

ORIGINAL RESEARCH

Fibromyalgia: a new facet of the post-COVID-19 syndrome spectrum? Results from a web-based survey

Francesco Ursini ^{1,2}, Jacopo Ciaffi,¹ Luana Mancarella,¹ Lucia Lisi,¹ Veronica Brusi,¹ Carlotta Cavallari,¹ Martina D'Onghia,¹ Anna Mari,¹ Elena Borlandelli,³ Jacopo Faranda Cordella,⁴ Micaela La Regina,⁵ Pasquale Viola,⁶ Piero Ruscitti,⁷ Marco Miceli,⁸ Roberto De Giorgio,⁹ Nicola Baldini,¹⁰ Claudio Borghi,¹¹ Alessandro Gasbarrini,¹² Annamaria Iagnocco,¹³ Roberto Giacomelli,¹⁴ Cesare Faldini,¹⁵ Maria Paola Landini,¹⁶ Riccardo Meliconi ^{1,17}

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For numbered affiliations see end of article.

Correspondence to

Professor Francesco Ursini; francesco.ursini2@unibo.it

ABSTRACT

Objective Postacute COVID-19 syndrome (PACS) is an emerging entity characterised by a large array of manifestations, including musculoskeletal complaints, fatigue and cognitive or sleep disturbances. Since similar symptoms are present also in patients with fibromyalgia (FM), we decided to perform a web-based cross-sectional survey aimed at investigating the prevalence and predictors of FM in patients who recovered from COVID-19.

Methods Data were anonymously collected between 5 and 18 April 2021. The collection form consisted of 28 questions gathering demographic information, features and duration of acute COVID-19, comorbid diseases, and other individual's attributes such as height and weight. The American College of Rheumatology (ACR) Survey Criteria and the Italian version of the Fibromyalgia Impact Questionnaire completed the survey.

Results A final sample of 616 individuals (77.4% women) filled the form 6±3 months after the COVID-19 diagnosis. Of these, 189 (30.7%) satisfied the ACR survey criteria for FM (56.6% women). A multivariate logistic regression model including demographic and clinical factors showed that male gender (OR: 9.95, 95% CI 6.02 to 16.43, $p<0.0001$) and obesity (OR: 41.20, 95% CI 18.00 to 98.88, $p<0.0001$) were the strongest predictors of being classified as having post-COVID-19 FM. Hospital admission rate was significantly higher in men (15.8% vs 9.2%, $p=0.001$) and obese (19.2 vs 10.8%, $p=0.016$) respondents.

Conclusion Our data suggest that clinical features of FM are common in patients who recovered from COVID-19 and that obesity and male gender affect the risk of developing post-COVID-19 FM.

INTRODUCTION

Since its first appearance in December 2019, SARS-CoV-2—the pathogen responsible for COVID-19—exhibited all its devastating potential,¹ causing more than three million deaths

Key messages**What is already known about this subject?**

- Postacute COVID-19 syndrome (PACS) is emerging as a complex condition with a wide range of clinical manifestations.
- Clinical features of PACS include musculoskeletal pain, fatigue, cognitive impairment and sleep disturbances.

What does this study add?

- Our study suggests that up to 30% of patients with PACS may satisfy criteria for fibromyalgia (FM).
- Obesity and male gender represent the strongest risk factors for post-COVID-19 FM.

How might this impact on clinical practice or further developments?

- It is reasonable to expect that rheumatologists will soon face up with a sharp rise of cases of this new entity that we defined 'FibroCOVID'.

worldwide. Apart from the clinical manifestations of the acute disease, the long-term consequences of COVID-19 are emerging as a new, overwhelming challenge for clinicians and healthcare systems. A postacute COVID-19 syndrome (PACS)² is now clearly recognised and, in the near future, is expected to impose a serious burden on different medical specialties, given the pleiotropic nature of its clinical manifestations. Of note, musculoskeletal pain—the cardinal symptoms of fibromyalgia (FM), reported in one-third of patients with acute COVID-19³—is part of the complex spectrum of PACS, along with pulmonary, cardiovascular, haematological, renal, gastroenteric, dermatological, endocrine and neuropsychiatric sequelae.²

The diagnosis of FM historically relied on the 1990 American College of Rheumatology (ACR) criteria,⁴ including widespread pain of at least 3 months' duration and tenderness on pressure at 11 or more of 18 specific tender points. In 2010, the ACR proposed a new set of clinical criteria for the diagnosis of FM based on a Widespread Pain Index (WPI) and a Symptom Severity (SS) scale; however, the tender point examination was withdrawn.⁵ The 2010 criteria underwent a revision in 2016⁶ that combined physician and questionnaire criteria and eliminated the previous recommendation regarding diagnostic exclusions. Furthermore, the ACR 2010 criteria have also been adapted for administration as a self-report questionnaire ('survey' criteria) to be used in epidemiological studies with good reliability and convergent and discriminant validity.⁷

The pathogenesis of FM is still far to be fully understood. Pain augmentation/dysperception seems associated with exquisite neuromorphological modifications and imbalance between pronociceptive and antinociceptive pathways arising from an intricate interplay between genetic predisposition, stressful life events, psychological characteristics and emerging peripheral mechanisms, such as small fibre neuropathy or neuroinflammation.⁸ Strikingly, a role for infectious triggers—viral infections, in particular—has been also postulated.⁹

Internet-based surveys have gained growing popularity in the past years among healthcare researchers because of their clear advantages, such as the ability to reach a large pool of potential participants within a short period of time and to involve subjects that may be geographically dispersed or otherwise may be difficult to access; this is in conjugation with other practical reasons, such as the inexpensiveness and the easiness of data extraction, management and analysis.¹⁰ The emergence of COVID-19 even emphasised the use of web-based surveys, and more than 2000 records can now be retrieved on PubMed when applying the search string "COVID-19" AND "online survey".

On this basis, here we report the results of a web-based survey aimed at investigating the prevalence of FM developed after symptomatic COVID-19; the secondary aim was to investigate predictive factors of post-COVID-19 FM syndrome development.

METHODS

Design of the study

The present study was carried out as a web-based cross-sectional survey. Data were collected between 5 and 18 April 2021 through an online form built using the Google Forms platform.¹¹ Google Forms is a free survey administration tool that has been largely used in medical research.¹¹ Reporting was compliant with the Checklist for Reporting Results of Internet E-Survey.¹²

FM classification and assessment

To define the presence of FM in survey respondents, the ACR survey criteria¹³—developed as a modification of the ACR 2010 criteria to be used as a self-administration tool—were applied after linguistic validation as detailed in online supplemental methods. FM survey criteria have been successfully applied in web-based survey research.¹⁴ A Fibromyalgiasness Scale or Fibromyalgia Symptom Scale (FS) is obtained by summing up the modified WPI and SS scores. An FS score of ≥ 13 has been largely adopted as the best cut-off for FM classification.¹³ The FS can be also used as a continuous measure of symptom burden.¹³ To quantify FM severity, the Italian version of the Fibromyalgia Impact Questionnaire (FIQ-I)¹⁵ was used. FIQ-I was modified by excluding one question (item 3, 'work missed', and item 4, 'do work') expressly recalling a past FM diagnosis. The overall score was adjusted to account for the reduced number of questions according to the suggestion for managing non-responses to individual questions.

Survey development

A group of senior researchers (FU, RM and LM), including a medical psychotherapist (LL), designed the survey draft. The content was further reviewed by all study researchers. Pilot testing investigating the understandability of questions and technical usability was performed on a pool of healthy individuals ($n=20$) and patients with post-COVID-19 ($n=20$) who did not participate in developing the survey. The final version was consequently modified following their suggestions and was approved by consensus. The survey was found to require a total of 10min to be completed. Results were transmitted to the database only if the participant clicked on 'survey completed' at the end of the questionnaire. Questions were listed in the same order for all participants. The survey could not be submitted unless all mandatory questions were completed.

Survey structure

The survey consisted of a single page including a total of 28 questions. Questions were preceded by a preface stating the overall goal of the survey, information on how to contact the research group and that collected data could be anonymously used for research purposes and publication. Details of the survey structure are reported in online supplemental table S1.

Briefly, a first part of the survey (Q1–Q14) was used to collect demographic characteristics, marital and occupational status, symptoms and duration of acute COVID-19, comorbid diseases and other individual characteristics such as height and weight.

A second part of the survey (Q15–Q19) was dedicated to the ACR Survey Criteria for FM.¹³ Finally, the third and last parts of the survey (Q20–Q28) contained the FIQ-I.¹⁵

Target population and survey administration

The target population comprised adult individuals (≥ 18 years) who developed COVID-19 3 or more months before the survey publication. To reach this population,

Table 1 General characteristics of the study population

	Overall (n=616)	Weight- adjusted (n*=591)
Age (years)	45±12	45±12
Female gender, n (%)	477 (77.4)	–
Marital status		
Single, n (%)	116 (18.8)	126 (21.3)
Married, n (%)	437 (70.9)	417 (70.6)
Separated, n (%)	51 (8.3)	39 (6.6)
Widowed, n (%)	12 (1.9)	9 (1.6)
Employment status		
Student, n (%)	29 (4.7)	35 (5.9)
Employed, n (%)	494 (80.2)	469 (79.4)
Unemployed, n (%)	66 (10.7)	56 (9.4)
Retired, n (%)	27 (4.4)	31 (5.3)
COVID-19 symptoms		
Fever, n (%)	429 (69.6)	421 (71.3)
Cough, n (%)	292 (47.4)	282 (47.7)
Dyspnoea, n (%)	237 (38.5)	225 (38.1)
Headache, n (%)	390 (63.3)	352 (59.5)
Myalgia, n (%)	436 (70.8)	398 (67.3)
Arthralgia, n (%)	404 (65.6)	358 (60.6)
Anosmia or ageusia, n (%)	437 (70.9)	399 (67.4)
Abdominal pain, n (%)	229 (37.2)	198 (33.5)
Treatment setting		
No treatment, n (%)	165 (26.8)	165 (27.9)
Home treatment, n (%)	375 (60.9)	342 (57.8)
Hospital admission, n (%)	66 (10.7)	73 (12.3)
ICU admission, n (%)	10 (1.6)	12 (2.0)
COVID-19 duration (days)	13 (7–20)	10 (7–20)
COVID-19 treatment		
Analgesics/NSAIDs, n (%)	422 (68.5)	401 (67.8)
LMWH, n (%)	131 (21.3)	132 (22.3)
Hydroxychloroquine, n (%)	53 (8.6)	48 (8.2)
Supplemental oxygen, n (%)	66 (10.7)	69 (11.6)
Pre-existent comorbid diseases		
Rheumatoid arthritis or other inflammatory arthritides, n (%)	30 (4.9)	24 (4.0)
Connective tissue diseases, n (%)	3 (0.5)	2 (0.3)
Anxiety, n (%)	108 (17.5)	104 (17.7)
Depression, n (%)	30 (5.8)	37 (6.3)
Diabetes, n (%)	10 (1.6)	9 (1.6)
High blood pressure, n (%)	97 (15.7)	101 (17.1)
Chronic pulmonary diseases, n (%)	52 (8.4)	46 (7.8)
History of myocardial infarction, n (%)	3 (0.5)	5 (0.8)
History of stroke, n (%)	2 (0.3)	1 (0.2)
Other neurological diseases, n (%)	3 (0.5)	2 (0.3)
Malignancy, n (%)	5 (0.8)	5 (0.8)

Continued

Table 1 Continued

	Overall (n=616)	Weight- adjusted (n*=591)
BMI (kg/m ²)	25.3±4.8	25.6±4.7
BMI category		
Underweight, n (%)	19 (3.1)	14 (2.3)
Normal weight, n (%)	335 (54.4)	302 (51.1)
Overweight, n (%)	160 (26.0)	172 (29.1)
Obese, n (%)	102 (16.6)	103 (17.5)
FM (FS score ≥13), n (%)	189 (30.7)	234 (39.5)
Additional symptoms		
Constipation, n (%)	68 (11)	71 (12.1)
Diarrhoea, n (%)	69 (11.2)	76 (12.9)
Nausea, n (%)	61 (9.9)	61 (10.4)
Heartburn, n (%)	90 (14.6)	105 (17.7)
Pain in upper abdomen, n (%)	108 (17.5)	127 (21.5)
Numbness and tingling in arms, n (%)	228 (37)	241 (40.7)
Dizziness or vertigo, n (%)	177 (27.1)	183 (31.0)
Insomnia, n (%)	199 (32.3)	223 (37.7)
Chest pain, n (%)	121 (19.6)	130 (22.0)
Blurred vision, n (%)	163 (26.5)	175 (29.7)
Fever, n (%)	151 (24.5)	158 (26.7)
Dry eyes, n (%)	154 (25.0)	163 (27.5)
Diffuse itching, n (%)	124 (20.1)	128 (21.7)
Excessive sweating, n (%)	108 (17.5)	116 (19.7)
Ringing in ear, n (%)	123 (20.0)	126 (21.4)
Change in smell and/or taste, n (%)	157 (25.5)	167 (28.3)
Shortness of breath, n (%)	270 (43.8)	284 (48.1)
Loss of appetite, n (%)	57 (9.3)	65 (11.1)
Hair loss, n (%)	156 (25.3)	165 (28)
Frequent or painful urination, n (%)	98 (15.9)	103 (17.5)

*Corresponding to the imputed total number of respondents calculated on the basis of gender-based weights.

BMI, body mass index; FM, fibromyalgia; FS, Fibromyalgia Symptom Scale; ICU, intensive care unit; LMWH, low-molecular-weight heparin; NSAIDs, non-steroidal anti-inflammatory drug.

members of the research group combined several lines of contact, mainly based on social network (Facebook and Instagram) interactions as detailed in online supplemental methods. No monetary or non-monetary incentives were offered for the voluntary completion of the survey. One follow-up reminder message was sent 1 week apart to all direct contacts; however, participants were explicitly asked to answer the survey only once.

Ethical considerations

By voluntarily taking part in the survey, each participant explicitly authorised the use of the anonymous data recorded in the questionnaire for research purposes and their publication, as clearly stated in the questionnaire preface.¹⁶

Table 2 Comparison between respondents with and without FM

	FM (n=189)	No FM (n=427)	P value	Weighted P value
Age (years)	46±11	44±12	0.312	0.04
Female gender, n (%)	107 (56.6)	370 (86.7)	<0.0001	–
Marital status				
Single, n (%)	37 (19.6)	79 (18.5)	0.739	1.000
Married, n (%)	137 (72.5)	300 (70.3)	0.631	0.519
Separated, n (%)	12 (6.3)	39 (9.1)	0.271	0.848
Widowed, n (%)	3 (1.6)	9 (2.1)	1.000	1.000
Employment status				
Student, n (%)	4 (2.1)	25 (5.9)	0.061	0.019
Employed, n (%)	158 (83.6)	336 (78.7)	0.188	0.047
Unemployed, n (%)	17 (9.0)	49 (11.5)	0.399	0.059
Retired, n (%)	10 (5.3)	17 (4.0)	0.523	0.187
COVID-19 symptoms				
Fever, n (%)	128 (67.7)	301 (70.5)	0.491	0.746
Cough, n (%)	100 (52.9)	192 (45.0)	0.069	0.046
Dyspnoea, n (%)	86 (45.5)	151 (35.4)	0.017	0.001
Headache, n (%)	112 (59.3)	278 (65.1)	0.165	0.108
Myalgia, n (%)	133 (70.4)	303 (71.0)	0.882	0.749
Arthralgia, n (%)	122 (64.6)	282 (66.0)	0.719	0.246
Anosmia or ageusia, n (%)	134 (70.9)	303 (71.0)	0.988	0.352
Abdominal pain, n (%)	71 (37.6)	158 (37.1)	0.910	0.696
Treatment setting				
No treatment, n (%)	45 (23.8)	120 (28.1)	0.279	0.019
Home treatment, n (%)	103 (54.5)	272 (63.7)	0.032	0.233
Hospital admission, n (%)	36 (19.0)	30 (7.0)	<0.0001	<0.0001
ICU admission, n (%)	5 (2.6)	5 (1.2)	0.185	0.234
COVID-19 duration (days)	14 (7–21)	12 (7–20)	0.424	0.820
COVID-19 treatment				
Analgesics/NSAIDs, n (%)	131 (69.3)	291 (68.1)	0.755	0.364
LMWH, n (%)	58 (30.7)	73 (17.1)	<0.0001	<0.0001
Hydroxychloroquine, n (%)	22 (11.6)	31 (7.3)	0.074	0.029
Supplemental oxygen, n (%)	34 (18)	32 (7.5)	<0.0001	<0.0001
Pre-existent comorbid diseases				
Rheumatoid arthritis or other inflammatory arthritides, n (%)	7 (3.7)	23 (5.4)	0.371	0.294
Connective tissue diseases, n (%)	0 (0)	3 (0.7)	0.248	0.251
Anxiety, n (%)	37 (19.6)	71 (16.6)	0.375	0.198
Depression, n (%)	11 (5.8)	25 (5.9)	0.986	0.414
Diabetes, n (%)	3 (1.6)	7 (1.6)	0.962	0.706
High blood pressure, n (%)	51 (27)	46 (10.8)	<0.0001	<0.0001
Chronic pulmonary diseases, n (%)	16 (8.5)	36 (8.4)	0.989	0.575
History of myocardial infarction, n (%)	1 (0.5)	2 (0.5)	0.920	0.983
History of stroke, n (%)	0 (0)	2 (0.5)	0.346	0.418
Other neurological diseases, n (%)	1 (0.5)	2 (0.5)	0.920	0.763

Continued

Table 2 Continued

	FM (n=189)	No FM (n=427)	P value	Weighted P value
Malignancy, n (%)	2 (1.1)	3 (0.7)	0.650	0.349
BMI (kg/m ²)	30.4±4.4	23±2.9	<0.0001	<0.0001
BMI category				
Underweight, n (%)	0 (0)	19 (4.4)	0.001	0.001
Normal weight, n (%)	19 (10.1)	316 (74)	<0.0001	<0.0001
Overweight, n (%)	77 (40.7)	83 (19.4)	<0.0001	<0.0001
Obese, n (%)	93 (49.2)	9 (2.1)	<0.0001	<0.0001

P values <0.05 are highlighted in bold.

BMI, body mass index; FM, fibromyalgia; ICU, intensive care unit; LMWH, low-molecular-weight heparin; NSAID, non-steroidal anti-inflammatory drug.

Statistical analysis

A sample size of at least 457 patients was needed to estimate a prevalence of 2%–5% with precision set at 0.02 and confidence level set at 0.95. A sample size of at least 385 patients was needed to estimate a prevalence of 10%–50% with precision set at 0.05 and confidence level set at 0.95.

Data are expressed as mean±SD, median (25th–75th percentiles) or number (percentage) as appropriate.

Poststratification weighting was used to adjust for self-selection bias as previously suggested.¹⁷ Weighting is a family of techniques that allow improvement of the accuracy of survey estimates by using auxiliary information, that is, a set of variables (eg, age and gender) that have been measured in the survey, and for which the value for the population is available. By comparing the response distribution of an auxiliary variable, it can be extrapolated whether or not the sample is representative for the whole population. If these distributions differ considerably, adjustment weights are computed and assigned to all records in order to align the representation of various subpopulation groups to match that of the known population. The weight is calculated as the ratio between the population (N) and the sample (n) proportion for the auxiliary variable: $W=N/n$. The weight variable is created for each record of the data table and applied in analysis using the ‘weight cases’ function of the statistical analysis software.

Table 3 Univariate correlation between FS score and other continuous variables with poststratification weights applied

	FS score	
	R	P value
Age (years)	0.028	0.742
BMI (kg/m ²)	0.763	<0.0001
Time since COVID-19 (months)	0.068	0.423
In-COVID-19 duration (days)	0.054	0.523

P values <0.05 are highlighted in bold.

BMI, body mass index; FS, Fibromyalgia Symptom Scale.

Student’s t-test was used for comparing means of continuous variables between groups; highly skewed variables were ln-transformed before the analysis. Fisher’s exact test was used to compare categorical variables. Univariate and multivariate logistic regression models were built to assess the predictivity of continuous or categorical variables for a dichotomic dependent variable, expressed as OR and 95% CI.

All tests were two tailed. Analyses were performed using the Statistic Package for Social Sciences software V.23 (IBM).

RESULTS

General features of the survey respondents

A total of 937 individuals (76.7% women) completed the survey form. Of these, 321 were excluded from the analysis for different reasons: 37 did not report a diagnosis of COVID-19 confirmed by a physician; 61 did not perform a nasopharyngeal swab or reported a negative result; 23 had a pre-existent diagnosis of FM; 12 declared a history of chronic musculoskeletal pain and 188 did not meet the symptom duration criteria (≥3 months) for FM classification. The final cohort comprised 616 patients. General characteristics of the population are reported in [table 1](#). Most patients (77.4%) were women with a mean age of 45±12 years; median COVID-19 duration was 13 days with 10.7% of patients requiring hospital admission. The most common pre-existent comorbid diseases were anxiety (17.5%), obesity (16.6%), high blood pressure (15.7%), chronic pulmonary diseases (8.4%), depression (5.8%) and inflammatory arthritides (4.9%). Comparison between our cohort and official data released from the Italian Ministry of Health¹⁸ describing the cumulative Italian population of patients with COVID-19 showed a major difference in gender distribution (women: 77.4% vs 51.1%); to account for this potential source of self-selection bias, poststratification weights were assigned to each gender (weight for female gender=0.66, weight for male gender=1.99), and weighted values for all variables were calculated ([table 1](#)).

Table 4 Univariate and multivariate logistic regression analyses with poststratification weights applied

	Univariate OR	P value	Multivariate OR	P value
Age	1.015 (1.001 to 1.029)	0.036	1.006 (0.986 to 1.026)	0.584
Male gender	4.975 (3.332 to 7.426)	<0.0001	9.951 (6.025 to 16.435)	<0.0001
COVID-19 duration	1.000 (0.991 to 1.009)	0.962	–	–
Had COVID-19 during the first wave	1.537 (0.944 to 2.502)	0.084	1.126 (0.523 to 2.426)	0.762
Had cough	1.397 (1.003 to 1.945)	0.048	1.090 (0.677 to 1.755)	0.723
Had dyspnoea	1.786 (1.272 to 2.506)	0.001	1.627 (0.922 to 2.870)	0.093
Had myalgia	0.944 (0.665 to 1.341)	0.749	–	–
Had arthralgia	0.822 (0.587 to 1.151)	0.254	–	–
Had anosmia	0.842 (0.593 to 1.195)	0.336	–	–
Intensity of COVID-19 treatment setting	1.727 (1.344 to 2.218)	<0.0001	1.548 (0.937 to 2.58)	0.088
Treated with analgesics/NSAIDs	1.182 (0.828 to 1.686)	0.357	0.983 (0.592 to 1.631)	0.947
Treated with LMWH	2.180 (1.472 to 3.229)	<0.0001	1.897 (0.944 to 3.812)	0.072
Treated with HCQ	1.969 (1.088 to 3.562)	0.025	1.313 (0.464 to 3.713)	0.607
Treated with supplemental oxygen	2.531 (1.514 to 4.229)	<0.0001	0.482 (0.181 to 1.283)	0.144
Pre-existent arthritis	0.664 (0.273 to 1.614)	0.366	–	–
Pre-existent anxiety	1.314 (0.858 to 2.013)		–	–
Pre-existent depression	1.255 (0.642 to 2.452)	0.507	–	–
Pre-existent high blood pressure	3.061 (1.964 to 4.770)	<0.0001	1.511 (0.781 to 2.925)	0.220
Pre-existent diabetes	0.849 (0.219 to 3.290)	0.813	–	–
Pre-existent pulmonary disease	1.164 (0.634 to 1.164)	0.624	–	–
Pre-existent obesity	41.192 (18.003 to 98.879)	<0.0001	82.823 (32.192 to 213.084)	<0.0001

P values <0.05 are highlighted in bold.

HCQ, hydroxychloroquine; LMWH, low-molecular-weight heparin; NSAIDs, Nonsteroidal anti-inflammatory drugs.

Prevalence of FM after COVID-19 and comparison between respondents with FM and without FM

A total of 189 individuals (30.7%) fulfilled the criteria for FM classification after an average of 6±3 months from COVID-19 diagnosis (table 1). Of these, a total of 79 patients have contracted COVID-19 during the first pandemic wave (February–April 2020) and 491 during the second wave (October 2020–January 2021); prevalence of FM was 39.2% (31 cases) and 28.9% (142 cases), respectively (p=0.066). The remaining 46 got COVID-19 in between the two waves.

Respondents with FM were predominantly women (56.6%), admitted to hospital more frequently than counterparts without FM (19.0% vs 7.0%, p<0.0001) and reported significantly higher proportions of cough (52.9% vs 45.0%, p=0.046) and dyspnoea (45.5% vs 35.4%, p=0.017) during acute COVID-19 (table 2). Accordingly, a higher proportion of patients with FM were treated with supplemental oxygen (18.0% vs 7.5%, p<0.0001). The body mass index (BMI) was significantly higher in patients with FM (30.4±4.4 kg/m² vs 23.0±2.9 kg/m², p<0.0001) as well as the proportion of obese individuals (49.2% vs 2.1%, p<0.0001). Furthermore, among self-reported pre-existent comorbidities, high blood pressure was significantly more common in individuals with FM (27.0% vs 10.8%, p<0.0001).

To explore the possible role of COVID-19 severity in FM symptom burden, we compared the Fibromyalgia Impact Questionnaire (FIQ) scores obtained from patients with FM who were admitted to the hospital versus those who were not and from patients treated with supplemental oxygen versus those who were not (online supplemental table S2). No significant differences, except for a slightly lower FIQ–stiffness score in patients treated with oxygen were observed.

Predictors of FM after COVID-19

In an attempt to identify factors associated with FM development, we performed correlation and regression analyses to ascertain potential predictors among demographic, anthropometric and COVID-19-related variables. After gender-based poststratification weighting, the FS score (table 3) was positively correlated with BMI (R=0.763, p<0.0001).

Further, variables obtaining a p value of <0.10 in the comparative analysis reported in table 2 were entered in univariate and multivariate logistic regression models with poststratification weights applied. Results of the logistic regression analyses are shown in table 4. Age (OR: 1.015, 95% CI 1.001 to 1.029, p=0.036), male gender (OR: 4.975, 95% CI 3.332 to 7.426, p<0.0001), cough (OR: 1.397, 95% CI 1.003 to 1.945, p=0.048), dyspnoea

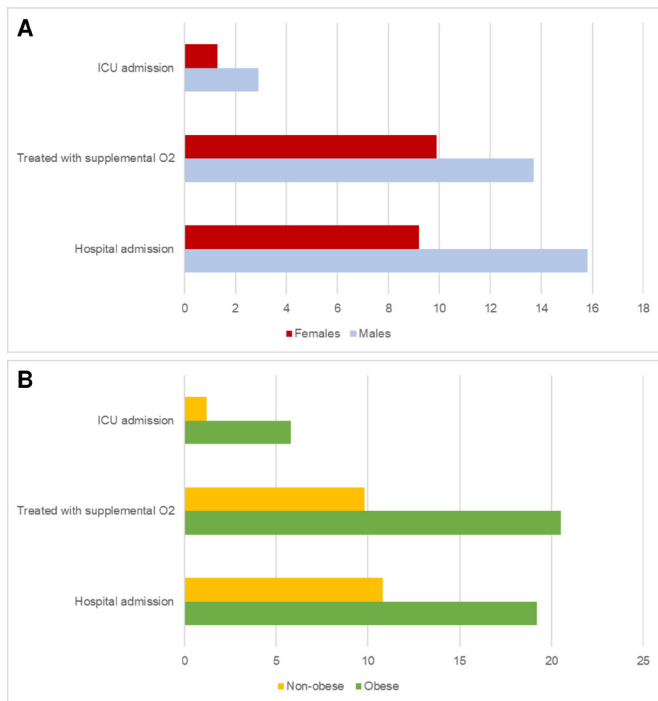


Figure 1 Percentage of patients admitted to the hospital, treated with supplemental oxygen and admitted to ICU according to gender (A) or obesity (B). ICU, intensive care unit.

(OR: 1.786, 95% CI 1.272 to 2.506, $p=0.001$), intensity of treatment setting (OR: 1.727, 95% CI 1.344 to 2.218, $p<0.001$), treatment with antibiotics (OR: 1.440, 95% CI 1.033 to 2.008, $p=0.031$), LMWH (OR: 2.180, 95% CI 1.472 to 3.229, $p<0.0001$), supplemental oxygen (OR: 2.531, 95% CI 1.514 to 4.229, $p<0.0001$), high blood pressure (OR: 3.061, 95% CI 1.964 to 4.770, $p<0.0001$) and obesity (OR: 41.192, 95% CI 18.003 to 98.879, $p<0.0001$) significantly predicted the fulfilment of FM criteria. In a multivariate model including all the aforementioned variables, only male gender (OR: 9.951, 95% CI 6.025 to 16.435, $p<0.0001$) and obesity (OR: 82.823, 95% CI 32.192 to 213.084, $p<<0.0001$) predicted FM classification. Finally, as depicted in figure 1, male sex (figure 1A) and obesity (figure 1B) were associated with surrogate measures of COVID-19 severity, including higher hospital admission, treatment with supplemental oxygen and intensive care unit (ICU) admission.

Given the significant role of male gender as a predictor of FM, we compared the clinical characteristics of male versus female respondents (online supplementary table S3). Although women had more symptoms during acute COVID-19, the rate of hospital admission was significantly higher in male respondents (15.8% vs 9.2%, $p=0.001$). Moreover, BMI was higher in men (26.3 ± 4.3 kg/m² vs 24.9 ± 4.9 kg/m²) as was the percentage of overweight individuals (36.0% vs 23.1%, $p=0.002$). Similarly, when comparing non-obese versus obese individuals, the latter showed a higher rate of hospital admission (19.2% vs 10.8%, $p=0.016$) and treatment with supplemental

oxygen (20.5% vs 9.8%, $p=0.002$) and admission to ICU (5.8% vs 1.2%, $p=0.003$).

DISCUSSION

In this study, we demonstrated that self-reported clinical features of FM are common after symptomatic COVID-19, with an estimated prevalence of ~31%. Notably, this figure is similar to that found in other chronic painful disorders¹⁹ and comparable to the 30% recently reported for PACS after a similar follow-up.²⁰

Globally, respondents with FM exhibited features suggestive of a more serious form of COVID-19, including a higher rate of hospitalisation and more frequent treatment with supplemental oxygen. Unfortunately, the study design did not allow accurate definition of the clinical severity of COVID-19,²¹ and thus, our evaluation relies solely on surrogate measures. However, when a multivariate model was built, obesity and male gender were identified as independent, strong predictors of being classified as FM. Notably, both male gender²² and obesity²³ have been consistently associated with a more severe clinical course in patients with COVID-19, including a significantly increased mortality rate.

Strikingly, we found a high percentage of men (43%) in respondents meeting criteria for FM. Subanalysis of our data revealed that male gender was associated with surrogate measures of COVID-19 severity, as suggested by a significantly higher rate of patients requiring hospital admission. Thus, the most intuitive explanation for the increased prevalence of FM in men is the overall tendency to a more aggressive disease course. However, other speculative mechanisms may contribute to this phenomenon. Although it is a common belief that FM is a female-predominant disorder, an elegant study by Wolfe *et al*²⁴ questioned this assumption, suggesting that gender specificity may be the consequence of several biases. Indeed, similarly to what we observed, they demonstrated that women represent ~59% of cases of FM when classification criteria are applied to individuals, as opposed to 'traditional', biased cohorts where they account for nearly 90% of patients. Similar figures have been reported in other studies,²⁵ including a web-based survey.¹⁴

The second, perhaps strongest, predictor of FM in our cohort was obesity. The relationship between obesity and FM is mutual and bidirectional; in a recent systematic review and meta-analysis,²⁶ our group demonstrated that BMI can influence nearly all domains of the syndrome.

Taken together, our data suggest a speculative mechanism in which obesity and male gender synergistically affect the severity of COVID-19 that, in turn, may rebound on the risk of developing post-COVID-19 FM syndrome and determine its severity. Interestingly, individuals who got COVID-19 during the first pandemic wave—when a prejudicial mix of hospital overloading and extremely limited knowledge of the disease affected the management of the disease—showed a tendency towards an increase in FM prevalence when compared with those

who got COVID-19 during second wave, further emphasising a possible association with the disease severity and proper management.

Despite their usefulness in healthcare research, online surveys are affected by well-known intrinsic limitations.^{27,28} First, the respondents are not selected through probability sampling, and this may impair the generalisability of the findings; in addition, information about non-respondents is not available. Thus, a self-selection bias may arise because some individuals are more likely than others to complete online surveys. Several factors have been associated with non-response in health surveys,²⁹ including male gender, younger age, lower socioeconomic status, and poorer health and health behaviours. Thus, the lower rate of male respondents in our survey is not surprising and reflects a well-known gender bias in survey-based research, with women being more prone to participate in online surveys.^{30,31}

In an attempt to ascertain the presence of self-selection bias, we used a classical approach based on comparing study results with auxiliary information available from official government data. In Italy, the median age of confirmed COVID-19 cases is 47 years,¹⁸ and analysing official Italian Ministry of Health data collected from 24 February 2020 to 25 April 2021, we found that the mean daily percentage of hospitalised patients was 11.1% of all confirmed cases, while 1.4% were admitted to ICU. The figures observed in our sample, with 10.7% and 1.6% of patients, respectively, treated in a non-critical hospital setting or in an ICU, are therefore comparable to the general COVID-19 population. Moreover, also additional sociodemographic characteristics are similar. For instance, in 2020, taking into account the age range 30–59 years, 63.2% of the Italian population was married and 31% was single. Correspondingly, the majority of participants in our sample were married (70.9%), while only 18.8% were single.¹⁷

The only major difference between our study sample and the COVID-19 population in Italy is the female predominance of survey respondents. Although the overall gender ratio of COVID-19 is thought to be ~1:1, official data from the Italian Ministry of Health demonstrate an uneven distribution of COVID-19 cases, with a female predominance in the age range of 30–59 years and, on the contrary, a male predominance in individuals >60 years of age¹⁸; female predominance is perhaps more evident in certain populations, such as healthcare professionals. Similarly, a female predominance has been reported in other countries according to The Sex, Gender and COVID-19 Project, an online database of gender-disaggregated data on COVID-19.³² It is important to note that these data refer to the overall population of SARS-CoV-2-positive individuals and do not distinguish between mildly symptomatic (the vast majority in our sample) and fully asymptomatic individuals. Interestingly, available literature suggests that, taking into consideration only mild cases of COVID-19, women seem to be more represented than men.^{33–36}

However, to account for this potential source of bias, we applied gender-calculated poststratification weights to all analyses; no major differences emerged when compared with raw data.

In conclusion, clinical features of FM are common in patients who recovered from symptomatic COVID-19. Preliminary evidence from clinical and preclinical studies suggests that several disease-specific mechanisms may explain the pathophysiology of this musculoskeletal syndrome, including virus-induced injury to endothelium³⁷ or neuromuscular structures,³⁸ immunological derangement and smouldering inflammation. Regarding the latter, it is interesting to note that some of the proinflammatory cytokines involved in COVID-19 and PACS manifestations, such as interleukin (IL)-1 and IL-6,^{2,39} may contribute to the pathogenesis of FM.^{40,41} Unfortunately, our data do not provide a mechanistic support in understanding the pathophysiology of fibromyalgias in these patients and other, indirect and non-specific processes—for example, prolonged bed rest, deconditioning, post-traumatic stress disorder—may actually prevail. Moreover, the lack of a control group impairs the possibility to ascertain a possible contribution of the psychophysical distress associated with lockdown measures and other pandemic-related constraints on the susceptibility to FM-like symptoms in patients without a known SARS-CoV-2 exposure.

In the light of the overwhelming numbers of the SARS-CoV-2 pandemic, it is reasonable to forecast that rheumatologists will face up with a sharp rise of cases of a new entity that we defined ‘FibroCOVID’ to underline potential peculiarities and differences, such as the male involvement. From the rheumatology perspective, some open questions need to be addressed in the near future. First, how are patients, in general practice and other specialty settings (eg, infectious disease clinics), who deserve referral to the rheumatologist after COVID-19 identified for suspected FM? Easy-to-use, inexpensive and quick instruments, such as the Fibromyalgia Rapid Screening Tool⁴² questionnaire or the London Fibromyalgia Epidemiology Study Screening Questionnaire,⁴³ may be the answer, but they need adequate validation in this new population of patients. Second, what is the optimal treatment strategy for FibroCOVID? Although no definitive protocols are still available for FM treatment, it is possible to hypothesise that a traditional approach including graded exercise, cognitive behavioural therapy and pain modulators may still help patients. On the other hand, given the suspected viral trigger, other treatments (eg, immune-modulating agents) or SARS-CoV-2 vaccines may provide specific benefits. Finally, what is the clinical course of post-COVID-19 musculoskeletal symptoms? Prospective studies, including comparative analysis with primary FM cohorts, will shed light on this topic.

Author affiliations

¹Medicine and Rheumatology Unit, IRCCS Istituto Ortopedico Rizzoli, Bologna, Italy

²Department of Biomedical and Neuromotor Sciences (DIBINEM), Alma Mater Studiorum University of Bologna, Bologna, Italy

³Department of Experimental, Diagnostic and Specialty Medicine, IRCCS Azienda Ospedaliero-Universitaria di Bologna Policlinico S Orsola-Malpighi, Bologna, Italy

⁴Clinical Trial Centre, IRCCS Istituto Ortopedico Rizzoli, Bologna, Italy

⁵Risk Management Unit, ASL 5 La Spezia, La Spezia, Italy

⁶Department of Experimental and Clinical Medicine, Magna Graecia University of Catanzaro, Catanzaro, Italy

⁷Department of Biotechnological and Applied Clinical Sciences, Università degli Studi dell'Aquila, L'Aquila, Italy

⁸Diagnostic and Interventional Radiology, IRCCS Istituto Ortopedico Rizzoli, Bologna, Italy

⁹Department of Morphology, Surgery and Experimental Medicine, University of Ferrara, Ferrara, Italy

¹⁰Biomedical Science and Technology Unit, IRCCS Istituto Ortopedico Rizzoli, Bologna, Italy

¹¹Department of Medical and Surgical Sciences, Alma Mater Studiorum University of Bologna, Bologna, Italy

¹²Department of Oncological and Degenerative Spine Surgery, IRCCS Istituto Ortopedico Rizzoli, Bologna, Italy

¹³Academic Rheumatology Centre, MFRU and Dipartimento Scienze Cliniche e Biologiche, Università degli Studi di Torino, Torino, Italy

¹⁴Unit of Allergology, Immunology, Rheumatology, Department of Medicine, Università Campus Bio-Medico di Roma, Roma, Italy

¹⁵1st Orthopaedic and Traumatologic Clinic, IRCCS Istituto Ortopedico Rizzoli, Bologna, Italy

¹⁶Scientific Direction, IRCCS Istituto Ortopedico Rizzoli, Bologna, Italy

¹⁷Department of Biomedical and Neuromotor Sciences, University of Bologna, Bologna, Italy

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ORCID iDs

Francesco Ursini <http://orcid.org/0000-0001-8194-8642>
 Riccardo Meliconi <http://orcid.org/0000-0001-5216-5219>

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