Japanese Journal of Medical Research

Chronic Medical Disorders during COVID-19: Differences in Morbidities, Work, Habits and Attitudes

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Received: 04 Nov 2024; Accepted: 12 Dec 2024; Published: 24 Dec 2024

Citation: Paiva T, Luzeiro I Reis C. Chronic Medical Disorders during COVID-19: Differences in Morbidities, Work, Habits and Attitudes. Japanese J Med Res. 2024; 2(1): 1-14.

ABSTRACT

Background: Chronic medical disorders (CMD) are highly prevalent and are often associated with multiple mental and physical comorbidities, which amplify health risks and pose significant challenges for long-term treatment, contributing to their substantial burden on individuals and healthcare systems. During the pandemic CMD were particularly affected. Portugal besides having a higher prevalence of CMD is among the countries which used more strict protective pandemic measures with subsequent negative impacts.

Objectives: The paper aims to identify different dysfunctional patterns in 14 CMD: diabetes, hypertension, cardiovascular (CVD), fibromyalgia, rheumatologic, respiratory, allergic, autoimmune, gastrointestinal and dermatologic disorders, chronic pain, fatigue, tinnitus and dizziness.

Methods: Online surveys were used for data collection gathering 5479 individuals, mean age 48.5 years, ranging from 18 to 90 years, 67.7% were females. Demographics, work before COVID, health status, confinement, attitudes and behaviors, mental health, sleep, physical activity, multimedia use, nutrition, toxics and additions. ANOVA and linear discriminant analysis (LDA) were used for data analysis and significance was set at p < 0.05.

Results: The most important features to differentiate CMD were comorbidities, work stress before COVID, Sleep, Mental Health and Attitudes. The disorders with higher canonical correlations with the discriminant function in LDA were: Chronic Pain, Fibromyalgia, Autoimmune, CVD, Fatigue, Diabetes, Dermatologic, Dizziness. Correct classification of the original group varied between 79.2 and 92.0%. Correct classification for "Not having" a specific CMD varied between 79.7 and 92.5%. Correct classification for "Having" a specific CMD provided lower values and varied between 60.0 and 82.4%.

Conclusions: The CMD studied exhibited distinct characteristics both among themselves and compared to individuals without the disease. LDA achieved high accuracy in classifying each CMD. These findings provide valuable insights for guiding strategies to manage CMD's during future public heath disasters.

Keywords

Chronic medical disorders, Comorbidities, COVID19, Sleep, Work Stress.

Abbreviations

AID: Auto Immune Disorders, BMI: Body Mass Index, CMD: Chronic Medical Disorders, CVD: Cardiovascular disorders, EU: European Union, GDP: Gross Domestic Product, GID: Gastro-Intestinal Disorders, LDA: Linear Discriminant Analysis, WHO: World Health Organization.

Introduction

COVID was a societal and a health care marker. Despite the main symptoms being similar to a common flue (dry cough, shortness of breath, headaches, muscle aches, fever or chills) other symptoms pointed to upper airway and digestive tract involvement (loss of taste or smell, fatigue, upset stomach, vomiting, diarrhea) and threatening complications occurred, such as severe pneumonia and death.

A death rate around 1% was devastating and urged efficient measures worldwide [1]. Death and serious complications varied among countries and regions [2], but globally they increased with age, affecting particularly the elderly. Multiple morbidities were associated with severe COVID-19 illness; obesity, diabetes with complications, and anxiety and fear-related disorders had the strongest association with death [3,4].

Long-term consequences and immunologic dysfunction were further serious consequences which health care services are still handling. In Scotland older age, higher BMI, severe COVID-19 infection, female sex, social deprivation and comorbidities were predictors of long COVID, while vaccination against COVID-19 and testing positive, while Delta or Omicron variants were dominant, predicted reduced risk [5]. In a Chinese study significant predictors of long-term fatigue were length of hospitalization, nonuse of antiviral drug, immune-related serum markers of IL-6 and CD16+CD56⁺ NK cell levels, neurologic diseases and a lack of vaccination [6]. However, the variability of symptoms included in long COVID is quite high together with the presence of preexisting morbidities [7].

In Portugal the predictors of long COVID after discharge from a specialized centre were fatigue and persistent cough at 3 months and pain and cognitive disturbances at 6 months; symptoms had significant negative impact upon quality of life. Concerning long COVID risk factors this study provided somewhat different results since neither age, nor gender, nor previous morbidities were significant risks [8]. Symptoms cluster analysis of long COVID Portuguese patients was performed, and four clusters were found: Cluster 1 had older patients, 90% females and minor symptoms; Cluster 2 had a high female percentage and joint pain, myalgia, fatigue, insomnia, headache and palpitations; Cluster 3 symptoms had the highest prevalence of pre-existing health conditions, fatigue affected 100%, headache 86% of the patients, joint pain and tobacco consumption were frequent; in Cluster 4 memory loss

and concentration issues were the most frequent symptoms and the patients had a high prevalence of alcohol use [7].

According to the EU-SILC survey 41 % of Portuguese people aged 16 and over reported having at least one chronic condition – a higher proportion than in the EU (36 %). One third (30 %) of all deaths in Portugal in 2019 can be attributed to behavioural risk factors (smoking, dietary risks, alcohol and low physical activity); this share is however lower than the EU average (39%) [9]. Taking this data into consideration it is important to evaluate behavioural, environmental and health factors associated with chronic disorders during the COVID pandemic to get relevant knowledge, which can be potentially helpful in future pandemics, namely in preventing negative short- and long-term consequences of Chronic Medical Disorders (CMD). The aim of the present paper was to obtain differential behavioural and environmental factors of CMD during the first COVID outbreak, which can be integrated in future recommendations of more personalized health care strategies.

Methods

Online surveys were conducted with 5479 participants (mean age: 48.6 ± 14.3 years; range: 18-90 years), of whom 67.7% were female. The study included data collected from the Portuguese mainland and the Islands of Madeira and the Azores [10]. The overall project was approved by CENC's Ethical Committee 1/2020. There was no funding, public or private, and no conflict of interests.

The Survey Legend platform was utilized to conduct anonymous surveys targeting adults (aged >18 years). Participants provide their consent for data collection allowing data analysis and statistical use. Data were collected online during the first wave of COVID-19, spanning from April to August 2020. The surveys addressed the following topics: Demographics, Work before COVID; Health Status; Confinement/Lockdown Attitudes and behaviors; Mental health; Sleep; Physical activity; Multimedia use; Nutrition; Toxics and Additions. Health Status included yes/no questions: being healthy (subjective) or suffering from the following diseases: Sleep, Psychiatric, Neurologic, Cardiovascular, Respiratory, Allergies, Gastrointestinal, Rheumatologic, Endocrinologic/metabolic, Orthopedic, Autoimmune, Cancer, Renal, Dermatologic, Hematologic, Gynecologic, Urologic, Ear-Nose-Throat (ENT), Ophthalmologic, Chronic pain, Fatigue, Tinnitus and Dizziness.

The Morbidities Index (MI) represents the total number of reported morbidities at baseline. During the COVID-19 pandemic, participants indicated whether their morbidities had worsened or improved. Based on this, two additional indices were calculated: the COVID-19 Worsening (Morbidities Worsening Index (MWI)) and improvement (Morbidities Improvement Index (MII)).

Confinement attitudes and behaviors were evaluated by yes/ no answers [10]. The average and number of both, positive and negative attitudes and behaviors were computed per subject.

Calamity mood data were obtained by 1 to 10 visual analogue scales (VAS). The Calamity Experience Check List was computed

by averaging the scales of depression, anxiety, irritability, worries [11]. Sleep data included data relative to weekdays and weekends during COVID-19: Sleep schedules, Subjective Sleep duration in hours, sleep latency in minutes, number of awakenings and Sleep Quality (SQ) and Awakening quality (AQ) obtained in a 1-10 VAS. Physical activity: intensity (null, mild, moderate, intense) and hours/week. TV, Social Networks, Mobile Phone, Gaming use were quantified in hours/day. Nutrition included: meals/day. Scores for the recommended frequencies were calculated [12]. Smoking: yes/no; cigarettes/day. Alcoholic intake: Alcoholfree: Yes/No; glasses/day of beer, wine, aperitive wine, brandy. Drugs: no; occasionally; sometimes; regularly. The prevalence of chronic disorders and complaints among survey responders was calculated; only disorders with prevalences higher than 1.6% (86 subjects were used).

Quantitative variables were calculated by the mean and its confidence interval. Normality was tested by Kolmogorov-Smirnov test. Most continuous variables had a normal distribution except calamity scale, get up time for weekdays and weekends, cigarettes per/day. ANOVA (unidirectional analysis of variance) with Post-hoc Bonferroni tests was used comparing each chronic disease with the remaining population.

Linear Discriminant Analysis (LDA) was used in order to find a linear combination of features characterizing the different sub samples associated with chronic disorders or complaints (n=14) using the continuous variables with gaussian distribution as predictors: age, Body Mass Index (BMI), work stress variables before COVID, lockdown days, number of morbidities, and its evolution during COVID, variables related with daily living in confinement (attitudes and behaviors either positive or negative), humor VAS (depression, anxiety, irritability, worries), economic problems, sleep variables, screen time (TV, Mobile, Social networks, gaming), nutrition, physical activity. Discriminant functions, Structure Correlation Coefficients, Functions at Group Centroids, Eigen values and Canonical correlations, Wilks Lambda, Box M and Classifier (computed for cases without missing values (n=900) and for the total samples) were calculated.

Table	1
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All tests were performed using the SPSSv29. Statistical significance was assessed with an alpha level of 0.05.

Results

The number of patients and corresponding percentage in the total population are shown in table 1. Allergies, Hypertension, Fatigue and Respiratory disorders were the most prevalent. The prevalence of conditions showed some variation by sex. Males were more prevalent in certain groups, such as Diabetes and Hypertension. In Cardiovascular disorders (CVD) no significant sex differences were observed, while females predominated in the remaining groups.

Table 2 presents data on demographics, before COVID work stress, morbidities and confinement characteristics.

Patients with diabetes, hypertension, CVD, fibromyalgia, rheumatologic disorders, and tinnitus were older; in respiratory, allergies, autoimmune, and dermatologic disorders were younger; there were no age differences in gastrointestinal disorders (GID), chronic pain, fatigue and dizziness. Regarding work conditions before COVID, patients with hypertension reported higher levels of stress and greater responsibilities. Similarly, patients with respiratory disorders reported higher stress levels, increased responsibilities, more frequent conflicts, greater exposure to moral or sexual harassment, and intellectually demanding work. Patients with allergies reported significantly higher stress levels, more interruptions, multitasking, conflicts, responsibilities, and work that was both intellectually and physically demanding. Those with gastrointestinal disorders (GID) reported increased conflicts. Fibromyalgia patients experienced higher levels of conflicts and moral/sexual harassment, while patients with rheumatic disorders reported more interruptions and moral/sexual harassment. Patients with autoimmune disorders (AID) faced more interruptions, multitasking, responsibilities, conflicts and work that was both intellectually and physically demanding. Patients suffering from chronic pain and fatigue reported the highest levels across all work-related factors including stress, interruptions, multitasking, conflicts, responsibilities, moral/sexual harassment, and both intellectually and physically demanding work. Those with

		Fotal	I	Males	F	Females			
Disease	No n (%)	Yes n (%)	No n (%)	Yes n (%)	No n (%)	Yes n (%)	p-value		
Diabetes	5266 (98.1%)	104 (1.9%)	1678 (96.8	56 (3.2)	3608 (98.7)	49 (1.3)	<0.001		
Hypertension	4856 (90.4%)	514 (9.6%)	1513 (87.3)	221 (12.7)	3363 (92.0)	294 (8.0)	<0.001		
Cardiovascular	5224 (97.3%)	146 (2.7%)	1678 (96.8)	56 (3.2)	3566 (97.5)	91 (2.5)	0.119		
Respiratory	5048 (94.0%)	322 (6.0%)	1669 (96.3)	65 (3.7)	3400 (93.0)	257 (7.0)	<0.001		
Allergy	4659 (86.8%)	711 (13.2%)	1620 (93.4)	114 (6.6)	3059 (83.6)	598 (16.4)	<0.001		
Gastrointestinal	5202 (96.9%)	168 (3.1%)	1694 (97.7)	40 (2.3)	3529 (96.5)	128 (3.5)	0.019		
Fibromyalgia	5284 (98.4%)	86 (1.6%)	1727 (99.6)	7 (0.4)	3574 (97.7)	93 (2.5)	<0.001		
Rheumatic	5250 (97.8%)	120 (2.2%)	1707 (98.4)	27 (1.6)	3564 (97.5)	93 (2.5)	0.022		
Autoimmune	5162 (96.1%)	208 (3.9%)	1713 (98.8)	21 (1.2)	3470 (94.9)	187 (5.1)	<0.001		
Chronic Pain	5205 (98.9%)	166 (3.1%)	1716 (99.0)	18 (1.0)	3508 (95.9)	150 (4.1)	<0.001		
Fatigue	4980 (92.7%)	391 (7.3%)	1681 (96.9)	53 (3.1)	3320 (90.8)	338 (9.2)	<0.001		
Dermatologic	5256 (97.9%)	114 (2.1)	1714 (98.8)	29 (1.2)	3563 (97.4)	94 (2.6)	<0.001		
Tinnitus	5224 (97.3%)	147 (2.7%)	1699 (98.0)	35 (2.0)	3546 (96.9)	112 (3.1)	0.028		
Dizziness	5192 (96.7%)	179 (3.3%)	1710 (98.6)	24 (1.4)	3502 (95.7)	156 (4.3)	<0.001		

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			Diabetes	Hypertension	CVD	Respiratory	Allergies	GID	Fibromyalgia	Rheumatic	AID	Chronic Pain	Fatigue	Dermatologic	Tinnitus	Dizziness
		No	48.3	47.5	48.3	48.6	49.4	48.5	48.4	48.4	48.6	48.4	48.5	48.6	48.4	48.5
	Age	Yes	58.8	57.9	56.0	46.9	42.7	48.8	54.1	52.4	46.5	50.5	48.1	44.5	51.2	49.3
_	ANOVA	Sig	.000	.000	.000	.038	.000	.819	.000	.003	.040	.067	.591	.002	.018	.428
3M		No	25.7	25.5	25.7	25.8	25.9	25.8	25.7	25.7	25.8	25.8	25.7	25.8	25.7	25.7
L Dr	BMI	Ves	27.7	27.9	27.3	25.5	23.5	22.0	27.1	26.5	25.3	25.0	25.7	25.0	25.8	25.8
e ai	ANOVA	Sig	000	000	000	329	000	030	011	418	172	418	329	805	866	875
Ag		No	0.20	0.17	0.18	0.15	0.11	0.18	0.16	0.16	0.16	0.15	0.12	0.17	0.17	0.16
	Weight diff	Ves	-0.92	0.23	0.03	0.15	0.57	0.15	0.76	0.77	0.10	0.92	0.93	0.31	0.39	0.56
		Sig	0.52	699	587	0.37	0.57	917	0.70	0.77	111	0.92	0.95	672	441	115
	W atraag D	No	2.83	2.82	2.83	2.82	2.80	2.83	2.83	2.83	2.83	282	2.80	2.83	2.83	2.82
	Covid	Vos	2.05	2.82	2.05	3.06	3.06	3.01	3.09	3.04	2.05	3.06	3.26	3 11	3.05	3.18
		Sig	0.103	0.041	0.112	001	000	0.067	0.066	0.064	0.134	0.017	000	0.018	0.035	0.000
	W intermentions	No	2.24	2.24	2.24	2.001	2.21	2.24	2.24	2.004	2.22	2.22	2.000	2.24	2.24	2.24
	B Covid	Voc	2.24	2.24	2.24	2.23	2.21	2.24	2.24	2.23	2.23	2.23	2.22	2.24	2.24	2.24
		Sig	0.333	0.535	2.15 0.314	2.30	2.41	2.30	2.34	0.005	2.47	2.47	2.47	0.128	0.751	0.492
		No	2.04	2.05	2.04	2.02	2.00	2.02	2.04	2.04	2.02	2.02	2.01	2.04	2.02	2.02
	W multitask B	No.	2.04	3.03	2.04	2.17	2.00	2.12	2.04	2.16	2.05	2.05	2.20	2.12	2.14	2.05
		Sig	2.09	2.97	0.974	0.079	000	0.206	0.256	0.342	5.20 0.010	5.20 0.021	000	0.526	0.252	0.056
le.	W conflicts D	No	2.10	2.10	2 10	2.08	2.06	2.00	2.10	2.10	2.00	2.00	2.07	2.00	2.00	2.08
WORK before COV	Covid	Vos	2.10	2.10	2.10	2.00	2.00	2.09	2.10	2.10	2.09	2.09	2.07	2.09	2.09	2.08
		Sig	0.369	0.675	0.508	001	000	0.029	0.013	0.356	0.009	000	000	0.008	0.001	0.000
	Responsibilities	No	3 30	3 38	3 38	3 38	3 36	3 30	3 30	3 30	3 38	3 38	3 37	3 30	3 38	3 38
	R Covid	Voc	3.12	3.50	3.58	3.60	3.50	3.37	3.16	3.11	3.65	3.50	3.68	3.53	3.50	3.75
		Sig	0.846	0.042	0.087	0.005	000	0.830	0.665	0.901	0.005	0.012	000	0.267	0 101	0.001
	W1 and and D	No	1.09	1.08	1.09	1.09	1.09	1.09	1.09	1.09	1.00	1.09	1.07	1.09	1.09	1.09
	Covid	Vos	1.00	1.08	1.00	1.00	1.00	1.08	1.00	1.00	1.00	1.06	1.07	1.00	1.00	1.00
		Sig	0.540	0.820	0.703	0.002	0.068	0.687	0.000	0.010	0.521	000	000	0.038	0.005	0.002
	Wintelest	No	3 21	3.22	3.22	3.20	3.18	3 21	3 21	3 21	3 21	3 21	3 10	3 21	3 21	3.21
	heavy	Vos	3.21	3.22	3.22	3.20	3.16	3.21	3.21	3.21	3.46	3.21	3.19	3.21	3.21	3.21
		Sig	0.566	0.986	0.908	0.000	000	0.335	0.310	0.268	0.000	0.018	000	0.047	0.022	0.117
	W physicaly	No	2 33	2 32	2 33	2 32	2 32	2 33	2 33	2 33	2 32	2 32	2 30	2 33	2 33	2 32
	heavy	Vos	2.33	2.32	2.33	2.32	2.52	2.33	2.55	2.35	2.52	2.52	2.50	2.33	2.33	2.52
		Sig	0.681	0.242	0.107	0.127	0.045	0.324	0.164	0.706	0.001	0.008	0.000	0.615	0.385	0.016
	AIOIA	No	1.63	1.50	1.60	1.55	1.42	1.58	1.60	1.61	1.58	1.54	1.42	1.61	1.58	1.56
	N morbidities	Vos	3.48	3.10	3.60	3.44	3.27	1.50	5.12	1.01	3.74	5 20	1.72	3 00	1.50	1.50
	ANOVA	Sig	000	000	000	000	000	000	000	000	000	000	000	000	000	4.00 000
ties	Morbidities	No	1 73	1 71	1 72	1 71	1 69	1 72	1 72	1 72	1 71	1 70	1 66	1 72	1 71	1 70
idi	Worse	Ves	1.75	1.87	1.93	2.06	2.00	2 10	2 50	2 21	2.08	2 59	2.56	2 22	2 31	2 42
orb	ANOVA	Sig	0.423	0.103	0.215	0.003	0.000	012	000	0.007	0.008	0.000	0.000	0.000	0.000	0.000
Σ	Morbidities	No	0.42	0.44	0.43	0.43	0.45	0.43	0.41	0.42	0.43	0.42	0.43	0.42	0.42	0.42
	Better	Yes	0.32	0.23	0.21	0.30	0.23	0.19	0.76	0.34	0.26	0.28	0.25	0.18	0.27	0.27
	ANOVA	Sig	0.344	0.000	0.023	0.052	0.000	.007	0.007	0.424	0.044	0.099	0.002	0.000	0.097	0.078
	Number of days	No	47.04	47.27	47.03	47.04	47.32	46.91	46.99	46.82	46.88	47.00	47.05	47.14	47.01	46.94
	in lockdown	Yes	43.59	44.22	44.86	45.95	44.52	49.35	46.34	53.14	49.36	46.12	45.97	37.08	45.56	48.15
t	ANOVA	Sig	0.758	0.065	0.477	0.608	0.062	0.420	0.857	0.048	0.338	0.758	0.577	0.010	0.638	0.678
nen	Living in	No	6.58	6.58	6.58	6.59	6.60	6.58	6.58	6.58	6.59	6.59	6.61	6.58	6.59	6.59
nen	confinement	Yes	6.55	6.51	6.44	6.32	6.42	6.30	6.25	6.51	6.24	6.27	6.15	6.47	6.01	6.08
iluc	ANOVA	Sig	0.031	0.390	0.377	0.013	0.017	0.060	0.098	0.716	0.008	0.031	.000	0.530	.000	.000
Ŭ	Your economic	No	2,98	2,99	2,99	2,99	3,00	2,99	2,97	2,98	2,99	2,96	2,94	3,00	2,98	2,96
	problems	Yes	3,53	3,05	3,12	3,02	2,97	3,25	4,14	3,48	3,09	3,91	3,70	2,78	3,51	3,88
	ANOVA	Sig	0,000	0,516	0,492	0,847	0,788	0,127	0,000	0,015	0,535	0,000	0,000	0,295	0,004	0,000

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Ves firs 6.12 6.07 6.39 6.77 6.40 6.41 6.57 6.15 6.50 7.01 6.93 6.37 6.70 7.02 NoVA Sig 0.000 0.000 0.000 0.000 0.052 0.668 0.00 0.00 0.00 0.00 0.00 0.000
ANOVA Sig 0.000 0.200 0.000 0.000 0.052 0.668 0.009 0.000 0.168 0.002 0.003 Mow is frequency No 3.99 4.03 3.99 3.99 3.98 3.98 3.99 3.99 3.99 3.98 3.98 3.99 3.89 3.81 3.83 3.46 3.83 3.84 3.80 3.80 8.02 8.01 8.01 8.01 8.01 8.01 8.01 8.01 8.01 8.01 8.01 8.01 8.01 8.01 8.01 8.01 8.0
How is frequency vecked and activity No 3.99 4.03 3.99 3.99 3.98 3.98 3.99 4.00 4.02 3.99 4.00 3.99 ANOVA Sig 0.000 0.056 0.144 0.673 0.112 0.112 0.874 0.88 3.88 3.46 3.80 3.80 3.89 3.80
your sexual activity Yes 3.70 3.51 3.59 3.78 3.94 3.68 3.76 3.95 3.74 3.88 3.46 3.82 3.49 3.77 ADVA Sig 0.001 0.000 0.056 0.144 0.673 0.112 0.415 0.874 0.814 0.000 0.463 0.013 0.238 Time out of bed weekdays Yes 7.84 7.84 7.88 7.89 7.88 7.88 7.88 7.88 7.88 7.88 7.88 7.89 7.88 7.88 7.89 7.88 7.89 7.88 7.88 7.89 7.88 7.89 7.88 7.89 7.88 7.89 7.88 7.89 7.88 7.89 7.88 7.89 7.89 7.89 7.89 7.8
ANOVA Sig 0.001 0.000 0.056 0.144 0.673 0.112 0.415 0.874 0.148 0.001 0.000 0.633 0.238 Time out of bed weekdays Yes 7.89 7.89 7.89 7.90 7.71 7.77 8.08 7.88 8.92
Time out of bed weekdays No 7.89 7.89 7.90 7.91 7.89 8.02 8.02 8.02 8.02 8.9
weekdays Yes 7.84 7.84 7.96 7.71 7.74 7.77 8.08 7.98 8.02 8.01 8.04 7.67 7.98 8.15 ANOVA Sig 7.74 8.03 8.03 8.02 8.04 2.50 5.44 2.23 3.32 0.49 1.36 4.82 0.25 Weekends Yes 8.88 8.92 8.93 8.93 8.92 <th< td=""></th<>
ANOVA Sig 7.45 4.94 5.63 .043 .010 .335 .250 .544 .223 .332 .049 .136 .482 .025 Time out of bed weekends Yes 8.93 8.92 8.93 8.93 8.93 8.92 8.92 8.92 8.92 8.92 8.92 8.92 8.92 8.92 8.92 8.92 8.92 8.92 8.92 8.92 8.92 8.92 8.92 9.07 8.05 0.06 0.02 0.07 0.07 0.02 0.12 9.07 ANOVA Sig 0.042 0.470 0.038 0.324 0.431 0.954 0.391 0.067 0.07 0.26 0.26 0.26 0.26 0.26
Time out of bed weekends No 8.93 8.93 8.93 8.93 8.92 8.9
weekends Yes 8.88 8.98 9.21 8.84 8.88 8.92 9.08 9.20 8.99 9.18 8.96 9.08 9.12 9.07 ANOVA Sig 0.042 0.470 0.038 0.324 0.431 0.954 0.391 0.068 0.584 0.042 0.665 0.313 0.162 0.247 Time into bed weekdays No -0.06 -0.07 -0.06 -0.06 -0.06 -0.06 -0.07
ANOVA Sig 0.042 0.470 0.058 0.524 0.431 0.954 0.391 0.068 0.584 0.042 0.665 0.513 0.162 0.247 Time into bed weekdays No -0.06 -0.06 -0.06 -0.06 -0.070
Time into bed weekdays No -0.06 -0.06 -0.06 -0.06 -0.06 -0.07
Weekdays Yes -0.06 -0.11 -0.03 -0.17 -0.13 0.17 0.07 0.07 -0.02 0.11 -0.05 0.01 ANOVA Sig 0.306 0.560 0.828 0.021 0.454 0.420 0.704 0.129 0.252 0.306 0.574 0.257 0.934 0.542 Time into bed weekends No 0.27 0.26
ANOVA Sig 0.306 0.360 0.828 0.021 0.434 0.420 0.744 0.129 0.252 0.306 0.374 0.257 0.934 0.342 Time into bed weekends No 0.27 0.26 0.26 0.26 0.27 0.27 0.26 0.27 0.26
Imme into bed weekends No 0.27 0.26 0.26 0.27 0.27 0.26<
Weekends Fes 0.22 0.28 0.39 0.29 0.16 0.18 0.41 0.44 0.57 0.27 0.48 0.43 0.44 ANOVA Sig 0.424 0.791 0.435 0.077 0.804 0.447 0.637 0.381 0.148 0.424 0.949 0.184 0.220 0.196 Sleep duration weekdays No 6.67 6.68 6.67 6.67 6.67 6.67 6.67 6.68 6.69 6.69 6.69 6.69 6.69 6.69 6.69 6.69 <
ANOVA Sig 0.424 0.791 0.435 0.677 0.604 0.447 0.637 0.148 0.424 0.949 0.184 0.220 0.190 Sleep duration weekdays No 6.67 6.68 6.67 6.71 6.68 6.67 6.68 6.69 6.69 6.43 6.67 6.69 6.68 6.67 6.68 6.68 6.68 6.68 6.69 6.69 6.84 ANOVA Sig 0.874 0.661 0.615 0.891 0.000 0.964 0.248 0.972 0.065 0.874 0.336 0.985 0.700 0.210 Sleep duration weekends No 7.46 7.44 7.45 7.46 7.45 7.47 7.46 7.47 7.45 7.46 7.46
Size duration No 0.03
ANOVA Sig 0.874 0.661 0.615 0.891 0.000 0.964 0.248 0.972 0.056 0.874 0.336 0.985 0.700 0.210 Sleep duration weekends No 7.46 7.44 7.45 7.46 7.48 7.46 7.45 7.47 7.46 7.47 7.46 7.47 7.46 7.46 7.46 ANOVA Sig 0.723 0.150 0.398 0.863 0.000 0.964 0.248 0.972 0.065 0.874 0.336 0.985 0.700 0.210 Sleep duration weekends No 7.46 7.44 7.45 7.46 7.45 7.47 7.46 7.47 7.45 7.46 7.46 ANOVA Sig 0.723 0.150 0.398 0.863 0.041 0.546 0.073 0.149 0.087 0.723 0.240 0.304 0.811 0.998 Sleep latency weekdays No 32.25 32.28 32.21 32.31 </td
AROVA Sig 0.014 0.015 0.015 0.015 0.015 0.015 0.015 0.015 0.015 0.016 0
Sice dulation No
ANOVA Sig 0.723 0.150 0.398 0.863 0.041 0.546 0.073 0.149 0.087 0.723 0.240 0.304 0.811 0.998 Sleep latency weekdays No 32.25 32.28 32.21 32.31 32.07 32.16 32.09 32.34 32.27 32.12 31.74 32.18 32.18 32.04 Move Yes 29.21 31.29 31.50 0.586 0.772 0.161 0.059 0.452 0.514 0.003 0.940 0.963 0.129 Move Sig 0.514 0.575 0.823 0.370 0.586 0.772 0.161 0.059 0.452 0.514 0.003 0.940 0.963 0.129 Sleep latency No 31.72 31.82 31.71 31.81 31.47 31.57 31.54 31.77 31.75 31.63 31.63 31.63 31.63
Sleep latency weekdays No 32.25 32.28 32.21 32.31 32.07 32.16 32.09 32.34 32.27 32.12 31.74 32.18 32.18 32.04 Mo 29.21 31.29 31.50 30.32 32.92 33.03 38.00 25.79 30.17 34.05 37.82 32.46 32.33 36.52 Mov Sig 0.514 0.575 0.823 0.370 0.586 0.772 0.161 0.059 0.452 0.514 0.003 0.940 0.963 0.129 Sleep latency No 31.72 31.82 31.71 31.81 31.47 31.57 31.54 31.77 31.76 31.25 31.63 31.63 31.50
weekdays Yes 29.21 31.29 31.50 30.32 32.92 33.03 38.00 25.79 30.17 34.05 37.82 32.46 32.33 36.52 MovA Sig 0.514 0.575 0.823 0.370 0.586 0.772 0.161 0.059 0.452 0.514 0.003 0.940 0.963 0.129 Sleep latency No 31.72 31.82 31.71 31.47 31.57 31.54 31.77 31.56 31.25 31.63 31.63 31.50
ANOVA Sig 0.514 0.575 0.823 0.370 0.586 0.772 0.161 0.059 0.452 0.514 0.003 0.940 0.963 0.129 Steep latency No 31.72 31.82 31.71 31.47 31.57 31.54 31.77 31.56 31.25 31.63 31.63 31.50
Sleep latency No 31.72 31.82 31.71 31.81 31.47 31.57 31.54 31.77 31.77 31.56 31.25 31.63 31.63 31.50
weekends Yes 28.65 30.17 29.74 29.30 32.88 34.24 38.80 27.05 28.73 34.47 36.84 32.72 32.75 36.24
ANOVA Sig 0.335 0.362 0.547 0.270 0.373 0.387 0.089 0.183 0.285 0.335 0.007 0.771 0.738 0.114
Awakenings No 2.8 2
weekdays Yes 2.3 2.9 3.2 3.0 2.9 2.6 3.0 2.9 3.0 3.4 3.0 2.5 3.0 2.3
ANOVA Sig 0.018 0.557 0.161 0.174 0.182 0.482 0.639 0.724 0.350 0.018 0.172 0.319 0.532 0.067
Awakenings No 2.4 2
weekends Yes 2.1 2.5 2.4 2.6 2.5 2.4 2.8 2.4 2.5 2.6 2.6 2.2 2.4 2.2
ANOVA Sig 0.258 0.462 0.820 0.118 0.547 0.926 0.113 0.977 0.435 0.258 0.064 0.281 0.826 0.192
SleepQuality No 5.68 5.70 5.69 5.71 5.75 5.71 5.71 5.70 5.71 5.73 5.79 5.69 5.71 5.71
Yes 5./3 5.58 5.3/ 5.28 5.30 5.02 4.32 5.04 5.20 4.40 4.41 5.28 4.90 4.87
ANOVA Sig 0.000 0.291 0.108 0.002 0.000 0.000 0.000 0.003 0.003 0.0000 0.000 0.000 0.000 0.000 0.000 0
Steep Waking NO 5.79 5.80 5.85 5.82 5.81 5.82 5.84 5.89 5.82 5.82 5.81 5.82 5.84 5.89 5.82 5.82 5.81 5.82 5
Quality I CS 0.10 5.81 5.36 5.27 5.39 5.05 4.47 5.35 5.17 4.38 4.30 5.42 4.96 4.94 ANOVA Sig 0.000 0.972 0.000 0.000 0.000 0.000 0.022 0.000 0.000 0.000
House physical No. 2.75 2.77 2.75 2.75 2.76 2.75 2.74 2.76 2.77 2.75 2.78 2.75 2.74 2.74
activity Ves 2.60 2.53 2.6 2.71 2.68 2.58 2.00 2.35 2.74 2.70 2.71 2.75 2.76 2.75 2.74 2.74
ANOVA Sig 0.563 0.321 0.495 0.914 0.706 0.672 0.674 0.378 0.104 0.563 0.052 0.863 0.934 0.415

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	Mool's day	No	3.84	3.86	3.84	3.84	3.82	3.84	3.84	3.84	3.84	3.84	3.84	3.84	3.84	3.84
	wiear's day	Yes	3.74	3.68	3.73	3.91	3.97	3.84	3.99	3.88	3.94	3.89	3.83	3.82	3.75	3.81
	ANOVA	Sig	0.494	0.000	0.168	0.217	0.000	0.990	0.181	0.677	0.144	0.494	0.834	0.819	0.278	0.666
d Physical Activity	Food	No	1.3	1.3	1.3	1.3	1.3	1.3	1.3	1.3	1.3	1.3	1.3	1.3	1.3	1.3
	Recommended	Yes	1.3	1.3	1.3	1.3	1.3	1.3	1.3	1.3	1.3	1.3	1.3	1.3	1.3	1.3
	ANOVA	Sig	0.640	0.097	0.727	0.065	0.000	0.296	0.574	0.106	0.613	0.640	0.051	0.164	0.079	0.080
	Food REC YES	No	5.3	5.3	5.3	5.3	5.3	5.3	5.3	5.3	5.3	5.3	5.3	5.3	5.3	5.3
		Yes	5.4	5.4	5.4	5.1	5.0	5.5	5.3	5.6	5.0	5.4	5.0	4.9	5.6	5.6
	ANOVA	Sig	0.446	0.144	0.742	0.130	0.004	0.344	0.955	0.118	0.077	0.446	0.054	0.150	0.079	0.076
		No	11.6	11.6	11.6	11.6	11.5	11.6	11.6	11.6	11.6	11.6	11.6	11.6	11.6	11.6
an	Food REC NO	Yes	11.6	11.4	11.6	11.8	11.9	11.4	11.2	11.3	11.7	11.4	11.7	12.0	11.3	11.3
ion	ANOVA	Sig	0.384	0.062	0.985	0.147	0.003	0.335	0.226	0.161	0.430	0.384	0.274	0.096	0.187	0.140
trit	Food YES NO	No	0.52	0.52	0.52	0.52	0.53	0.52	0.52	0.52	0.52	0.52	0.52	0.52	0.52	0.52
ñ	proportion	Yes	0.52	0.55	0.56	0.49	0.48	0.55	0.51	0.58	0.48	0.53	0.49	0.47	0.56	0.58
	ANOVA	Sig	0.735	0.156	0.238	0.132	0.007	0.325	0.731	0.132	0.166	0.735	0.127	0.148	0.263	0.089
	Hours physical	No	2.75	2.77	2.75	2.75	2.76	2.75	2.74	2.76	2.77	2.75	2.78	2.75	2.74	2.74
	activity	Yes	2.60	2.53	2.46	2.71	2.68	2.58	3.00	2.35	2.19	2.52	2.21	2.66	2.78	3.08
	ANOVA	Sig	0.563	0.321	0.495	0.914	0.706	0.672	0.674	0.378	0.104	0.563	0.052	0.863	0.934	0.415

dermatologic disorders and with tinnitus experienced elevated levels of work stress, conflicts, moral/sexual harassment and intellectually demanding work. Individuals with dizziness complaints reported increased work stress, conflicts, responsibilities, moral/sexual harassment and physically demanding work. In contrast, no significant work-related issues were observed among patients with diabetes or CVD.

The number of comorbidities was significantly higher in all subgroups. Comorbidities worsening during COVID were reported by patients with respiratory disorders, allergies, GID, fibromyalgia, rheumatic disorders, AID, chronic pain, fatigue, dermatologic, tinnitus and dizziness. Comorbidities improvement during COVID was reported only for patients with fibromyalgia. The remaining disorder groups either had not significant differences or significant differences with lower improvements levels than the participants without disorders (control population). Patients with rheumatic disorders were more days in lockdown, while those with dermatologic disorders were the ones with less days in lockdown, for the remaining groups no differences were observed. The following groups were feeling less well during lockdown when compared to controls: respiratory, allergic, AID, chronic pain, fatigue, tinnitus and dizziness. Economic problems were more prevalent among individuals with diabetes, fibromyalgia, rheumatic disorders, chronic pain, fatigue, tinnitus, and dizziness. Additionally, lower sexual frequency was reported by patients with hypertension, chronic pain, fatigue and tinnitus.

Data from Mental Health and the health pillars (Sleep, Nutrition and Physical activity) are shown in Table 3. Higher depression levels were reported by patients with respiratory disorders, GID, fibromyalgia, AID, chronic pain, fatigue, dermatologic, tinnitus, dizziness. Higher anxiety levels were reported by patients with respiratory disorders, allergies, GID, AID, chronic pain, fatigue, dermatologic, tinnitus, and dizziness. Higher levels of irritability were reported by patients with hypertension, respiratory disorders, allergies, GID, AID, chronic pain, fatigue, tinnitus, and dizziness. Worries towards uncertainty were higher in respiratory, allergies, AID, chronic pain, fatigue, tinnitus, and dizziness. Patients with respiratory and allergic diseases get out of bed earlier during weekdays while those with fatigue and dizziness get out of bed later. During weekends the patients' groups getting latter out of bed were the ones with CVD and chronic pain. There were no relevant differences in bedtime during weekdays and weekends. Patients with allergies had reduced sleep duration for both weekdays and weekends. Patients with fatigue had longer sleep latency for both weekdays and weekends. Patients with chronic pain report more frequent awakenings during weekdays. Patients with respiratory disorders, allergies, GID, fibromyalgia, rheumatic, AID, chronic pain, fatigue, tinnitus and dizziness reported worse Sleep Quality during the COVID. Patients with allergies, respiratory disorders, GID, fibromyalgia, rheumatic and autoimmune disorders, chronic pain, tinnitus and dizziness reported worse Awakening Quality. The number of meals per day were similar in patients with diabetes, cardiovascular, respiratory, GID, fibromyalgia, rheumatologic, chronic pain, fatigue, tinnitus and dizziness. However, the number of meals per day was lower in patients with hypertension, on the other hand the number of meals in patients with allergies was higher. Patients with allergies ate less recommended food and more non recommended foods.

In diabetes, hypertension, CVD, respiratory, allergies, GID, fibromyalgia, rheumatic, AID, chronic pain, dermatologic, tinnitus, and dizziness there were no differences in Physical Activity during COVID.

Data from toxics' use: screen time, dependences and attitudes are shown in table 4. Patients with diabetes, hypertension and CVD drink more beer per day, while patients complaining of dizziness drink less. patients with hypertension, respiratory, allergic, GID, chronic pain, AID, fatigue and dizziness drink less wine per day and patients with allergies drink significantly less aperitives and brandies per day.

Patients with fibromyalgia and chronic pain saw more TV. Patients with fibromyalgia, chronic pain, fatigue and dizziness were the ones spending more time in social networks, and the ones with allergies spent more time in the mobile phone. GID had lower levels

			Diabetes	Hypertension	CVD	Respiratory	Allergies	GID	Fibro-	Rheumatic	Autoimmune	Chronic	Fatigue	Dermatologic	Tinnitus	Dizziness
	How many	No	12.36	12.01	12.28	12.21	12 53	12 30	12 37	12.40	12 30	12 34	12 37	12.48	12 41	12.34
	cigarettes	Yes	14 54	16.40	16.39	15.41	11.33	15.42	14.78	12.40	12.55	14 23	12.37	5 00	12.41	15 75
	ANOVA	Sig	0.319	0.001	0.049	0.038	0 374	0.125	0.412	0.892	0.772	0.319	0.710	0.056	0.987	0.180
		No	0.49	0.46	0.50	0.50	0.51	0.50	0.51	0.50	0.51	0.51	0.51	0.51	0.50	0.52
	Beer per day	Yes	1.10	0.90	0.87	0.64	0.48	0.56	0.23	0.62	0.36	0.51	0.40	0.33	0.56	0.25
	ANOVA	Sig	0.964	0.000	0.004	0.144	0.689	0.693	0.072	0.413	0.175	0.964	0.191	0.313	0.681	0.027
	XX7. 1	No	0.23	0.24	0.23	0.23	0.25	0.23	0.22	0.23	0.23	0.23	0.24	0.23	0.23	0.23
	Wine per day	Yes	0.09	0.08	0.08	0.07	0.03	0.04	0.26	0.03	0.02	0.03	0.04	0.03	0.08	0.02
	ANOVA	Sig	0.026	0.004	0.123	0.019	0.000	0.042	0.719	0.053	0.009	0.026	0.001	0.132	0.123	0.019
cs	Aperitive Wine	No	0.02	0.02	0.02	0.02	0.02	0.02	0.02	0.02	0.02	0.02	0.02	0.02	0.02	0.02
Toxio	per day	Yes	0.00	0.00	0.00	0.01	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
	ANOVA	Sig	0.286	0.127	0.314	0.410	0.034	0.301	0.406	0.341	0.230	0.286	0.101	0.471	0.314	0.269
	Brandy per day	No	0.02	0.02	0.02	0.02	0.02	0.02	0.02	0.02	0.02	0.02	0.02	0.02	0.02	0.02
	Drundy per day	Yes	0.00	0.00	0.00	0.01	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
	ANOVA	Sig	0.288	0.120	0.316	0.386	0.035	0.303	0.408	0.343	0.232	0.288	0.102	0.473	0.316	0.272
	Alcohol	No	9.19	8.99	9.23	9.33	9.71	9.36	9.36	9.34	9.50	9.39	9.62	9.37	9.34	9.51
		Yes	14.36	11.93	11.42	8.66	6.16	7.12	6.06	7.77	4.58	6.46	5.30	4.24	7.68	3.27
	ANOVA	Sig	0.189	0.023	0.353	0.69/	0.004	0.329	0.243	0.529	0.014	0.189	0.004	0.113	0.478	0.004
	Drugs	INO	1.02	1.02	1.02	1.02	1.02	1.02	1.02	1.02	1.02	1.02	1.02	1.02	1.02	1.02
	ΑΝΟΥΑ	res	1.00	0.118	0.264	0.062	0.421	0.206	1.00	0.817	0.702	0.058	0.681	0.204	0.210	0.208
	ANOVA	No	0.938	3.06	3.08	3.08	3.10	3.08	3.06	0.01/	3.06	0.938	3.07	0.394 3.08	0.519	0.298
Screens	TV h Day	Vec	3.58	3.00	3.08	3.08	2.05	3.08	3.00 4 30	3.07	3.00	3.00	3.07	2.05	3.07	3.07
	ANOVA	Sig	0.026	0.078	0.988	1 000	0.221	0.997	0.000	0.139	0.052	0.026	0.449	0.633	0.092	0.223
	Social Networks	No	2 43	2 42	2 44	2 44	2 41	2 44	2 43	2 43	2 43	2 42	2 41	2 45	2 44	2 42
	h Dav	Yes	2.81	2.65	2.54	2.44	2.61	2.38	3.20	2.68	2.76	3.03	2.81	2.13	2.40	3.03
	ANOVA	Sig	0.009	0.103	0.676	0.993	0.090	0.795	0.015	0.329	0.101	0.009	0.009	0.204	0.851	0.005
		No	2.57	2.59	2.56	2.55	2.51	2.55	2.57	2.56	2.55	2.55	2.56	2.56	2.57	2.56
	Mobile h Day	Yes	2.33	2.33	2.66	2.89	2.90	2.99	2.51	2.77	2.93	3.03	2.71	2.75	2.58	2.74
	ANOVA	Sig	0.054	0.084	0.705	0.056	0.002	0.067	0.873	0.456	0.089	0.054	0.350	0.514	0.968	0.468
	Course h Door	No	1.91	1.93	1.90	1.87	1.91	1.90	1.90	1.90	1.90	1.90	1.88	1.91	1.91	1.89
	Games n Day	Yes	1.31	1.66	1.63	2.25	1.86	2.02	2.03	1.99	1.94	1.96	2.12	1.64	1.70	2.09
	ANOVA	Sig	0.826	0.174	0.540	0.122	0.764	0.735	0.765	0.802	0.886	0.826	0.300	0.467	0.532	0.535
	TV dependence	No	3.31	3.31	3.32	3.31	3.30	3.33	3.31	3.32	3.32	3.31	3.32	3.32	3.32	3.33
	i v dependence	Yes	3.57	3.40	3.14	3.50	3.45	2.95	3.68	3.31	3.32	3.46	3.35	3.38	3.39	3.08
	ANOVA	Sig	0.453	0.426	0.369	0.169	0.115	0.043	0.171	0.949	0.993	0.453	0.810	0.798	0.718	0.180
ø	Social Networks	No	3.66	3.69	3.66	3.64	3.58	3.66	3.65	3.65	3.64	3.65	3.64	3.65	3.66	3.65
nce	dependence	Yes	3.31	3.33	3.29	3.86	4.12	3.52	3.56	3.77	3.87	3.60	3.80	3.94	3.56	3.77
nde	ANOVA	Sig	0.813	0.005	0.098	0.170	0.000	0.518	0.744	0.630	0.246	0.813	0.265	0.246	0.677	0.569
ebei	Games	No	1.67	1.67	1.67	1.66	1.66	1.67	1.66	1.67	1.65	1.66	1.66	1.67	1.66	1.66
Ď		1 es	0.140	0.902	1.43	0.205	0.451	1.5/	0.140	0.753	0.002	0.140	1.75	0.657	0.541	0.082
	Alcohol	No	1.47	1.45	1.46	0.295	1.40	1.440	1.47	1.48	1.47	1.47	1.495	1.47	1.47	1.48
	dependence	Yes	1.26	1.66	1.40	1.45	1.36	1.47	1.40	1.70	1.41	1.39	1.36	1.25	1.45	1.70
	ANOVA	Sig	0.429	0.001	0.084	0.815	0.020	0.964	0.617	0.054	0.485	0.429	0.115	0.079	0.817	0.046
	Number Positive	No	0.56	0.56	0.56	0.56	0.57	0.56	0.56	0.56	0.56	0.56	0.57	0.56	0.56	0.56
	Attitudes	Yes	0.51	0.53	0.61	0.47	0.49	0.38	0.43	0.40	0.50	0.43	0.42	0.52	0.50	0.45
	ANOVA	Sig	0.020	0.312	0.380	0.025	0.007	0.001	0.089	0.020	0.291	0.020	0.000	0.557	0.332	0.050
	N. Negative	No	1.15	1.16	1.15	1.13	1.12	1.14	1.15	1.14	1.14	1.14	1.12	1.14	1.14	1.13
	Attitudes	Yes	1.08	1.05	0.98	1.38	1.33	1.40	1.29	1.34	1.37	1.40	1.52	1.43	1.47	1.52
	ANOVA	Sig	0.001	0.027	0.052	0.000	0.000	0.002	0.215	0.045	0.002	0.001	0.000	0.005	0.000	0.000
des	Number Trauma	No	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01
titu	Violence	Yes	0.04	0.01	0.03	0.04	0.02	0.02	0.01	0.02	0.03	0.02	0.03	0.04	0.04	0.04
Att	ANOVA	Sig	0.686	0.399	0.167	0.000	0.292	0.273	0.833	0.790	0.064	0.686	0.004	0.055	0.006	0.005
	Number positive	No	1.94	1.94	1.95	1.94	1.94	1.95	1.94	1.95	1.94	1.95	1.96	1.95	1.95	1.95
	doings	Yes	1.93	1.94	1.90	1.93	1.95	1.79	2.15	1.87	1.95	1.86	1.73	1.80	1.74	1.70
	ANOVA	Sig	0.451	0.909	0.699	0.893	0.914	0.197	0.203	0.602	0.947	0.451	0.004	0.322	0.099	0.031
	Number	No	0.60	0.60	0.60	0.59	0.59	0.60	0.60	0.60	0.60	0.60	0.59	0.60	0.60	0.60
	negative doings	Yes	0.71	0.59	0.56	0.71	0.69	0.58	0.64	0.70	0.74	0.71	0.74	0.79	0.77	0.75
	ANUVA	Sig	0.071	0.708	0.492	0.014	0.002	0.749	0.632	0.187	0.014	0.071	0.001	0.012	0.010	0.015





	EIGI	EN VALUES	Wilks	Lambda (31	df)	CENT	ROID	Test R	esults	Classification			
	Eigen value	Canonical Correlation	Wilks Lambda	Chi- Square	Sig	Centroid No	Centroid Yes	Box M	Sig	% Correct No	% Correct Yes	% Correct original group	
Diabetes	.68	.261	.937	58.044	.002	038	1.773	.059	.810	82.7	78.9	82.6	
Hypertension	.227	.430	.815	181.457	<.001	166	1.365	5.184	.023	79.7	74.5	79.2	
CVD	.125	.333	.889	104.759	<.001	.063	-1.979	6.673	.010	87.6	75.0	87.2	
Respiratory	.100	.302	.909	84.921	<.001	083	1.199	7.551	.006	82.4	66.1	81.4	
Allergies	.258	.453	.795	204.273	<.001	205	1.258	4.111	.043	82.4	71.7	80.9	
GID	.120	.327	.893	100.553	<.001	.066	-1.807	3.736	.055	86.1	71.9	85.6	
Fibromyalgia	.155	.366	.866	127.903	<.001	.056	-2.761	55.530	<.001	92.5	66.7	92.0	
Rheumatic Dis	.120	.327	.893	100.340	<.001	071	1.674	11.967	<.001	87.5	62.2	86.4	
Autoimmune	.113	.318	.899	94.911	<.001	081	1.388	14.545	<.001	84.4	60.0	83.0	
Chronic pain	.180	.391	.847	147.345	<.001	080	2.245	20.585	<.001	87.9	71.0	87.3	
Fatigue	.444	.554	.693	326.231	<.001	.197	-2.247	44.696	<.001	87.9	77.0	87.0	
Dermatologic	.077	.267	.929	65.847	<.001	038	2.003	.471	.497	84.9	82.4	84.9	
Tinnitus	.129	.338	.886	107.481	<.001	.065	-1.970	31.237	<.001	89.3	72.4	88.8	
Dizziness	.177	.388	0.849	145.050	<.001	.079	-2.235	56.313	<.001	90.4	67.7	89.6	

Chronic Pain	Fibromyalgia	AID	CVD	Fatigue	Diabetes	Dermatologic	Dizziness	Hypertension	Respiratory	Tinnitus	Allergies	GID	Rheumatic
N morbidities .867	N morbidities 821	N morbidities .787	N morbidities 612	N morbidities 916	Age .441	N morbidities .518	N morbidities 721	N morbidities .627	N morbidities .777	N morbidities 726	N morbidities .698	N morbidities 732	N morbidities .748
Sleep Quality 334	Sleep Quality 352	Work conflicts B Covid .248	Age454	Sleep Waking Quality .285	N morbidities .385	Work conflicts B Covid .382	Worries vs uncertainty 280	Age .583	Number negative doings .276	Number negative doings301	Age304	Sleep Waking Quality .222	
Sleep Waking Quality268	Sleep Waking Quality .330	Work physically heavy .248	Beer per Day 280	Morbidities worse231	BMI .328	N days of lockdown 366	Number negative doings255	BMI .368	Sleep Quality 244	Sleep Waking Quality .229	Sleep Quality 224		
How is your depression .244	Social Networks h/ day282	Morbidities worse .244	Work multitask B Covid .242	How is your depression 226	Sleep Waking Quality .312	Work harassment B Covid .340	Work conflicts B Covid249	Beer per Day .328	Sleep Waking Quality227	Work responsibilities B Covid .222			
Social Networks h/ day .240	Morbidities worse250	How is your anxiety .244	BMI228	Sleep Quality .221	How is your irritability 261	Number Negative Attitudes .215	How are your economic problems 238	How is frequency your sexual activity205	Number Negative Attitudes .209				
Morbidities Worse .237	TV h/day 225	N days of lockdown .243	How is your anxiety .216	How is your anxiety219	Work multitask B Covid259	Work physically heavy .205	How is your anxiety213						
How are your economic problems .236	Work stress B Covid208	Work stress interruptions B Covid .239	Work stress interruptions B Covid .215	How are your economic problems 203	Sleep Quality .254	Work multitask B Covid .202							
How is your anxiety .217	How are your economic problems 207	Sleep Waking Quality200	Work conflicts B Covid .211										
N days of lockdown .214	Work physically heavy200												
How are your worries vs uncertainty .212													
Number Negative Attitudes .206													

of TV dependence, hypertension lower levels of social networks dependence, while allergies had higher social networks dependence levels, AID higher levels of game dependence, hypertension had higher levels of alcoholic dependence while allergies and dizziness lower levels of alcoholic dependence. Positive attitudes were less frequent in Respiratory patients, allergies, GID, rheumatic, chronic pain, fatigue, dizziness. Negative attitudes were more frequent in Respiratory patients, allergies, GID, rheumatologic, AID, chronic pain, fatigue, dermatologic, tinnitus and dizziness, and less frequent in hypertension. Positive doings were lower in fatigue and dizziness. Negative doings were higher in Respiratory patients, allergies, AID, fatigue, dermatologic, tinnitus and dizziness.

Discriminant analysis showed consistent and significant differences for the chronic disorders and complaints evaluated (see table 5). Wilks Lambda was highly significant for all of them, the centroids for having (yes) and not having a disorder/complaint are quite different; for some disorders the "yes" centroid is negative (CVD, GID, fibromyalgia, fatigue, tinnitus and dizziness). The centroids biggest differences are for fibromyalgia, fatigue, chronic pain, dizziness, CVD and tinnitus (see Figure 1).

The test results evaluated by the Box M are significant for all situations, except for diabetes, GID and dermatologic disorders. The classification correctness was high in general with higher results for the No (not having the disorder/complaint) than for the Yes (having the disorder/complaint).

The pooled within-groups correlations between discriminating variables and standardized canonical discriminant functions were computed for each disorder/complaint; since degrees of freedom (df) are higher than 900, correlations higher than .200 were ranked and are presented in table 6. Chronic pain, Fibromyalgia, AID, Fatigue and CVD have the highest number of features with significant correlations.

It is also clear that the number of morbidities and morbidities worsening is very relevant is most disorders. Sleep quality and/ or sleep waking quality were quite significant in chronic pain, fibromyalgia, fatigue, AID, respiratory disorders, diabetes, tinnitus, allergies and GID. Items from concerning mental health (depression, anxiety, irritability and worries) were significant in all entities, except hypertension and allergies. Work stress before COVID was significant fibromyalgia, AID, CVD, diabetes, dermatologic, dizziness and tinnitus. Other factors such as economic problems, negative attitudes, screen time and beer consumption were specific of specific entities.

Discussion

The obtained results must be discussed in two different perspectives. One is the importance of a multidimensional approach and the associated relevance of specific features, which range from demographics to attitudes and from work characteristics before the pandemic to confinement and habits during the pandemic; the other relates to the differences between common chronic disorders in a stressful situation. Age is a significant factor: in most chronic disorder's patients tend to be older, except for those with allergies, respiratory disorders, fatigue and dermatologic disorders. In the LDA analysis, older age showed a strong correlation with the discriminant function for CVD, diabetes and hypertension, while younger age was similarly correlated for patients with allergies. BMI varied significantly across several conditions, being higher in diabetes, hypertension, CVD, and fibromyalgia, and or lower in allergies and gastrointestinal disorders. However, in the LDA analysis, BMI was only a significant factor for diabetes, hypertension and CVD.

The evaluation of work stress before COVID is relevant since, the presence of significant effects during the pandemic is a putative indicator of its long-term effects. Indeed, the several parameters of work stress (the subjective evaluation, the presence of significant interruptions and multitasking, conflicts, responsibilities together with physical and intellectual load, were quite frequent in most disorders and confirmed its relevance in LDA for fibromyalgia, AID, CVD, diabetes, dermatologic disorders, dizziness and tinnitus. Particular attention must be addressed to CVD and Dermatologic disorders since they share multitasking and work conflicts associated either with interruptions or an increased physical load. Furthermore, in the dermatologic group harassment is also present.

Work stress is worldwide quite prevalent; in 2019 (prior to the COVID pandemic) 38% of workers globally reported experiencing high daily stress [13]. Psychosocial factors are a considerable source of stress at work while having significant associations with depression [14].

The chronic cortisol excess associated with work stress and mediated by the stress level, may precede the development of an allostatic load, while explaining multiple health effects [15]. Data from 27 cohort studies in Europe, the USA and Japan suggests that work stressors, such as job strain and long working hours, are associated with a moderately elevated risk of incident coronary heart disease and stroke [16]. Work stress can indeed increase the risk of cardiovascular disease by 50%; the effort-reward imbalance workplace stress model explains the increased risk of CVD by its association with organic factors, such as, increased hypertension, intima media thickness and fibrinogen [17]. Job strain is also a risk factor for diabetes [16,18]. Stress, namely work bullying, excessive workload and low decision latitude are possible contributing factors in the development of fibromyalgia [19]. For our knowledge information concerning the prevalence of autoimmune disorders, tinnitus and dizziness in relation with work stress is not available.

The average number of comorbidities in the observed CMD was in all sub-groups significantly higher than in controls, ranging between 3.19 and 5.29; in practice this means a high prevalence of comorbidities, and is in line with the concept that chronic disorders are not only co-morbid but also syndemic, especially when this term extends from synergistic epidemics to synergistic multilevel environments contributing to negative heath impacts [20]. The existence of comorbidities in chronic disorders and their bidirectional mutual influence with potential increased severity of both is well known. In LDA the correlation values for morbidities were quite high, ranging from .518 in dermatologic disorders to -.916 in fatigue. Furthermore, morbidities worsening during COVID, showed significant correlations with the discriminant function in chronic pain, fibromyalgia, AID and fatigue. These results obtained during the COVID pandemic are in line with comorbidities relevance. Chronic pain is rather prevalent worldwide $(\sim 20\%)$ and its frequent comorbidities are anxiety, depression, cognitive impairments, sleep disorders, migraine, diabetes type II, hypertension, CVD, fatigue, etc. [21]. Fibromyalgia common morbidities are irritable bowel syndrome, headache, paraesthesia's, depression, anxiety, obesity and multiple skin disorders (psoriasis, chronic urticaria, contact allergy, acneiform disorders, hidradenitis suppurativa, and vitiligo) [22].

AID include an extensive spectrum of diseases, since from to brain to the blood and skin, auto immunity aggression might occur. Immunodeficiency comorbidities and complications include infection, chronic lung disease, granulomatous lymphocytic interstitial lung disease, and autoimmune disorders [23]; sleep disturbances and mental disorders are frequently comorbid manifestations. AID, despite the marked diversity share common haplotypes and gut dysbiosis is, among them, a common comorbidity, occurring long before autoimmunity even begins [24].

Low vision, diabetes mellitus, back/neck problems, osteoarthritis, chronic obstructive pulmonary disease (COPD), and cancer are the most prevalent comorbid conditions in the different CVDs [25,26]. In diabetes the most common comorbid disorders are dyslipidaemia, hypertension, Non Alcoholic Fatty Liver Disease, Obesity, Polycystic ovary syndrome and Sleep apnea [27]. Respiratory diseases have as frequent comorbidities obesity, gastroesophageal reflux disease, and allergic rhinitis, sleep disorders, cardiovascular disorders and cancer [28]. For tinnitus again sleep and mental disorders are frequently comorbid [29].

The number of days in lock down was quite similar in the several groups; differences occurred in Rheumatic disorders (loner days in lockdown) and in dermatologic patients (shorter lockdown). The number of days in lockdown had, however, low but significant, canonical correlations with the discriminant function in Chronic pain and AID and dermatologic disorders. The subjective way people was living in lockdown was always slightly worse in the chronic disorder's patients with significant differences in diabetes, respiratory and allergic subgroups, AID, chronic pain and fatigue, tinnitus and dizziness; despite these differences they did not emerge as relevant in discriminant analysis.

Depression, anxiety, irritability and worries were frequently higher in CMD patients. In some disorders all of them (diabetes, chronic pain, fatigue, tinnitus and dizziness), in some others, none of them, such as CVD. However, in discriminant analysis mental health components showed significant correlations with the discriminant function in chronic pain, AID, fatigue, CVD, diabetes and dizziness. These results are in line with two known aspects of mental health, especially in what concerns anxiety and depression: it deteriorates in chronic disorders, and it was negatively impacted during COVID; according to the WHO the COVID-19 pandemic triggered a 25% increase in prevalence of anxiety and depression worldwide [30].

Economic problems were relevant in most disorders and emerged with significant correlations in LDA in chronic pain, fibromyalgia, fatigue and dizziness. According to the World Bank the pandemic introduced "shock waves through the world economy and triggered the largest global economic crisis in more than a century. Studies based on precrisis data suggest, for example, that more than 50 percent of households in emerging and advanced economies were not able to sustain basic consumption for more than three months in the event of income losses. Similarly, the average business could cover fewer than 55 days of expenses with cash reserves" [31]. These negative impacts were strongly felt in Portugal with the EU commission stating that Portuguese GDP was estimated to have fallen by 7.6% in 2020 [32]. Therefore, citizens' economic problems were expected, but it must be stressed that the most affected were those who complain with pain, fatigue and dizziness, often considered functional symptoms.

The average sleep latency was higher than 30 minutes in both groups and even slightly higher in non-patients; average number of awakenings was close to 3 per night and sleep duration in weekdays was around 6.5 h and a bit higher in weekends 7.5h. This may reflect the global sleep difficulties of the Portuguese adult population with high prevalence of poor sleep [33], short sleep [34], Insomnia [35,36], sleep inducing medication [37,38] and pain-related insomnia [39]. Despite the detailed information concerning sleep habits (into bed and out of bed during weekdays and weekends) and sleep parameters (latency, awakenings and duration) only the quality of both Sleep and Wakening provided significant differences.

COVID impact upon Sleep and Wakening quality of the general population has been described [10]. In all chronic disorders, with exception of CVD and hypertension, Sleep and Wakening quality deteriorated during COVID. One or both, have significant correlations with the discriminant function in Chronic pain, Fibromyalgia, AID, Fatigue, Diabetes, Respiratory diseases, Tinnitus, Allergies and GID.

In Brazil, during the pandemic, there was decreased physical activity, and an increase in symptoms of anxiety and depression and screen time [40]. In Canada avoiding excessive screen time and engaging in exercise, particularly outdoors, were important behaviours associated with better perceived mental and general health during the COVID-19 [41]. In Portugal data concerning physical activity during COVID showed some singularities; it was practised more by males than females, elderly practice more often and more outdoors than younger subjects, those that practise PA between 4 and 10h per week had better pandemic compliance; those with low PA levels had poorer nutrition habits and longer

screen time [42]. A systematic review and metanalysis confirmed an increase screen time during the pandemic in all ages, from children to adults; the associated correlates included adverse dietary behaviours, sleep, mental health, parental health, and eye health [43]. In Portugal identical results were observed in adults in what concerns nutrition, sleep and mental health [44]. Despite both Physical activity and Nutrition being considered, together with Sleep, the pillars of health, they did not differentiate the subgroups of CMD in our sample. Physical activity was not different in the subgroups evaluated. Nutrition was only different in Allergies. No correlations were found in LDA.

Patients with CVD, hypertension and respiratory disorders had a higher cigarette consumption; most patients with CMD had a lower consumption of wine and only beer per day emerged with significant correlations in LDA, negative in CVD and positive in Hypertension. Screen time, mainly for social networks, was higher in several CMD subgroups (Diabetes, Fibromyalgia, Chronic Pain, Fatigue and Dizziness) and they had significant correlations for Chronic Pain (+) and Fibromyalgia (-). The number of Positive attitudes was lower in several patients' groups, and the number of negative attitudes was significantly lower in many subgroups with exception of diabetes and hypertension, in which it was lower and CVD with no differences. Trauma and violence were significant in respiratory diseases, fatigue, tinnitus and dizziness. Positive doings were lower in fatigue and dizziness, and negative doings were higher in respiratory, allergies, autoimmune, fatigue, dermatologic, tinnitus and dizziness. LDA confirmed in importance of negative attitudes and doings in respiratory, dermatologic, tinnitus. The study limitations concern the subjective not confirmed responses related to internet surveys, but this drawback is compensated by the high number of responses over the entire territory. Another limitation relates to the non-inclusion of gender in discriminant analysis since LDA requires gaussian distributions.

Conclusions

The high prevalence of chronic medical disorders, their association with multiple mental and physical comorbidities, and the consequent health risks and long-term treatment difficulties contribute to their high burden in multiple dimensions: individual, societal, healthcare and economic.

During the pandemic chronic medical disorders were particularly affected. Portugal besides having a higher prevalence of chronic medical disorders, is among the countries which used more strict protective pandemic measures with subsequent quite negative socioeconomic impacts. This country is therefore a good model for studying the differences and similarities of chronic disorders during negative environmental periods.

An ecologic multilevel model of health and disease was used both to collect and to analyse the data [45]. LDA was used to identify differences between 14 chronic disorders. Correct classification of the CMD was achieved for each of them, with percentages ranging from 79.2% in Hypertension to 92% in Fibromyalgia; in all subgroups Lamba Wilks was highly significant; Box M test results were only not significant for diabetes and GID. Comorbidities, Sleep, Mental Health, Work stress, and Attitudes emerged as the most relevant features in CMD. Two complaints' subgroups, Tinnitus and Dizziness, often discarded due to their non-specificity, presented important relevance among the studied disorders. Future studies are still needed and Recommendations for detailed attention in Public Health services proposed. Two others, CVD and Hypertension, presented some specificities, with higher prevalence of toxics consumption, cigarettes, beer, wine, alcoholic beverages and alcohol dependence, no differences in sleep, in mental symptoms and less work stress before COVID. Taking in consideration the high associated health risks of both specific recommendations dealing with toxic habits is proposed in health literacy programs. Chronic Pain, Fibromyalgia, Autoimmune disorders, Fatigue and CVD had the highest numbers of significant correlations with the discriminant function. It must be stressed that the first four types of disorders are usually associated with subjective symptoms and/or complex clinical presentations. Consequently, these disorders deserve special care in environmental stressing situations. Altogether each CMD deserves a tailored attention, considering and caring the associated specificities.

Acknowledgments

Helena Canhão, Margarida Gaspar de Matos, Katie Almondes, Amélia Feliciano, Maria Raquel Silva, Tânia Gaspar, Conceição Pereira, Alexandra Carreiro, Aurora Lino, Susana Moreira, Ana Bernarda, Susana Gaspar, Lúcia Ramiro, Júlio Fonseca, Gabriela Videira, Maria Augusta Machado, Carla Bentes, Francisco Sampaio, Lara Guedes Pinho, Joana Pimentel, Ana Bernarda, Joana Vaz de Castro, Sofia Rebocho, Gina Tomé, Cátia Branquinho, Eleni Araújo, Hugo Canas-Simião, João Peixoto, Fábio Guedes, Richard Staats.

Endorsements/Dissemination: Ordem dos Médicos, Ordem dos Enfermeiros, Ordem dos Psicólogos Portugueses, European Sleep Research Society (ESRS), Associação Portuguesa de Sono (APS), Sociedade Portuguesa de Neurologia e Ordem dos Farmacêuticos.

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