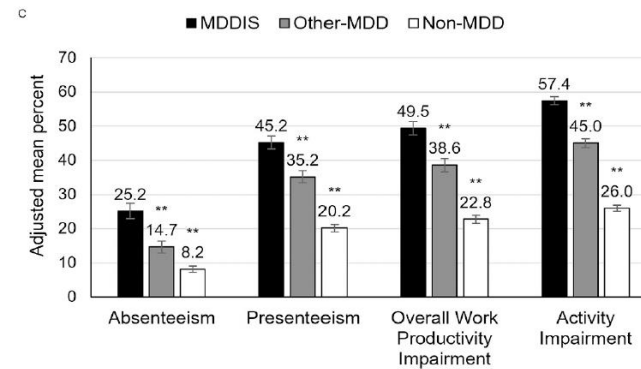
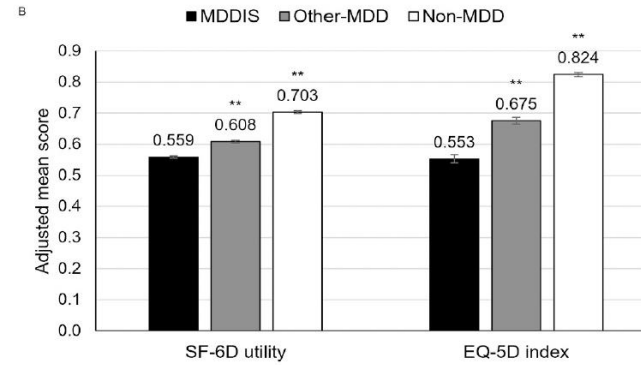
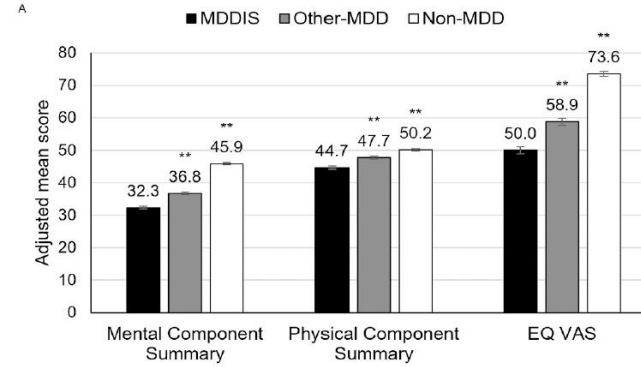
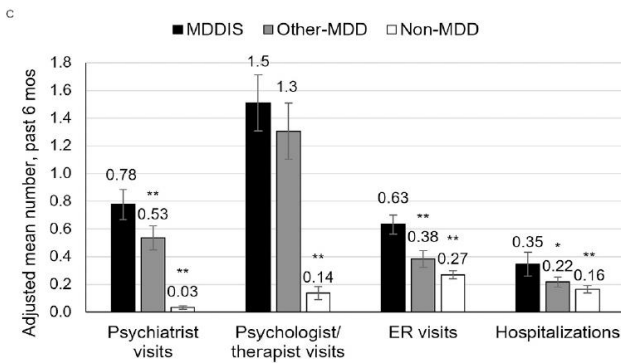
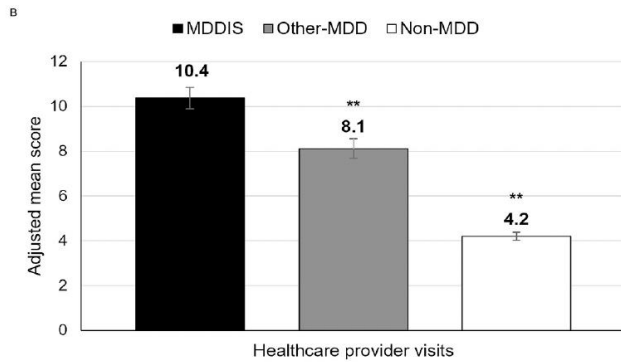
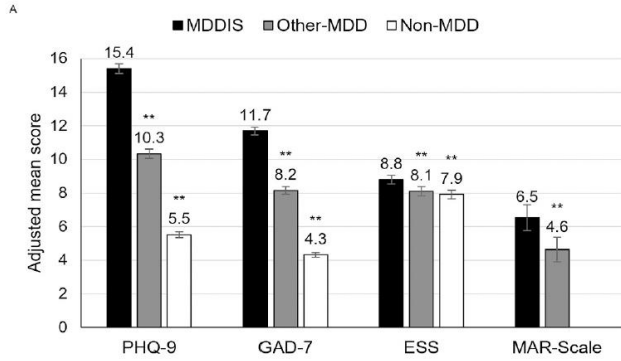
Population

Sleep disorders = ICD-10 codes, hypnotic prescriptions, or mentions in medical records.

CSD = Index of Length of Stays with Disorders (ILSD) >0.5,
No sleep disorders : ILSD=0

Comorbidities

- ↑ Psychiatric comorbidities
- ↑ Non-Psychiatric comorbidities
- ↑ Self-harm
- ↑ Psychoactive substance use
- ↑ Stress



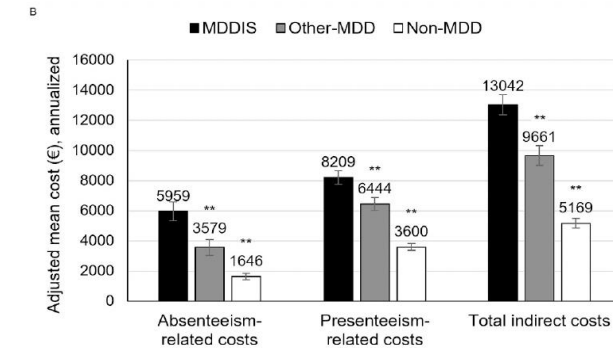
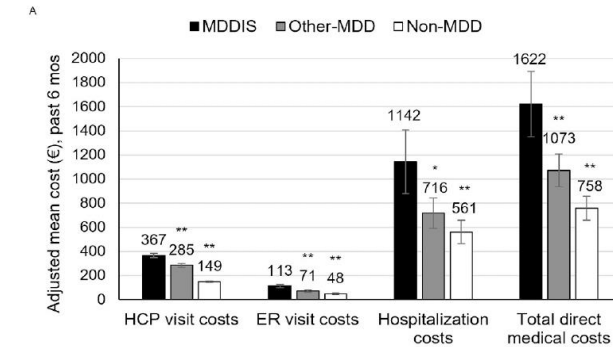
Research paper

Examining the burden of major depressive disorder with moderate-to-severe insomnia symptoms in five European countries

Jesper Gottorp Riise^a, M. Janelle Cambron-Mellott^b, Zhiheng Zhang^{c,*}, Véronique Huber^d, Alanna Pfau^c, Leiyu Yue^b, Nilanjana Dwibedi^c, Abisola Olopoenia^b, Pierre A. Geoffroy^{e,f,g}

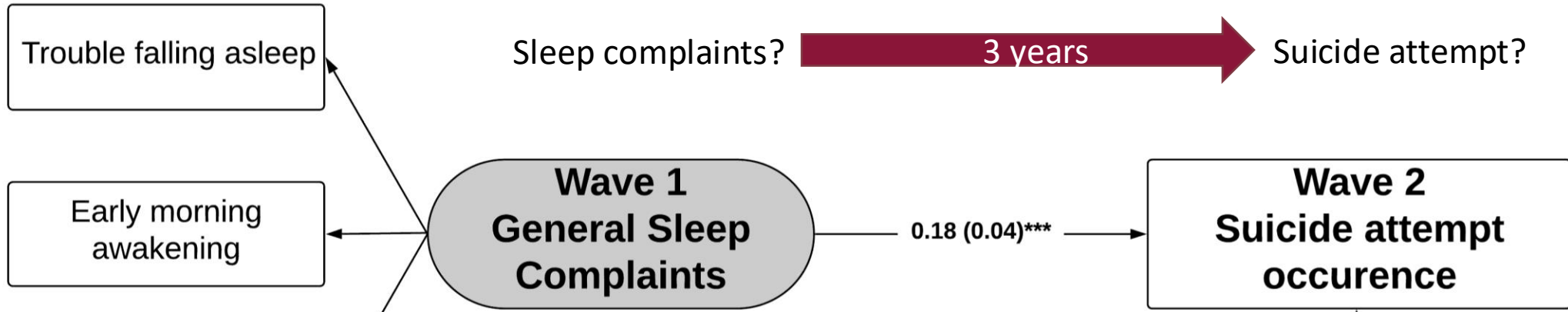
European (France, Germany, UK, Italy, Spain) 2020 National Health and Wellness Survey

1888 MDDIS, 3905 other-MDD, and 52,074 non-MDD participants





Sleep complaints predictive of suicidal behavior independently of any psychopathology



N Hoertel



H Peyre

Prospective 3-year study (NESARC) evaluating the impact of sleep complaints on the risk of suicide attempts (SA)

N=34,653 individuals representative of the general American population

Prevalence of SA over 3 years = 0.6% (SE=0.1, n=241).

Sleep complaints in TS vs noTS group: 59.9% vs 22.8%

- Difficulty falling asleep 46.6% vs 16.6%
- Early awakenings 38.9% vs 12.7%
- Hypersomnia 35.0% vs 10.7%

Wave 2 Suicide attempt occurrence

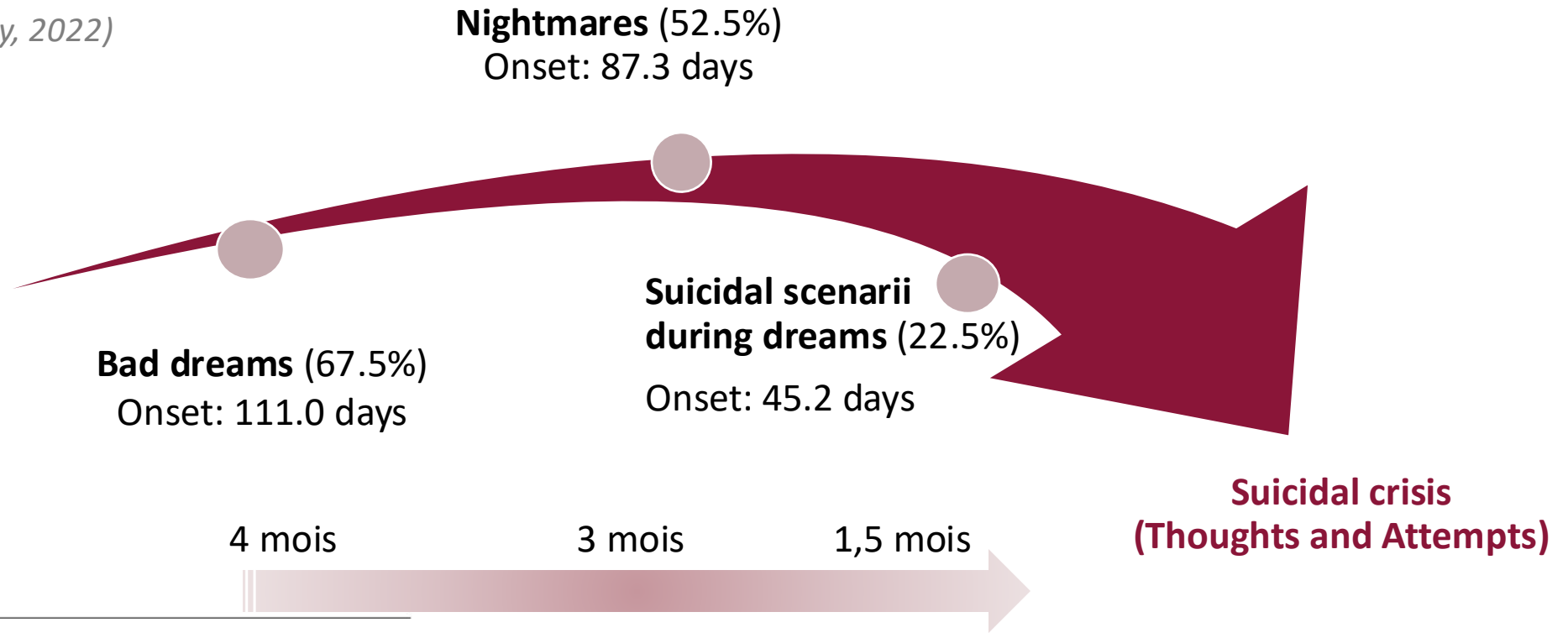
General Psychopathology Factor: 0.32 (0.04)***
 Prior suicide attempt: 0.99 (0.07)***
 Poverty: 0.22 (0.07)***
 Women: 0.18 (0.06)*
 Age:
 45-64: -0.30 (0.08)***
 65+: -0.69 (0.14)***

General psychopathology factor
 Prior history of suicide attempt
 Poverty
 Sex
 Age
 Race/ethnicity
 Marital status
 Obesity
 Number of stressful life events

* p-value<0.0125
 ** p-value<0.005
 *** p-value<0.001

Pierre A. Geoffroy, MD, PhD^{a,b,c,d,*}; Rodolphe Borand, MD^a; Marine Ambar Akkaoui, MD, MSc^{e,f}; Séverine Yung, MD^a; Yasmine Atoui, MD^a; Emeline Fontenoy, MD^a; Julia Maruani, MD, MSc^{a,c}; and Michel Lejoyeux, MD, PhD^{a,b,c}

(Geoffroy PA et al, J Clin Psy, 2022)



DOI: 10.1111/bdi.13325

LETTERS

BIPOLAR DISORDERS WILEY

How are you dreaming? A very simple question to screen for suicide risk

80% of individuals with suicidal behavior (suicidal thoughts or attempts) had **altered dream content**

Nightmares are a very common experience in the general population, with approximately 35%–45% of individuals experiencing at least one nightmare per month.¹ Nightmare disorder, which is characterized by repeated nightmares causing significant distress or impairment, is less frequent but still common, affecting 2%–8% of the general population.¹ Nightmares disorders are highly associated with

DATA AVAILABILITY STATEMENT

Data sharing is not applicable to this article as no new data were created or analyzed in this study.

Pierre Alexis Geoffroy^{1,2,3,4}

Prediction of mental health risk in adolescents

Received: 16 September 2024

Elliot D. Hill^{1,2}✉, Pratik Kashyap³, Elizabeth Raffanello³, Yun Wang⁴,
Terrie E. Moffitt^{5,6,7}, Avshalom Caspi^{5,6,7}, Matthew Engelhard^{1,2,8} &
Jonathan Posner^{3,8}

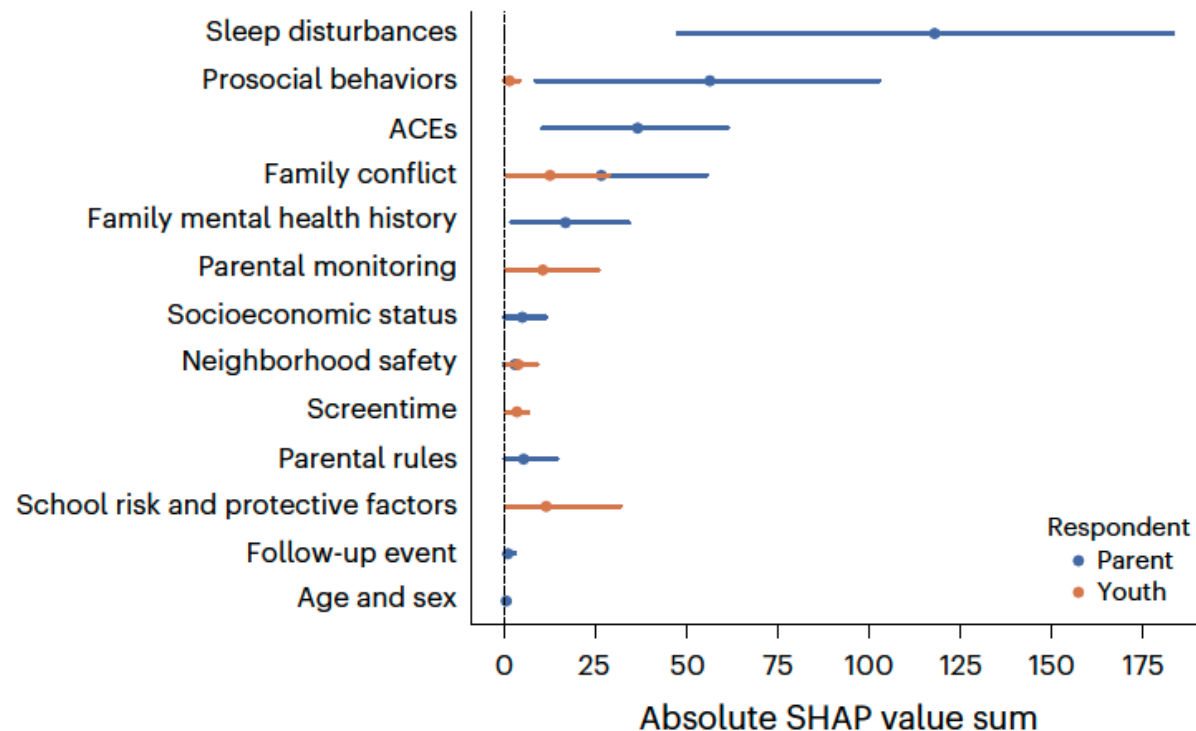
Accepted: 3 February 2025

Published online: 05 March 2025

Prospective prediction of mental health risk in adolescence can facilitate early preventive interventions. Here, using psychosocial questionnaires and neuroimaging measures from over 11,000 children in the Adolescent Brain and Cognitive Development Study, we trained neural network models to stratify general psychopathology risk. The model trained on current symptoms accurately predicted which participants would convert into the highest psychiatric illness risk group in the following year (area under the receiver operating characteristic curve = 0.84). The model trained solely on potential etiologies or disease mechanisms achieved an area under the receiver operating characteristic curve of 0.75 without relying on the child's current symptom burden. **Sleep disturbances emerged as the most influential predictor of high-risk status, surpassing adverse childhood experiences and family mental health history. Including neuroimaging measures did not enhance predictive performance.** These findings suggest that artificial intelligence models trained on readily available psychosocial questionnaires can effectively predict future psychiatric risk while highlighting potential targets for intervention. This is a promising step toward artificial intelligence-based mental health screening for clinical decision support systems.



Predictor category





Sleep disturbances and incident risk of major depressive disorder in a population-based cohort

Geoffroy Solelhac^{a,*}, Théo Imler^a, Marie-Pierre F. Strippoli^b, Nicola Andrea Marchi^a, Mathieu Berger^a, Jose Haba-Rubio^{a,c}, Tifenn Raffray^{a,c}, Virginie Bayon^a, Anne Sophie Lombardi^a, Setareh Ranjbar^b, Francesca Siclari^{a,i,j}, Peter Vollenweider^h, Pedro Marques-Vidal^h, Pierre-Alexis Geoffroy^{d,e,f}, Damien Léger^g, Aurélie Stephan^{a,1}, Martin Preisig^{b,1}, Raphaël Heinzer^a

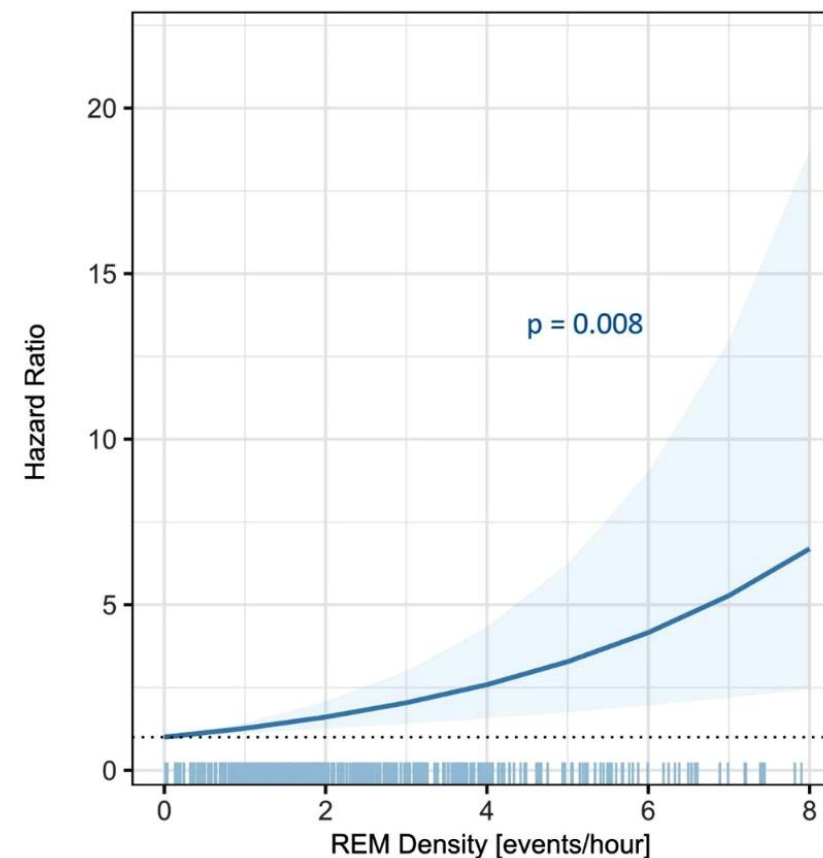
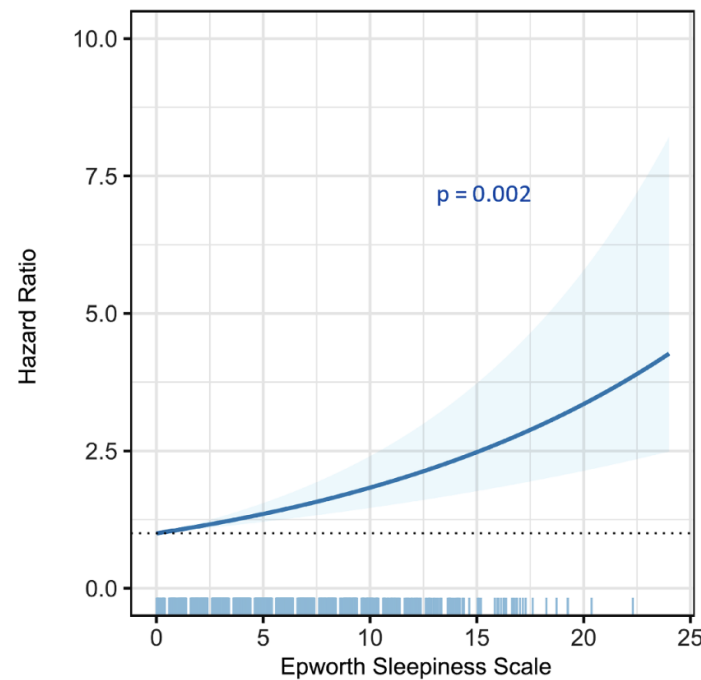
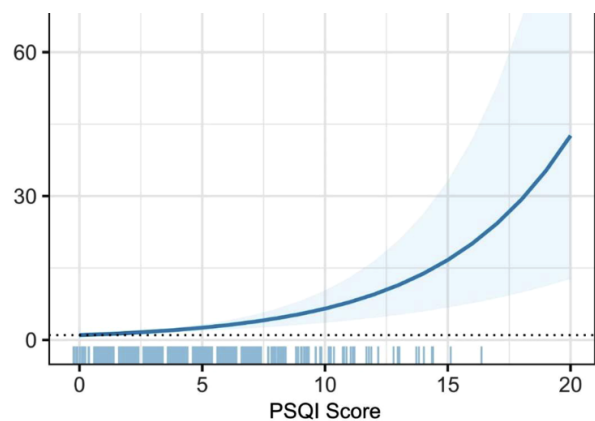
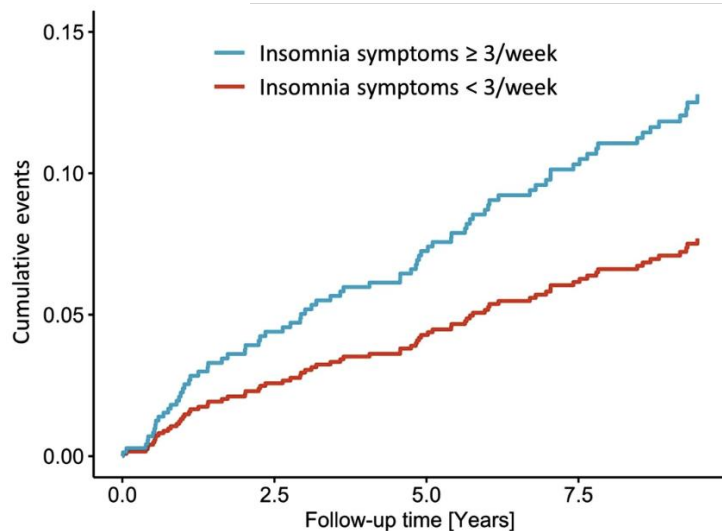
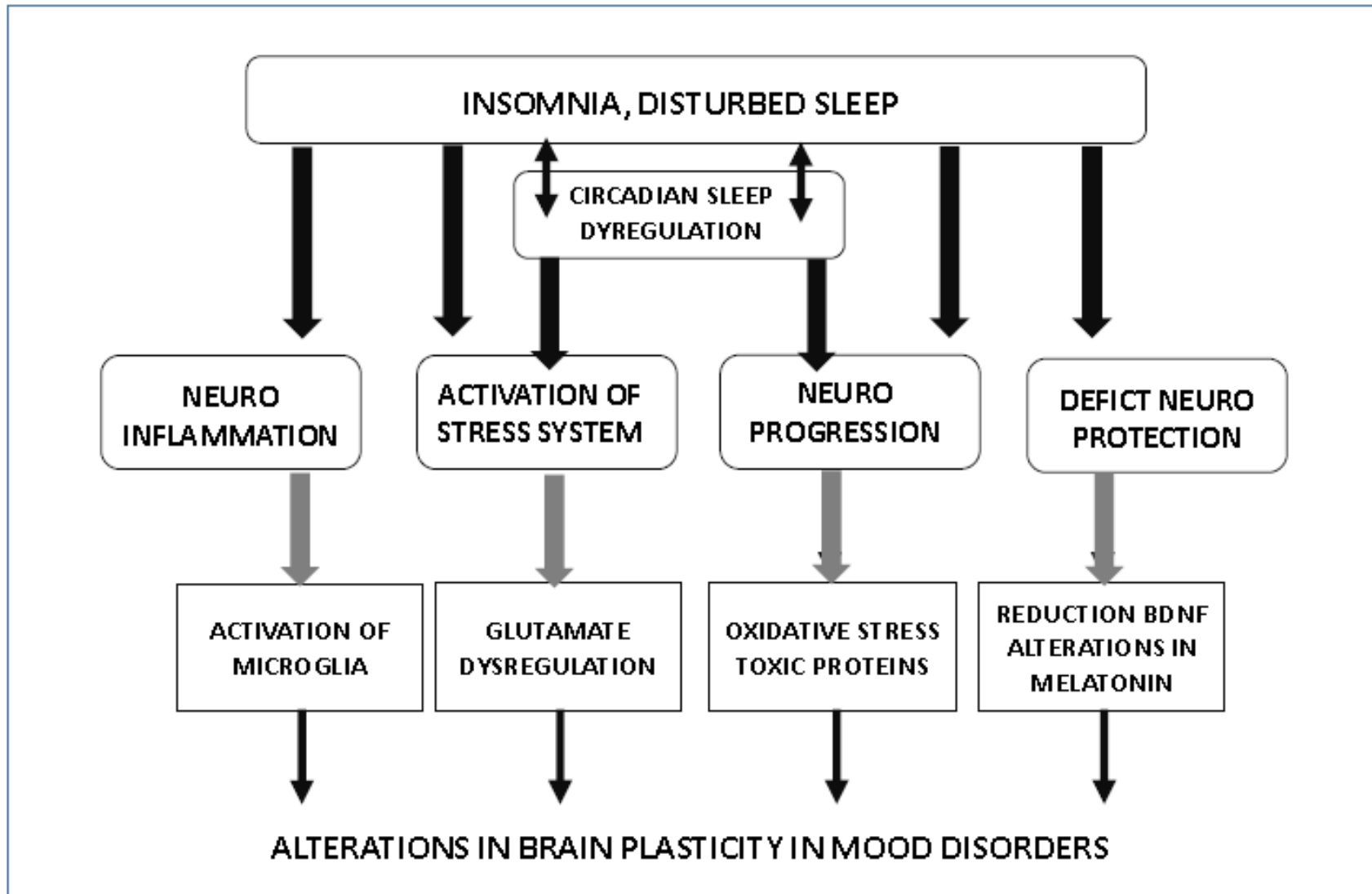


Fig. 6. Association between major depressive disorder incidence risk and rapid eye movements (REM) density in men. Data are multiple-adjusted Cox proportional hazards regression models showing simulated hazard ratio values with 95 % confidence intervals. Distribution of hazard ratio values are shown as a function of PSQI score and are derived from 1000 simulations using the approach described by King et al. (2000) (King et al., 2000). Models are adjusted for age, sex, and current smoking status. Distribution of REM density is shown on the rug plot along the x-axis. The p -value shown relates to the hazard ratio value in the original Cox proportional hazards regression model (1.270 [95 % confidence interval 1.064–1.516]).

(Solelhac G et al., Psychiatry Res, 2024)

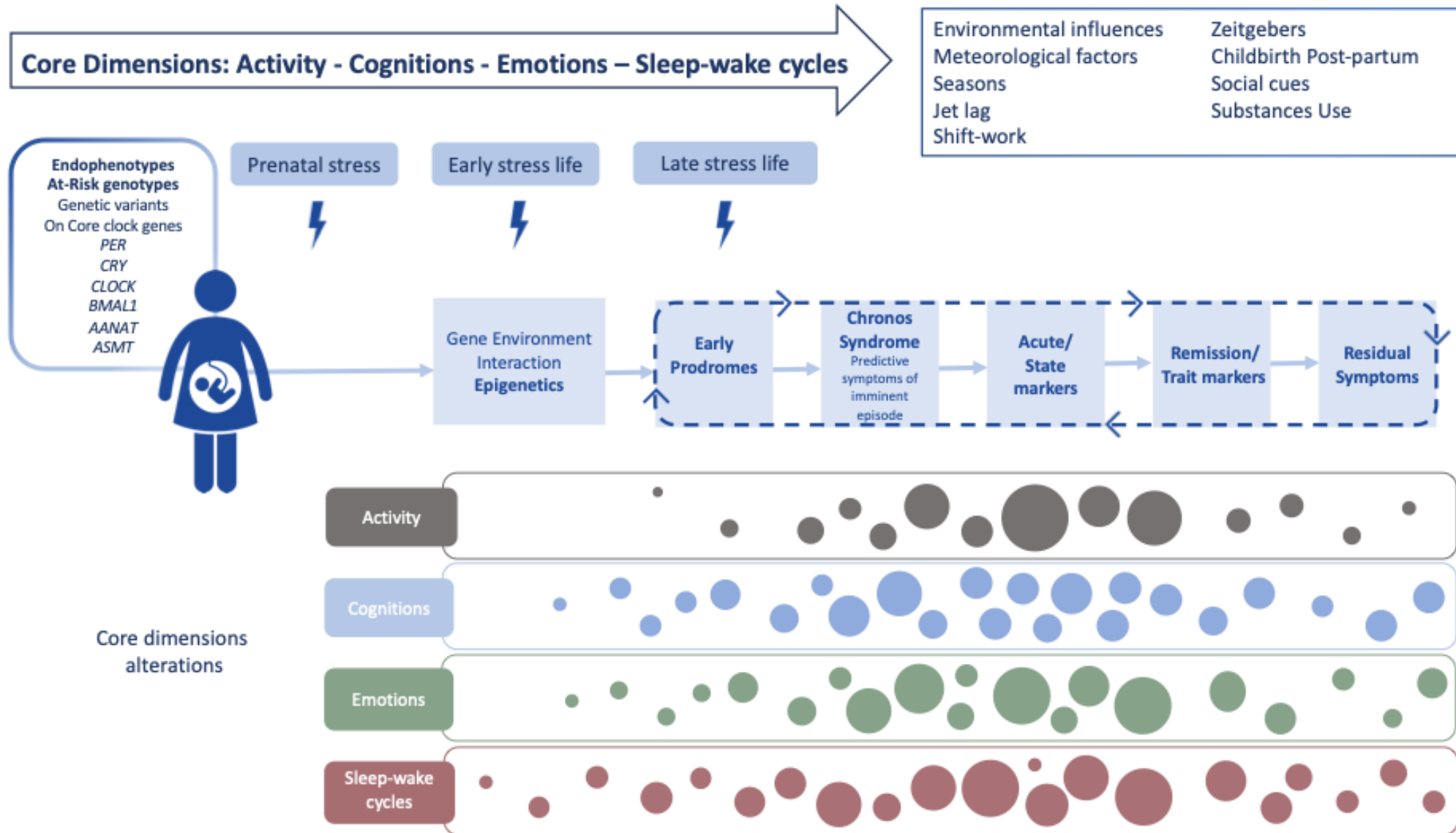


L Palagini



D Riemann

(Palagini L, Geoffroy PA, Riemann D, Current Medical Chemistry, 2022)



Dark skies before the storm: The Chronos syndrome

Pierre A. Geoffroy^{1,2,3,4} 



Chronos Syndrome / Depression

“The very first signs are usually a slightly restless night. I wake up and think ‘oh damn, something’s wrong,’ and typically in the morning I wake up with a knot in my stomach and I know I’m not doing well... Then come the cognitive problems...”

The classic early symptoms are loss of confidence, cognitive difficulties (I feel disconnected from the world, I don’t understand well what people are saying to me, it’s very unpleasant, I feel slowed down), I lose social ease, become a bit paranoid and just want to sleep. I go to bed around 11 p.m., sleep 9 hours instead of my usual 6.5 to 7 hours, but the sleep is of poorer quality. I sometimes wake up anxious during the night, and every morning it’s the same: I don’t feel well... and I’m a little scared; I’ve got this knot in my stomach...”

Chronos Syndrome / Mania

“The warning signs are when I get excited, I feel talkative, very reactive on social media, which normally isn’t like me—I don’t usually respond quickly or post much.

That’s often how it starts: I go out more, drink and smoke, and sleep less.

The decreasing sleep is my red flag, the sign that I need to slow down, otherwise I completely lose control and become unstoppable...”

Jean, 55 years old, BD1

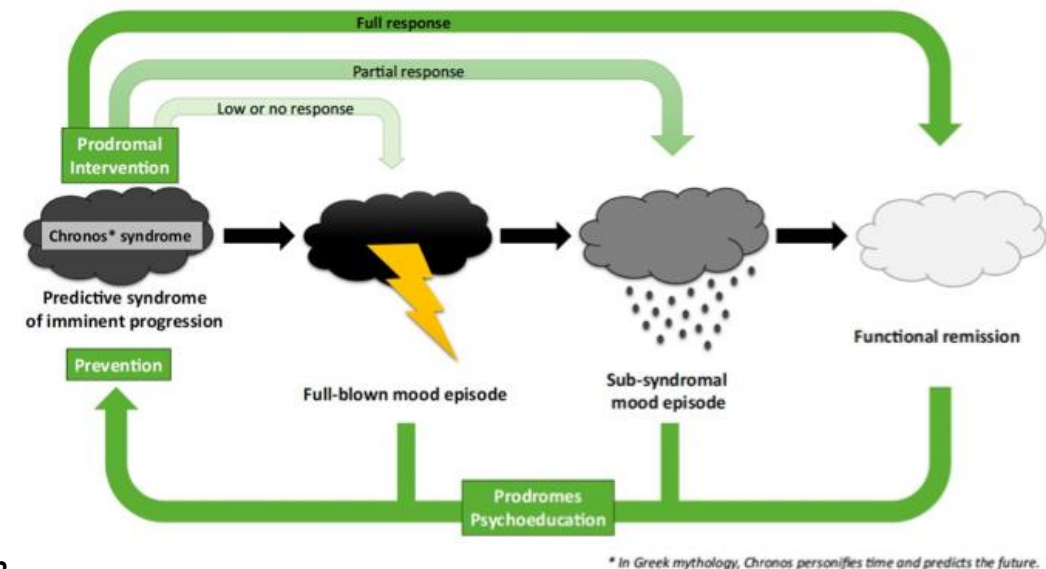
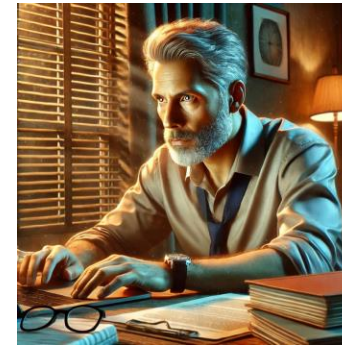
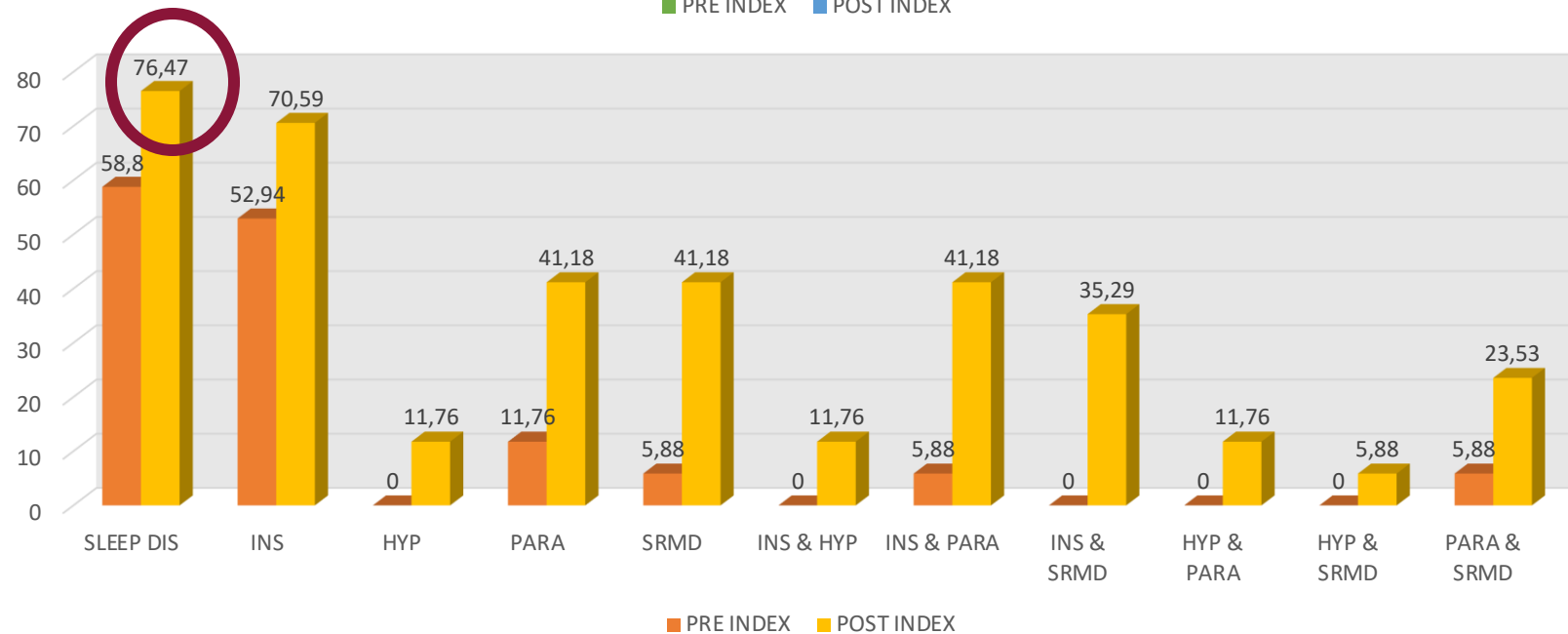
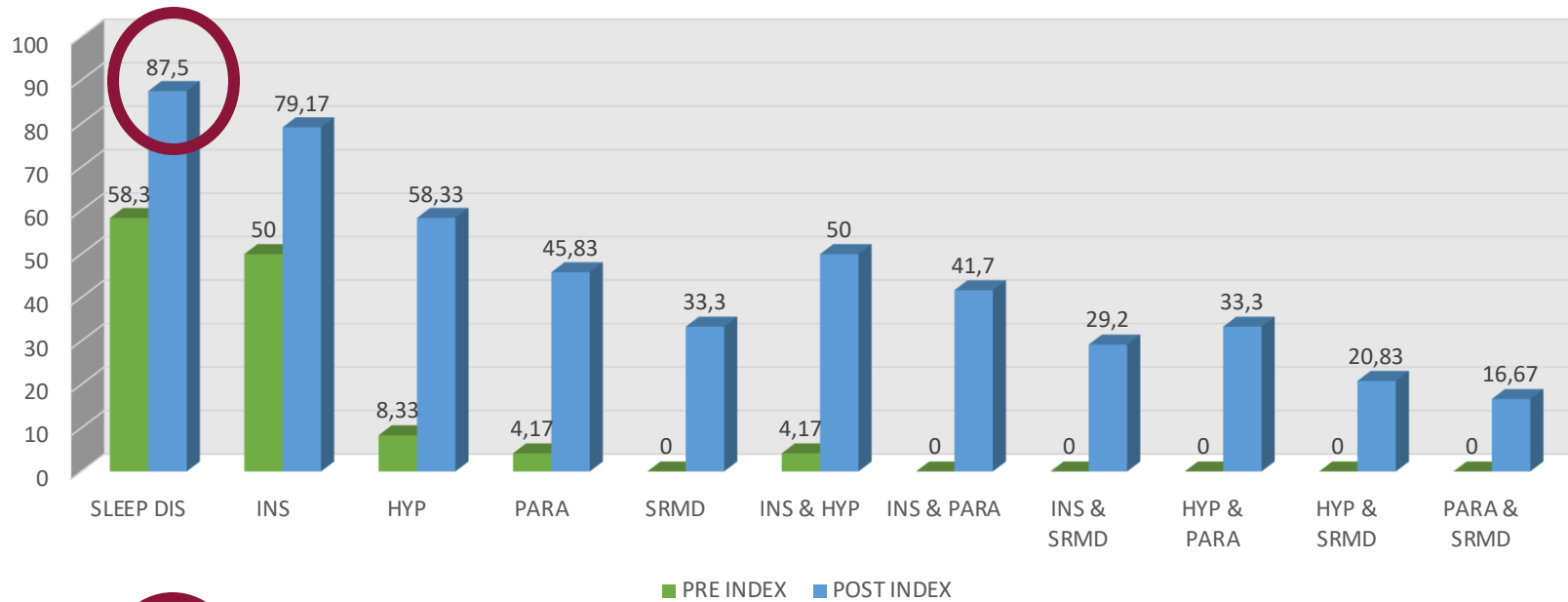


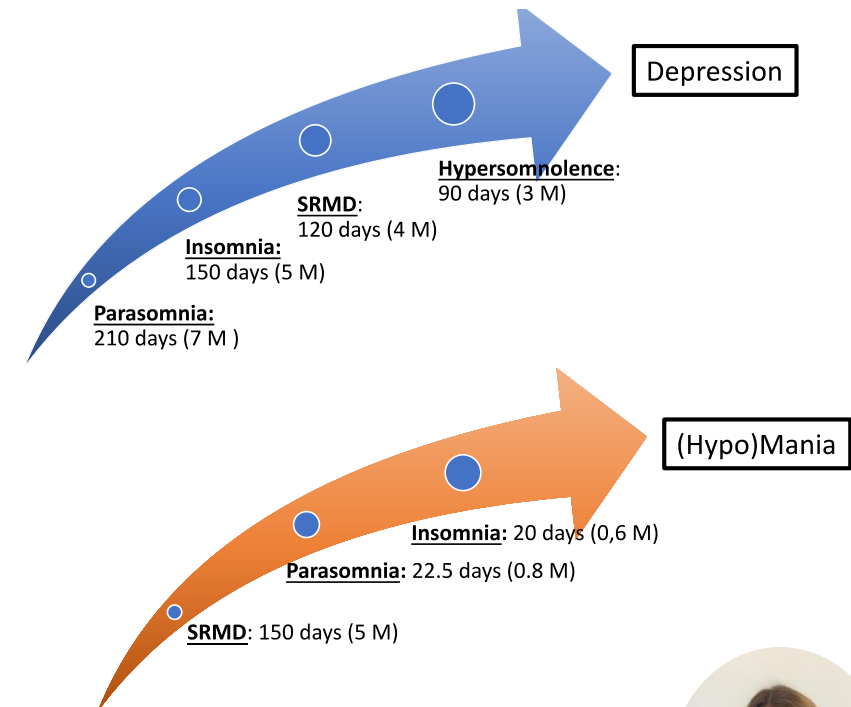
FIGURE 1 “Chronos syndrome”: a clinical syndrome reliably predicting an imminent transition to a mood episode. Addressing the “Chronos syndrome” during the prodromal phase could lead to more effective interventions, akin to anticipating dark skies before the storm, with the potential to limit or stop the progression to a full-blown mood disorder while empowering psychoeducation and prevention strategies.



(Geoffroy PA, Bipolar Disord, 2023)



87,5% report sleep disturbances prior to their **depressive episode**
76,5% prior to **Mania**



Louise Basquin

TABLE 1 Percentage and numbers of individuals experiencing each symptom of sleep alterations after interview.

	Yes (n, [%])	Onset (median in days [months])	Depression (24 patients with depression) (n, [%])	Mania (17 patients with (hypo)mania) (n, [%])	Onset for depression (median in days [months])	Onset for mania (median in days [months])
Insomnia	31 (75.6)	60 (2)	19 (79.2)	12 (70.6)	150 (5)	20 (0.7)
Nocturnal awakenings	19 (46.3)	42 (1.4)	13 (54.2)	6 (35.3)	60 (2)	20 (0.7)
Early awakenings	18 (43.9)	36 (1.2)	10 (41.7)	8 (47.1)	81 (2.7)	20 (0.7)
Onset insomnia	18 (43.9)	42 (1.4)	11 (45.8)	7 (41.2)	227,5 (7.6)	10 (0.3)
Poor quality sleep	16 (39.0)	150 (5)	12 (50)	4 (23.5)	180 (6)	95 (3.2)
Hypersomnolence disorders	16 (39.0)	90 (3)	14 (58.3)	2 (11.8)	90 (3)	80 (2.6)
Increased total sleep time	13 (31.7)	60 (2)	12 (50)	1 (5.9)	51 (1.7)	150 (5)
Sleepiness	8 (19.5)	45 (1.5)	6 (25)	2 (11.8)	45 (1.5)	80 (2.7)
Sleep inertia	6 (14.6)	105 (3.5)	6 (25)	0 (0.0)	105 (3.5)	0
Parasomnia	18 (43.9)	150 (5)	11 (45.8)	7 (41.2)	210 (7)	22.5 (0.8)
Epic dreaming	11 (26.8)	51 (1.7)	7 (29.2)	4 (23.5)	111 (3.7)	37.5 (1.3)
Nightmares	8 (19.5)	210 (7)	6 (25)	2 (11.8)	370 (12.3)	77 (2.6)
Bad dreams	8 (19.5)	120 (4)	6 (25)	2 (11.8)	135 (4.5)	77 (2.6)
Sleep talking	5 (12.2)	42 (1.4)	3 (12.5)	2 (11.8)	365 (12)	16.5
Suicidal scenarios among dreams	4 (9.8)	197,5 (6.5)	4 (16.7)	0 (0.0)	197,5 (6.5)	0
Hypnagogic hallucinations	3 (7.3)	150 (5)	2 (8.3)	1 (5.9)	165 (5.5)	2 (0.06)
Sleepwalking	2 (4.9)	1801,5 (60)	1 (4.2)	1 (5.9)	3600 (120)	3 (0.1)
Hypnopompic hallucinations	1 (2.4)	180 (6)	1 (4.2)	0 (0.0)	180 (6)	0
SRMD	15 (36.6)	150 (5)	8 (33.3)	7 (41.2)	120 (4)	150 (5)
Bruxism	9 (22.0)	150 (5)	3 (12.5)	6 (35.3)	60 (2)	257.5 (8.5)
Restless legs syndrome	7 (17.1)	180 (6)	5 (20.8)	2 (11.8)	180 (6)	95 (3.2)

Multiple therapeutic targets and highly personalized approaches +++

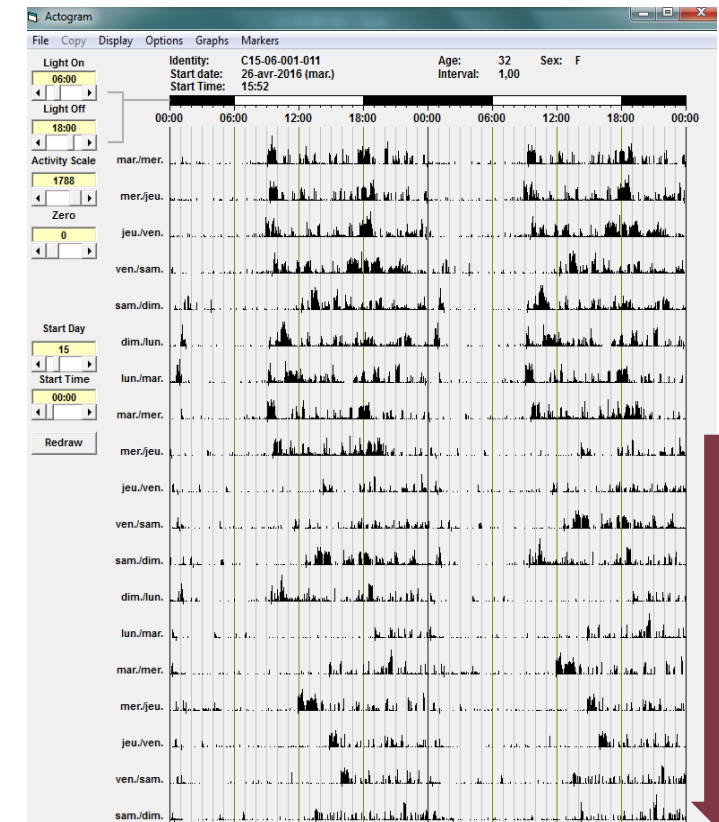
Note: Survival medians were calculated from Kaplan Meier curves.
Abbreviations: SRMD, Sleep-related movement disorders.

Exploring actigraphy as a digital phenotyping measure: A study on differentiating psychomotor agitation and retardation in depression

Julia Maruani^{1,2,3} | Sibylle Mauries^{1,2,3} | Ferial Zehani³ | Michel Lejoyeux^{1,2,3} | Pierre A. Geoffroy^{1,2,3}



Inserm
La science pour la santé
From science to health



Depressive relapse

TABLE 4 Multivariate analysis—Comparison of patients with major depressive episode (MDE) and psychomotor retardation (PMR) or agitation (PMA) regarding objective sleep and circadian rhythm markers assessed with actigraphy.

Variables	Groups mean (SD) or n (%)		Comparison PMR vs PMA	
	PMR (n = 58; 78.4%)	PMA (n = 16; 21.6%)	Z-score	p*
M10 average	10463.26 (±4069,59)	15046.75 (±3425.03)	4.12	0.002
Relative amplitude	0.80 (±0.13)	0.89 (±0.05)	277	0.027
Rest per 24 h%	46.62 (±9.13)	41.61 (±6.88)	-2.03	0.069
Interdaily stability	0.36 (±0.12)	0.48 (±0.09)	3.67	0.006
Intradaily variability	0.91 (±0.17)	0.75 (±0.14)	-2.94	0.012

Note: Bold values indicate a statistically significant difference with a p-value less than 0.05.

*p-value adjusted for current tobacco smoking and current alcohol use disorder.

A subtle clinical presentation, not easy to evaluate, with major therapeutic consequences!

Predictive biosignature of major depressive disorder derived from physiological measurements of outpatients using machine learning

Nicolas Ricka , Gauthier Pellegrin, Denis A. Fompeyrine, Bertrand Lahutte & Pierre A. Geoffroy

Seeking an **individual biosignature for depressive symptoms.**

Daily measurements over 6 months (medical device):
100 physiological markers physical activity, heart rate, heart rate variability, respiratory rate, sleep

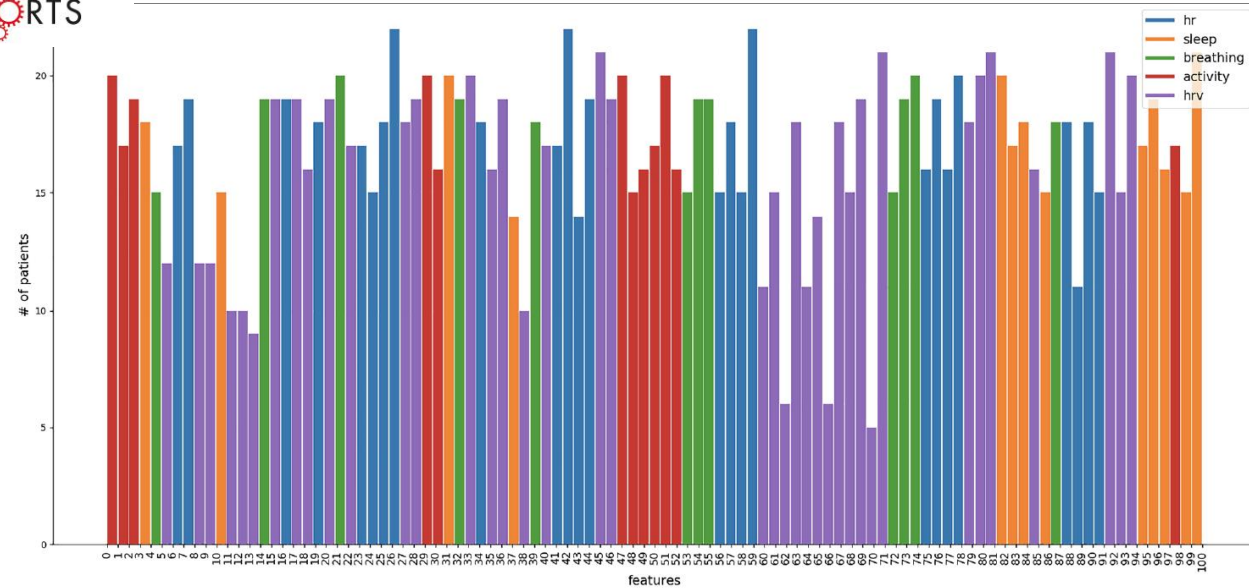


Figure 6. Histogram of features apparition in biosignatures. Histogram of unique features included in the biosignature of each patient. No feature was selected only for a single patient, and no feature was selected for all patients in the cohort. The colors represent the feature group to which the physiological variable belongs.

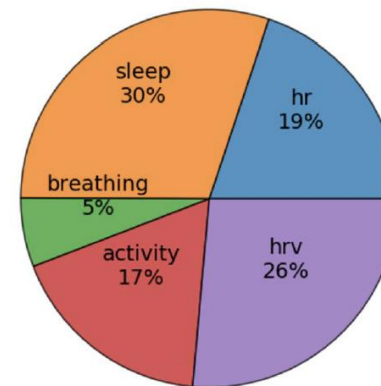
Method	2-class accuracy (TPR; TNR)	Predicted MADRS MAE
Constant (baseline) model	51.7%	9.63 (95% CI 7.83–11.43)
Optimistic model (prediction without physiological data)	74.4% (33.3%; 97.8%)	7.21 (95% CI 4.7–9.7)
Our model (baseline + deep learning approach)	86% (79%; 94%)	6.7 (95% CI 3.4–10.1)

Prediction for the next 3 months

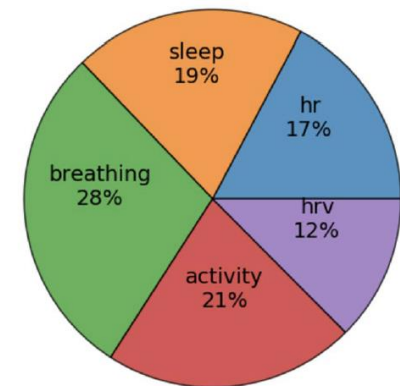
Sensitivity = 79%

Specificity = 94%

signature of patient 1



signature of patient 5





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Predictors of cognitive behavioral therapy for insomnia (CBT-I) effects in insomnia with major depressive episode

Julia Maruani^{a,b,c,*}, Emilie Stern^{c,d}, Charlotte Boiret^b, Jeanne Leseur^a, Alix Romier^{a,b}, Michel Lejoyeux^{a,b,c}, Pierre A. Geoffroy^{a,b,c,e,*}



J Maruani

Effectiveness of CBT-I in patients with a major depressive episode (MDE) versus Sleep Education (SE).

CBT-I compared to SE:

↓ significantly reduces the severity of insomnia: greater effect on early awakenings,
 ↑ increases sleep satisfaction,
 ↓ decreases the perception that sleep problems impact quality of life and daily functioning.

↑ increases the proportion of participants changing ISI category (CBT-I 54.6% vs. 23.8%)

↑ increases responses (CBT-I 30.3% vs. 0%)
 ↑ increases remissions (CBT-I 21.21% vs. 0%)

Greater reduction in the use of hypnotic medications +++

Greater improvement in depressive symptoms +++

Table 2
Comparison of the effects of CBT-I and Sleep education (SE) groups on sleep outcomes and on depression and anxiety symptoms in patients with major depressive episode (MDE) and insomnia.

	CBT-I (n = 30)	SE (n = 21)	Khi2, T- Student, Mann-Whitney U	P-value	Δ mean (Δ SD)	Effect size
<i>Sleep parameters, mean (SD)</i>						
ISI						
Δ ISI	6.30 (5.95)	1.95 (3.58)	3.01	0.004	4.35 (1.44)	0.84
Δ ISI1a	0.68 (1.06)	0.20(0.77)	248	0.114	2.35E-05	0.248
Δ ISI1b	0.67(1.16)	0.35(0.99)	273	0.275	4.95E-05	0.173
Δ ISI1c	0.86(1.38)	0.11(0.88)	190	0.022	1.000	0.375
Δ ISI2	1.09(0.28)	0.50(0.61)	225	0.045	1.000	0.318
Δ ISI3	1.03(1.36)	0.30(0.92)	219	0.047	1.000	0.317
Δ ISI4	1.00(1.24)	0.45(0.61)	234	0.089	1.39E-05	0.269
Δ ISI5	1.03(1.21)	0.20(0.70)	201	0.013	1.000	0.392
ISI response				0.005		
Yes	10 (30.3%)	0 (0%)				
No	23 (69.7%)	21 (100%)				
ISI remission				0.024		
Yes	7 (21.21%)	0 (0%)				
No	26(78.79%)	21(100%)				
ISI change of category				0.026		
Yes	18 (54.6%)	5 (23.8%)				
No	15 (45.4%)	16 (76.2%)				
PSQI						
ΔPSQI total score	2.37 (4.13)	1.33(2.55)	0.93	0.357	1.033 (1.11)	0.2645
ΔPSQI Sleep subjective quality	0.76(1.00)	0.24(0.77)	2.03	0.048	0.519 (0.25)	0.566
ΔPSQI sleep latency	0.27(1.01)	0.29(0.90)	339	0.88	3.51 e-5	0.0231
ΔPSQI Sleep duration	0.033(1.16)	0.29(1.19)	313	0.966	-3.97 e-5	0.008
ΔPSQI Daytime dysfunction	0.36(0.90)	0.24(1.04)	335	0.83	2.78 e-5	0.0346
ΔPSQI Sleep efficiency	0.40(1.57)	0.24(1.38)	274	0.41	4.09 e-5	0.009
ΔPSQI Sleep disturbance	0.21(0.70)	0.14 (0.57)	331	0.75	1.31 e-5	0.111
ΔPSQI Sleep promoting medication	0.75 (0.20)	-0.09 (0.18)	216	0.003	5.77 e-5	0.167
Excessive daytime sleepiness (ESS) ΔESS total score	1.09 (4.22)	-0.28(3.20)	1.27	0.208	1.379 (1.08)	0.3585
Home and Ostberg	-1.10 (5.52)	0.48 (9.03)	270	0.497	-1.00 (2.06)	0.12
Δ Home and Ostberg total score						
<i>Mood parameters, mean (SD)</i>						
ΔCESD	7.81(9.68)	2.38 (7.99)	2.13	0.038	5.43	0.599
ΔGAD7	3.13 (3.78)	1.700 (5.18)	227	0.148	2	0.243

Abbreviations: ISI, Insomnia severity index; PSQI, Pittsburgh Sleep Quality; ESS, Epworth Sleepiness Scale; CESD, Center for Epidemiologic Studies-Depression scale; GAD7, Generalized Anxiety Disorder-7.

Predictors of a decrease in ISI severity with CBT-I:

- Treatment-resistant depression (TRD)
- Seasonal fluctuations in depressive symptoms and sleep cycles throughout the year
- Daytime dysfunction (especially for insomnia related to early awakenings)

Factors of poor response to CBT-I:

- Shorter sleep duration
- Less improvement in daytime dysfunction and worries related to sleep disorders

Hypnotic, anxiolytic, antidepressant, and mood stabilizer medications does not predict a less favorable response to CBT-I.



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CLINICAL REVIEW

Cognitive behavioral therapy for insomnia in patients with mental disorders and comorbid insomnia: A systematic review and meta-analysis

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Efficacy on insomnia symptoms across psychiatric disorders

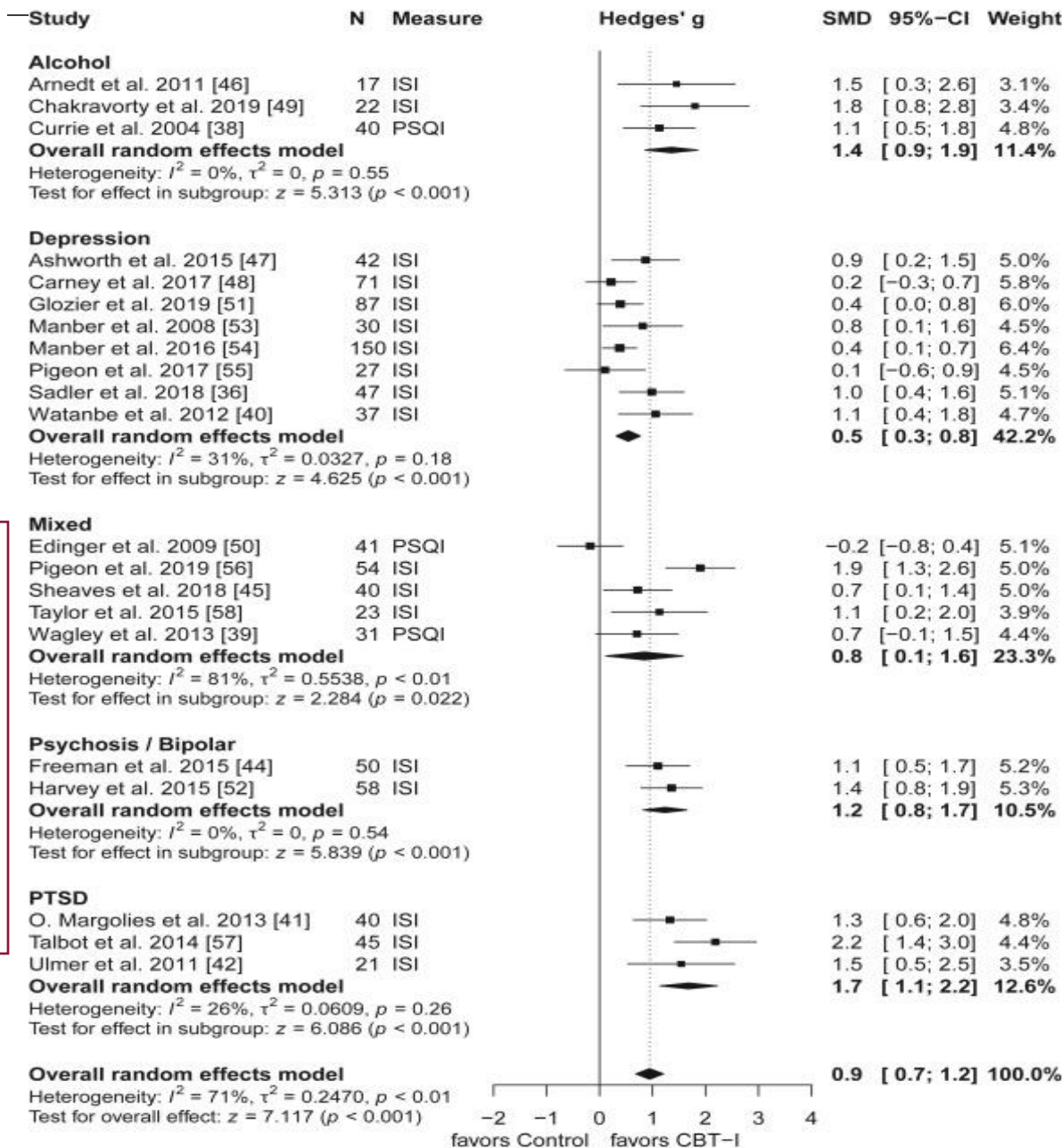
→ Moderate to large effect sizes

Efficacy also observed on psychiatric symptoms

CBT-I is effective in these populations

It can also be used as an adjunctive (add-on) treatment

→ To improve mental health outcomes in these disorders

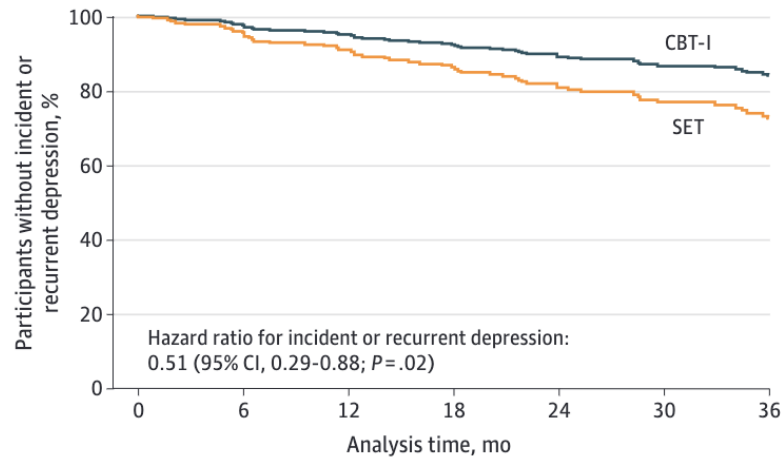




Prevention of Incident and Recurrent Major Depression in Older Adults With Insomnia A Randomized Clinical Trial

Michael R. Irwin, MD; Carmen Carrillo, MA, MHS; Nina Sadeghi, BS; Martin F. Bjurstrom, MD; Elizabeth C. Breen, PhD; Richard Olmstead, PhD

Figure 2. Time to Incident or Recurrent Depression Event by Treatment Group



No. at risk	0	6	12	18	24	30	36
CBT-I	156	129	123	116	92	88	81
SET	135	128	123	120	86	82	77

Older adults without depression but with insomnia were randomized to receive cognitive behavioral therapy for insomnia (CBT-I) or sleep education therapy (SET).

(Irwin et al., JAMA Psy, 2022)



CBT-I for prevention and early intervention in mental disturbances: A systematic review and meta-analysis

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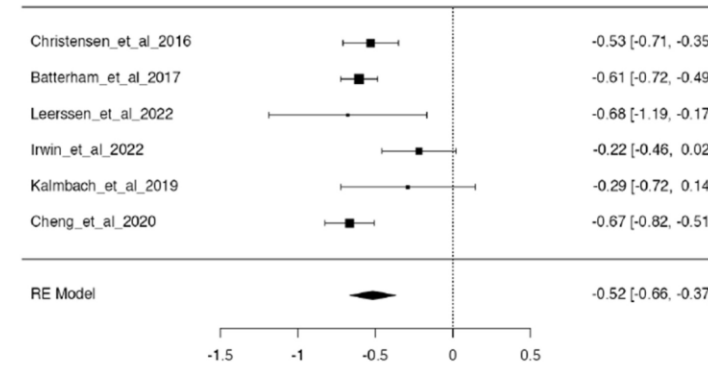


Fig. 2. A total of k = 6 studies were included in the analysis. CBT-I for insomnia was effective in reducing depressive symptoms ($\alpha = -6.8466$, $p < 0.0001$; RE Model = -0.5168 (95 % CI: 0.6648 to -0.3689)).

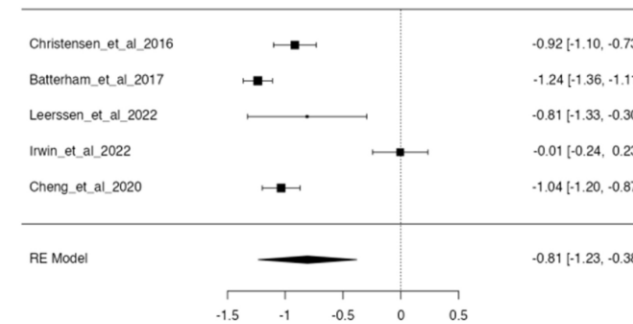


Fig. 4. Forest Plot for CBT-I on insomnia symptoms
CBT-I for insomnia was effective in reducing depressive and insomnia symptoms as well ($\alpha = -3.7126$, $p = 0.0002$; RE Model = -0.8074 (95 % CI: 1.2336 to -0.3811); k = 5).

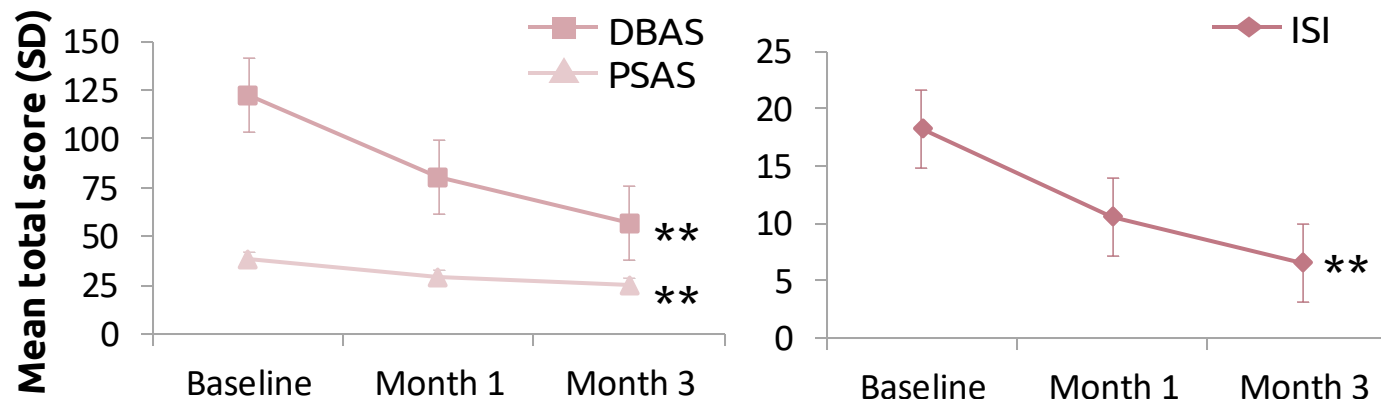
(Palagini et al., Sleep Med, 2024)



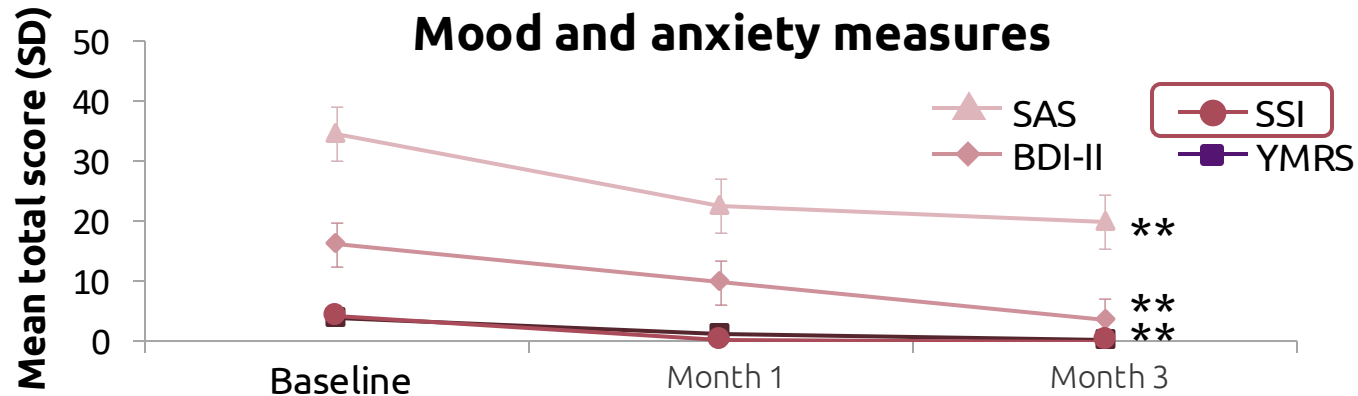
Study characteristics

Study design	Naturalistic observational, uncontrolled
Female (%)	54
Mean age (years ± SD)	60.0 ± 13.6
Observation period	1 and 3 months
Patients	N = 66
Daridorexant dose	50 mg
Sedative hypnotic use disorder, n (%)	24 (36.3)
Panic disorder (with anxiety), n (%)	30 (45.4)
Unipolar/bipolar depression, n (%)	32 (48.4)

Insomnia measures and insomnia severity



Mood and anxiety measures



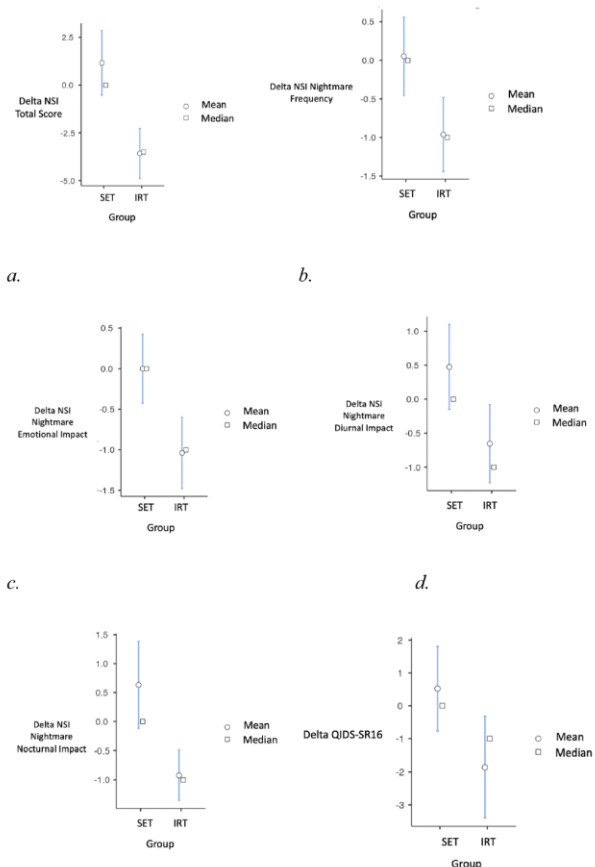
Analysis of variance; **P<0.001. N=66 at baseline and Month 1 only, at Month 3 N=60.

BDI, Beck Depression Inventory; DBAS, Dysfunctional Beliefs and Attitudes About Sleep Scale; ISI, Insomnia Severity Index; PSAS, Pre-Sleep Arousal Scale; SAS, Self-Rating Anxiety Scale; SD, standard deviation; SSI, Scale for Suicidal Ideation; YMRS, Young Mania Rating Scale.



Imagery Rehearsal Therapy (IRT) is associated with reduced nightmare severity and depressive, anxiety and suicidal symptoms in adults with Major Depressive Episode

Julia MARUANI^{a,b,c,*}, Nathan MARTINS^{a,b,c}, Emmanuelle CLERICI^c, Michel LEJOYEUX^{a,b,c}, Pierre A. GEOFFROY^{a,b,c,*}



Imagery Rehearsal Therapy (IRT) in Major Depressive Episode

Adults with:

- Major Depressive Episode
- Nightmare Disorder
- High suicidal risk



Imagery Rehearsal Therapy (IRT)

4 weekly group sessions



Rescripting nightmares while awake

Sleep Education Therapy (SET)
(waitlist control)

Main Outcomes

↓ Nightmare severity

- Frequency
- Emotional impact
- Nocturnal impact
- Daytime impact

↓ Depressive symptoms

↓ Anxiety symptoms

↓ Suicidal ideation

Moderators:

- ✓ Treatment-resistant depression → greater emotional improvement
- ⚠ Higher nightmare frequency → reduced response to IRT

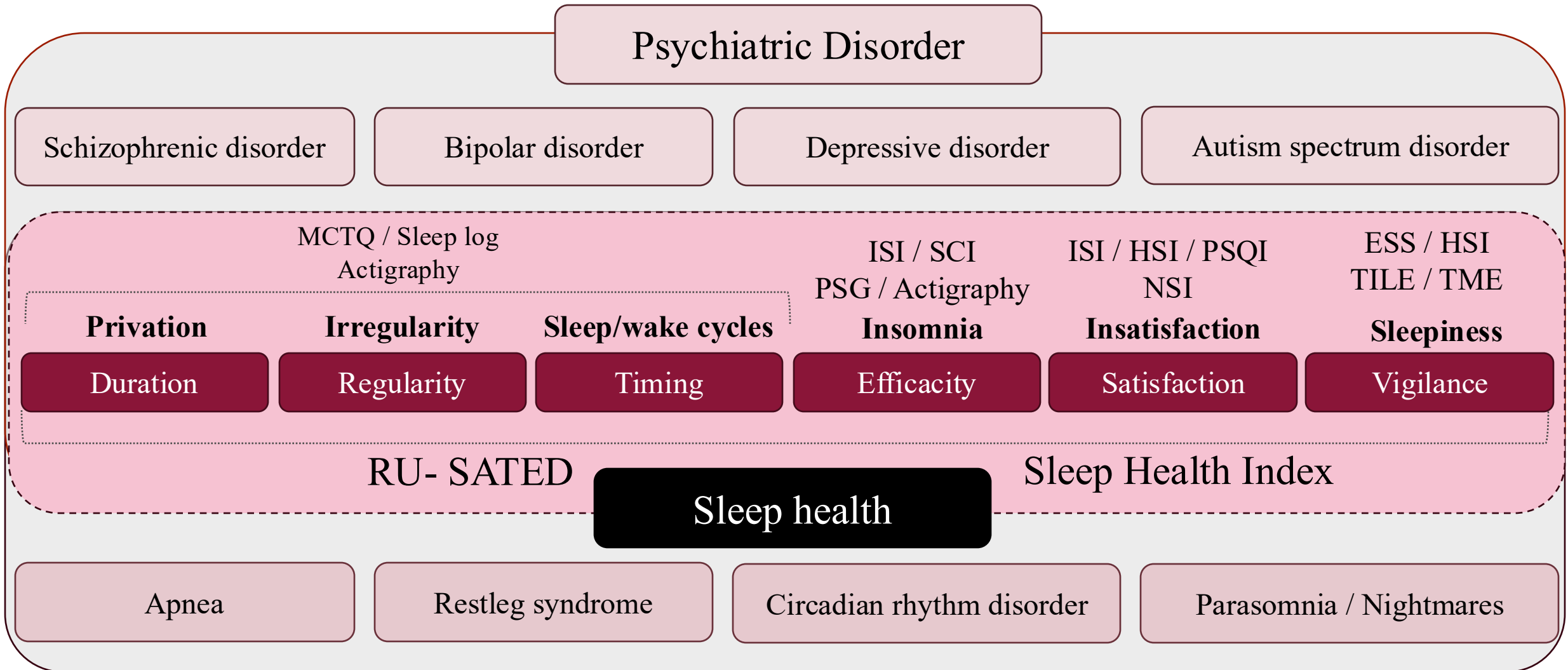
Centre Chronos – Hôpital Bichat (Paris)

Non-randomized controlled study | n = 53

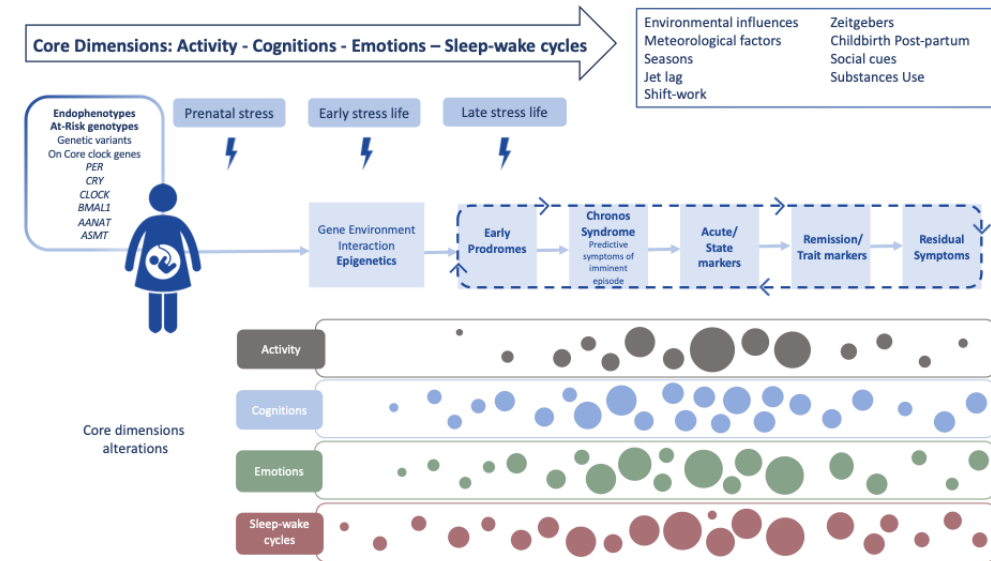
- Sleep disturbances and subscores PSQI accross BS, CZ, TRD and relations with clinical and biological factors
- Better characterize daytime symptoms and sleepiness in SZ, BD and TRD, and the impact on psychiatric (including suicide) and non-psychiatric disorders (including metabolic alterations)
- Links between cognitive alterations and sleep disturbances (night and daytime symptoms including sleepiness)
- Anhedonia and Sleep
- Study longitudinal sleep trajectories and links with psychiatric progressions (trajectories, crosslag)
- Psychotropics impact on sleep and psychiatric symptoms

Sleep as a transdiagnostic marker and intervention target in psychiatric health

Julien Coelho, Ludovic Samalin, Antoine Yroni, Anton Iftimovici, Pierre Philip, Jean-Arthur Micoulaud-Franchi



- Alterations in sleep–wake cycles are present across all phases of mood disorders: prodromes, transitions (**Chronos syndrome**), remission, recurrences, and progression → a **core dimension**
- Characterizing this “sleep dimension”
 → enables truly personalized treatment approaches
- Targeting sleep
 → offers an opportunity for early intervention and relapse prevention



(Geoffroy PA & Maruani J, *Biological Psychiatry*, 2025)



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Benjamin Dudoignon



Nathalie Couque



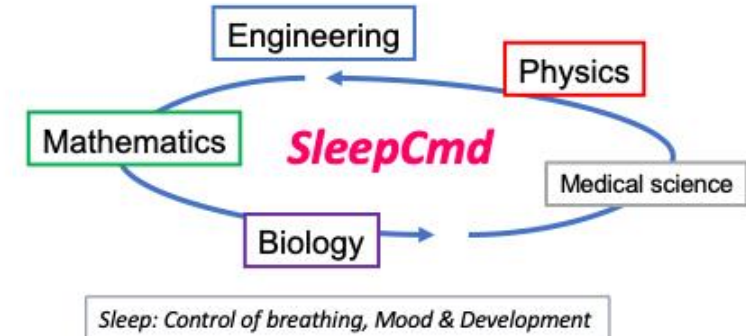
Michel Lejoyeux



Lily Vissoze



Julia Maruani



Factors associated with insomnia symptoms: A cross-sectional study during a Covid-19 fully restrictive lockdown

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 Justine Frija-Masson^{1,4,5} | Marie-Pia d'Ortho^{1,4,5} | Michel Lejoyeux^{1,2,6} |
 Dieter Riemann⁷ | Julia Maruani^{2,4} | Pierre A. Geoffroy^{2,4,6,8}

J Sleep Res, 2022

Variable	p	B	Odds ratio	95 C.I.	
				Lower	Upper
(a) Variables included in the final BSLR model					
Older age	0.06	0.14	1.15	0.99	1.33
Female	0.10	0.23	1.26	0.96	1.65
Housing size m ² (>100 m ²)	0.03	-0.264	0.77	0.60	0.98
Sleeping in a noisy environment	0.07	-0.46	1.60	0.96	2.62
Less regularity of sleep schedules	0.00	-0.22	1.25	1.18	1.32
Longer screen exposure in the morning	0.04	0.12	1.13	1.00	1.27
Longer strong light exposure in the evening	0.04	0.16	1.17	1.00	1.37
HAD-anxiety scores	0.00	0.22	1.24	1.19	1.29
HAD-depression scores	0.00	0.14	1.15	1.10	1.20
(b) Classification table and summary statistics					
Observed	Predicted		% Correctly classified		
	No insomnia	Insomnia			
No insomnia	614	187	76.77%		
Insomnia	230	589	71.92%		

TABLE 4 Backward stepwise logistic regression (BSLR) showing (a) the best combination of variables for correctly classifying study participants as insomniac or non-insomniac and (b) the overall classification rate by group

Note: % of all participants correctly classified 74.26% – Chi-square = 533.534, df = 9; p < 0.005.

Habitual sleep duration, if decreased <7 h or increased >8 h:

- ↑ mortality in the general population
(Cappuccio et al., Sleep, 2010)
- ↑ diabetes, stroke, coronary artery disease, and myocardial infarction
(Nuyujukian et al., Sleep health, 2019 ; Yin et al., J Am Heart Assoc, 2017)
- ↑ obesity (decreased sleep duration)
(Li et al., J Paediatr Child Health, 2017)

Habitual sleep duration and 12-month prevalences of Psychiatric disorders

Cohort representative of the US general population
(N = 36,309)

Prevalence of psychiatric disorder

<5H/night = 55%
7-8h = 28%
>9h = 48%

Prevalence of mood disorder

<5H/night = 23%
7-8h = 9%
>9h = 15%

Prevalence of suicide attempt

<5H/night = 11%
7-8h = 3%
>9h = 7%

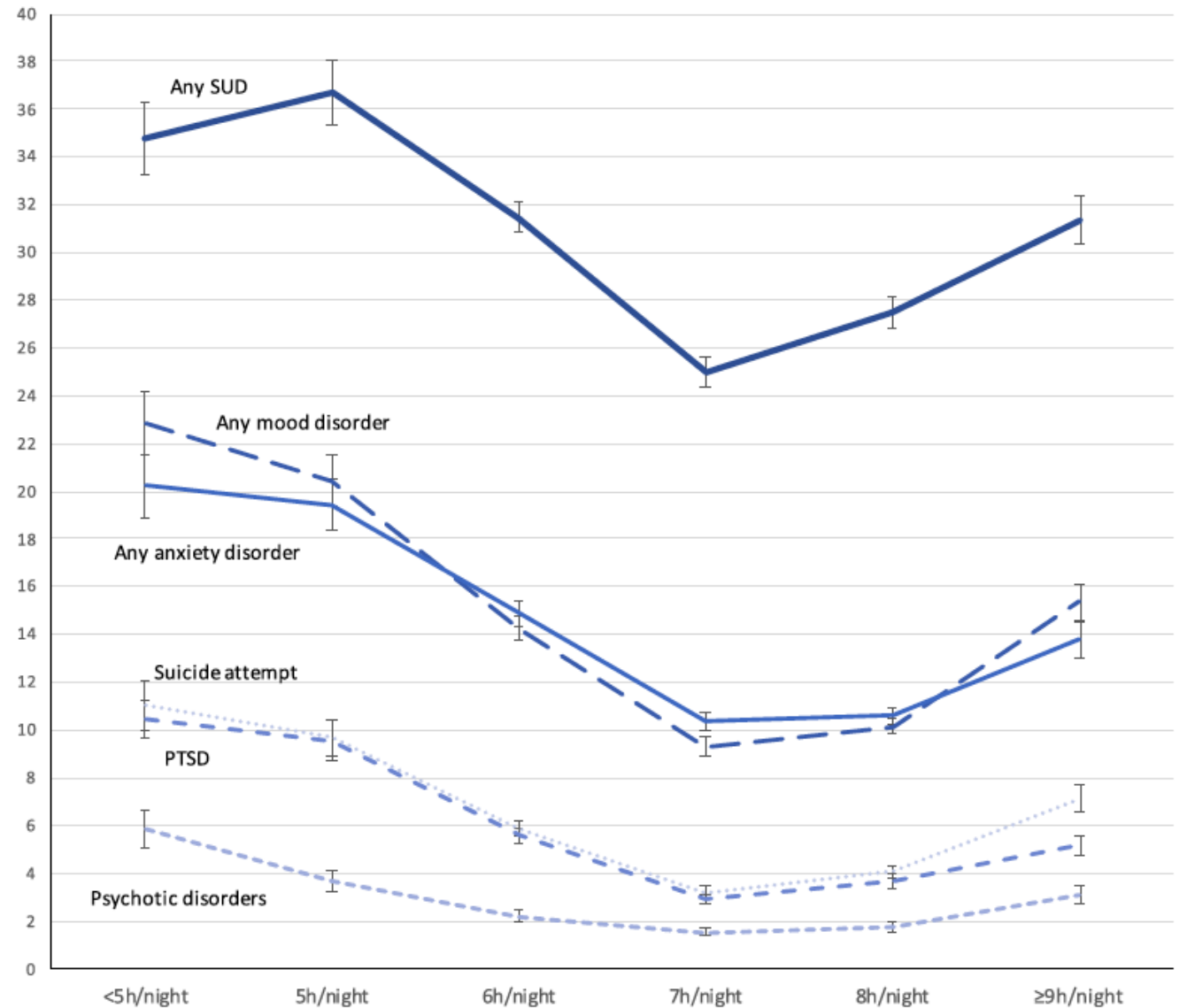


Fig. 1. The 12-month prevalences of psychiatric and substance use disorders depending on the habitual sleep duration

PTSD: Post-Traumatic Stress Disorder; SUD: Substance Use Disorder

“Any SUD” included alcohol use disorder, tobacco use disorder, and drug use disorder (cannabis, club drugs, cocaine, amphetamine, hallucinogen, heroin, opioid, sedative, tranquilizer, solvent or inhalant). “Any mood disorder” included major depressive disorder, bipolar disorder type I or II. “Any anxiety disorder included panic disorder, social phobia, specific phobia, generalized disorder. Psychotic disorder included schizophrenia or a psychotic illness or episode in the past 12 months. PTSD and suicide attempt were categories by their own.