COVID-19 and its continuing burden after 12 months: a longitudinal observational prospective multicenter trial

Supplementary Material

CovILD study team

# Supplementary Methods

## Data transformation, variable stratification, descriptive statistic

The study variables were transformed, analyzed and visualized with R version 4.0.5 with *tidyverse* (1,2) and *cowplot* (3).

Participant age during acute COVID-19 was stratified with the 50 and 65 year cutoffs. Participants were stratified according to the severity of acuet COVID-19 as ambulatory (outpatients), moderate (hospitalized at COVID-19 ward, no mechanical ventilation) and severe (mechanical ventilation and/or intensive care unit stay). Lung function testing (LFT) parameters were stratified by 80% predicted value (FEV1: forced expiratory volume in 1 second; FVC: forced vital capacity; DLCO: diffusion lung capacity for carbon monoxide; TLC: total lung capacity) or 70% predicted value cutoffs (FEV1:FVC: FEV1 to FVC ratio). Abnormal LFT was diagnosed when at least one of FEV1, FVC, DLCO, RV, TLC or FEV1:FVC was reduced. Chest computed tomography was assessed and rated with the CT severity score as described (4,5); moderate-to-severe CT abnormalities were defined by CT severity score > 5. Anemia was defined as hemoglobin < 140 g/dL or < 120 g/dL for men and women, respectively. Elevated ferritin (FT) was defined as 300 µg/L or 150 µg/L for men and women, respectively. Reduced transferrin saturation (TSAT) was defined as < 20% or < 15% for men and women, respectively. Elevated N-terminal pro - brain natriuretic peptide (NT-proBNP) was defined as > 150 pg/mL. Elevated D-dimer, C-reactive protein (CRP), procalcitonin (PCT) and interleukin 6 (IL6) were defined with the 500 µg/L, 0.5 mg/L, 0.15 µg/L and 7 pg/mL cutoffs, respectively. Anti-S1/S2 SARS-CoV-2 immunoglubulin gamma (anti-S1/S2 IgG) was stratified by quartiles (cutoffs: 54.35, 109, 168 arbitrary units) (4). For clustering, the sub-scores of the EQ-5D-5L questionnaire (6) addressing impairment of usual activities, mobility and self-care as well as pain/discomfort, depression/anxiety were stratified with the cutoff of 1 (1: no impairment or absence, > 1:impairment or presence). Visual analogue scale (VAS) of the EQ-5D-5L tool describing the self-perceived general health was binarized with the cutoff of 73.2 as published before for the general elderly German population (7). Elevated stress levels were defined by the median split of the 4-item PSS score (> 5: elevated stress). Reference values of the six-minute walking distance (SMWD) (8) were calculated with the participants sex, age, weight and height as described previously (9). Low SMWD was defined as a value below the patient’s specific reference value. The full list of the study variables and their stratification scheme is presented in **Supplementary Tables 1** and **S2**.

## Hypothesis testing and correlation

To compare differences in distribution of categorical features, test with Cramer V effect size statistic was applied. Since multiple numeric variables were strongly non-normally distributed as identified by Shapiro-Wilk test, Mann-Whitney U test with Wilcoxon r effect size statistic or Kruskal-Wallis test with effect size statistic was applied to assess differences between two or more groups, respectively. Association of the readouts of clinical, cardiopulmonary, mobility and quality of life deficits at the one-year follow-up visit were determined by Kendall’s B correlation. Binary variables were recoded as 1 and 2 for absent and present, respectively. P values were corrected for multiple comparisons with Benjamini-Hochberg method (10). R packages *rstatix*, *rcompanion* and *Hmisc* and the in-house developed package *ExDA* (<https://github.com/PiotrTymoszuk/ExDA>) were used for explorative data analysis, statistical hypothesis testing and correlation analysis.

## Modeling of risk and paramater value in time

To model recovery kinetics for categorical variables, second-order mixed-effect logistic (categorical features) modeling was applied (packages: *lme4* and *lmerTest*) (11–13). Each model followed the general formula:

where indicates the random effect of the individual and and indicate the first- and second-order time effect terms. The first-order term estimate was interpreted as a measure of the recovery speed and the second-order term estimate was used to assess the plateau/rebound effect. Significance of the accuracy gain of the full second-order model compared with the nested null model was determined by likelihood ratio test (LRT) versus the nested first-order and null models, respectively. Likelihood ratio statistic was used as an effect size measure.

To assess the longitudinal changes of a numeric dependent variable in time, Friedman test with Kendall’s W effect was used (packages: *rstatix* and *ExDA*: <https://github.com/PiotrTymoszuk/ExDA>). Results of the kinetic modeling and Friedman tests were adjusted for multiple comparisons with Benjamini-Hochberg method (10).

## Clustering analysis

To identify patterns of COVID-19 recovery defined by 19 binary symptom, cardiopulmonary and psychosocial variables recorded at the 1-year follow-up visit (**Supplementary Tables S1** and **S2**) the subset of the study participants with the complete variable record was clustered using the PAM (partition around medoids) algorithm and the simple matching distance statistic (14,15). The data pre-processing included conversion of binary features to the numeric format (absent: 1, present: 1). The choice of the clustering procedure was motivated by the analysis of the clustering variance (ratio of the total between-cluster to total sum of squares) and clustering structure stability in 10-fold cross-validation (metric: rate of correct cluster assignment, cluster assignment predicted by k = 5 nearest neighbors label propagation algorithm) (16,17) for several clustering algorithms as presented in **Supplementary Figure S4A**. The optimal number of clusters was determined by the bend of the total within-cluster sum of squares curve (**Supplementary Figure S4B**). The clustering analysis was accomplished with packages *cluster*, *dbscan* and the in-house developed package *clustTools* (<https://github.com/PiotrTymoszuk/clustTools>).

## Univariable modeling

Association of 34 candidate co-variates recorded during acute COVID-19 or at the 60-day follow-up (**Supplementary Table S1**) with the risk of persistent COVID-19 symptoms, LFT or CT abnormalities or diastolic dysfunction at the 1-year follow-up was investigated with a series of univariable logistic models (**Supplementary Table S6**). Model estimate (odds ratio, OR) significance was assessed by Wald Z test and p values were corrected for multiple comparisons with Benjamini-Hochberg method (10). Model estimate extraction and quality control was accomplished with the in-house-developed package *lmqc* (<https://github.com/PiotrTymoszuk/lmqc>).

## Multi-parameter modeling

The risk of persistent COVID-19 symptoms, LFT or CT abnormalities or diastolic dysfunction at the 1-year follow-up was modeled with multi-parameter logistic LASSO (least absolute shrinkage and selection operator) regression (18) and 34 independent variables recorded during acute COVID-19 and at the 60-day follow-up (**Supplementary Table S1**). The value of the shrinkage parameter were determined by 10-fold cross-validation (CV) repeated 100 times; the optimal value for the model with the lowest deviance was chosen (19). The model performance at predicting the 1-year event in the training data set and in 10-fold CV was investigated by receiver-operating characteristic (ROC) (20), Cohen’s (21) and R-squared statistics. Model assumption control was accomplished by a visual control of model residuals plots (residuals versus fitted and quantile-quantile plots). The multi-parameter modeling, model evaluation and model feature visualization tasks was accomplished with *glmnet* (19), *caret* (20), *plotroc* packages (22) and the in-house development package *caretExtra* (<https://github.com/PiotrTymoszuk/caretExtra>).

## Source code availability

The raw study data will be made available upon request. The R analysis pipeline is available at <https://github.com/PiotrTymoszuk/CovILD-Plus>.

# Supplementary Tables

**Supplementary** **Table S1:** Study variables.

| **R variable** | **Label** | **Collection time points** | **Independent modeling variable** | **Correlation variable** | **Clustering variable** |
| --- | --- | --- | --- | --- | --- |
| smwd | SMWD, m | 1-year FUP | no | no | no |
| smwd\_dref | SMWD vs ref., m | no | yes | no |
| smwd\_low | SMWD < ref., % | no | no | yes |
| BRCS | Resilience (BRCS) | no | no | no |
| BRCS\_class | Resilience (BRCS) | no | no | no |
| EQ5DL\_p | General health (EQ5D5L VAS), % | no | yes | no |
| EQ5DL\_mobility | Mobility impairment (EQ5D5L) | no | yes | no |
| EQ5DL\_selfcare | Self-care impairment (EQ5D5L) | no | yes | no |
| EQ5DL\_activities | Activity impairment (EQ5D5L) | no | yes | no |
| EQ5DL\_pain | Pain/discomfort (EQ5D5L) | no | yes | no |
| EQ5DL\_anxiety | Anxiety/depression (EQ5D5L) | no | yes | no |
| EQ5DL\_low | Imp. general health (VAS < 73, EQ5D5L), % | no | no | yes |
| EQ5DL\_mobility\_bi | Imp. mobility (EQ5D5L), % | no | no | yes |
| EQ5DL\_selfcare\_bi | Imp. self-care (EQ5D5L), % | no | no | yes |
| EQ5DL\_activities\_bi | Imp. usual activity (EQ5D5L), % | no | no | yes |
| EQ5DL\_pain\_bi | Pain/discomfort present (EQ5D5L), % | no | no | yes |
| EQ5DL\_anxiety\_bi | Anxiety/depression present (EQ5D5L), % | no | no | yes |
| Chalder\_FS | Fatigue (likert CFS) | no | yes | no |
| Chalder\_FS\_bimodal | Fatigue (bimodal CFS ≥ 4), % | no | no | yes |
| SSD12 | Somatic symptom disorder (SSD-12) | no | no | no |
| Stress | Stress (PSS) | no | yes | no |
| Stress\_hi | Elevated stress (PSS > 5), % | no | no | yes |
| SES | Self-efficacy (GSES) | no | no | no |
| KW\_IPQ | Illness perception (IPQ) | no | no | no |
| SOCL9 | Sense of coherence loss (SOCL-9) | no | no | no |
| FVC\_p | FVC, % | 60- till 1 year FUP | no | no | no |
| FEV1\_p | FEV1, % | no | no | no |
| FEV1\_FVC\_p | FEV1:FVC, % | no | no | no |
| DLCO\_p | DLCO, % | no | yes | no |
| TLC\_p | TLC, % | no | no | no |
| FVC\_red | reduced FVC, % | no | no | no |
| FEV1\_red | reduced FEV1, % | no | no | no |
| FEV1\_FVC\_red | reduced FEV1:FVC, % | no | no | no |
| TLC\_red | reduced TLC, % | no | no | no |
| DLCO\_red | reduced DLCO, % | no | no | no |
| lufo\_red | LFT abnormality, % | no | yes | yes |
| Hb | Hb, g/dL | no | no | no |
| anemia | Anemia, % | yes | no | no |
| ferritin | FT, µg/L | no | no | no |
| FT\_elv | elevated FT, % | yes | no | no |
| TSAT | TF-Sat, % | no | no | no |
| TSAT\_red | reduced TF-Sat, % | no | no | no |
| sTFR | sTFR, mg/L | no | no | no |
| Hepcidin | Hepcidin, ng/mL | no | no | no |
| NTproBNP | NT-proBNP, pg/mL | no | no | no |
| NtproBNP\_elv | elevated NT-proBNP, % | yes | no | no |
| DDimer | D-dimer, µg/L | no | no | no |
| Ddimer\_elv | elevated D-dimer, % | yes | no | no |
| CRP | CRP, mg/L | no | no | no |
| CRP\_elv | elevated CRP, % | yes | no | no |
| PCT | PCT, µg/L | no | no | no |
| PCT\_elv | elevated PCT, % | no | no | no |
| IL6 | IL6, pg/mL | no | no | no |
| IL6\_elv | elevated IL6, % | yes | no | no |
| HbA1c | HbA1c, % | no | no | no |
| HbA1c\_elv | elevated HbA1c, % | no | no | no |
| EF\_red | reduced LVEF, % | no | no | no |
| diastolic\_dysf | Diastolic dysfunction, % | no | yes | yes |
| mmrc | mMRC | no | yes | no |
| ECOG | ECOG | no | no | no |
| ct\_severity\_score | CT severity score | no | yes | no |
| ct\_severity\_any | CT abnormality, % | no | no | yes |
| ct\_severity\_5 | CT severity score > 5, % | no | no | no |
| ab\_quant | anti-S1/S2 IgG | 60-day FUP | yes | no | no |
| sex | Sex | acute COVID-19 | yes | no | no |
| age | Age, years | yes | no | no |
| cat\_WHO | COVID-19 severity | yes | no | no |
| weight\_class | Weight class | yes | no | no |
| smoking | Smoking | yes | no | no |
| no\_comorb | # comorbidities | yes | no | no |
| comorb\_present | Comorbidity present, % | no | no | no |
| cardiovascular\_comorb | CVD, % | yes | no | no |
| hypertension\_comorb | Hypertension, % | yes | no | no |
| pulmonary\_comorb | Pulmonary disease, % | yes | no | no |
| copd\_comorb | COPD, % | yes | no | no |
| asthma\_comorb | Asthma, % | yes | no | no |
| intenst\_lung\_comorb | ILD, % | no | no | no |
| endometabolic\_comorb | Metabolic disease, % | yes | no | no |
| hyperchol\_comorb | Hypercholesterolemia, % | yes | no | no |
| diabetes\_comorb | Diabetes, % | yes | no | no |
| ckd\_comorb | CKD, % | yes | no | no |
| gastro\_comorb | GID, % | yes | no | no |
| cldis\_comorb | CLD, % | no | no | no |
| malingancy\_comorb | Malignancy, % | yes | no | no |
| immdef\_comorb | Immune deficiency, % | yes | no | no |
| sympt\_present | Symptoms present, % | acute COVID-19 till 1-year FUP | no | no | yes |
| sympt\_number | # symptoms | yes | yes | no |
| sleep\_sympt | Sleep problems, % | yes | yes | yes |
| dyspnoe\_sympt | Dyspnea, % | yes | no | yes |
| cough\_sympt | Cough, % | yes | yes | yes |
| fever\_sympt | Fever, % | no | no | no |
| night\_sweat\_sympt | Night sweat, % | yes | yes | yes |
| gastro\_sympt | Gastrointestinal, % | yes | no | no |
| anosmia\_sympt | Hypo/anosmia, % | yes | yes | yes |
| fatigue\_sympt | Reduced performance, % | yes | yes | yes |
| pain\_sympt | Pain, % | yes | no | no |
| hair\_loss\_sympt | Hair loss, % | no | no | no |
| derma\_sympt | Dermatological, % | no | no | no |
| rehabilitation | Rehabilitation, % |  | no | no | no |

**Supplementary** **Table S2:** Physical performance, fatigue, quality of life, psychosocial and mental health assessment battery.

| **Evaluated parameter** | **Tool** | **Stratification scheme** |
| --- | --- | --- |
| Dyspnea | Modified Medical British Research Council (mMRC) | Dyspnea: > 0 |
| Physical performance | Eastern Cooperative Oncology Group (ECOG) | Reduced performance: > 0 |
| Mobility | Six-minute walking distance (SMWD) |  |
| Six-minute walking distance versus reference (SMWD vs ref.) | Impaired mobility: < 0 |
| Fatigue | Likert 11-item Chalder Fatigue Score (CFS) |  |
| Bimodal 11-item Chalder Fatigue Score (CFS) | Increased fatigue: ≥ 4 |
| Self-perceived general health | European quality of life 5 dimensions, visual analogue scale (EQ5D5L VAS) | Impaired general health: < 73 |
| Mobility impairment | European quality of life 5 dimensions, mobility sub-score (EQ5D5L mobility) | Impaired mobility: > 1 |
| Self-care impairment | European quality of life 5 dimensions, self-care sub-score (EQ5D5L self-care) | Impaired self care: > 1 |
| Usual activity impairment | European quality of life 5 dimensions, activity sub-score (EQ5D5L activity) | Impaired activity: > 1 |
| Pain/discomfort | European quality of life 5 dimensions, pain/discomfort sub-score (EQ5D5L pain/discomfort) | Pain/discomfort present: > 1 |
| Anxiety/depression | European quality of life 5 dimensions, anxiety/depression sub-score (EQ5D5L anxiety/depression) | Anxiety/depression present: > 1 |
| Resilience | Brief resilient coping scale (BRCS) |  |
| Somatic symptom disorder | Somatic symptom disorder – B criteria scale, 12 items (SSD-12) |  |
| Psychosocial stress | 4-item perceived stress score (PSS) | Elevated stress: > 5 |

**Supplementary** **Table S3:** Demographic and clinical characteristics of the COVID-19 recovery clusters.

| **Variable** | **Cluster #1** | **Cluster #2** | **Cluster #3** | **Significancea** | **Effect sizea** |
| --- | --- | --- | --- | --- | --- |
| N number | 36 | 33 | 18 |  |  |
| Sex | female: 42% (n = 15) male: 58% (n = 21) | female: 18% (n = 6) male: 82% (n = 27) | female: 72% (n = 13) male: 28% (n = 5) | p = 0.0025 | V = 0.41 |
| Age, years | median: 55 [IQR: 53 - 61] range: 19 - 77 | median: 63 [IQR: 53 - 71] range: 36 - 80 | median: 50 [IQR: 42 - 58] range: 32 - 87 | ns (p = 0.059) | η² = 0.061 |
| Weight class | normal: 22% (n = 8) overweight: 56% (n = 20) obesity: 22% (n = 8) | normal: 36% (n = 12) overweight: 39% (n = 13) obesity: 24% (n = 8) | normal: 44% (n = 8) overweight: 44% (n = 8) obesity: 11% (n = 2) | ns (p = 0.54) | V = 0.15 |
| Smoking | never: 64% (n = 23) ex: 36% (n = 13) active: 0% (n = 0) | never: 61% (n = 20) ex: 36% (n = 12) active: 3% (n = 1) | never: 72% (n = 13) ex: 17% (n = 3) active: 11% (n = 2) | ns (p = 0.32) | V = 0.19 |
| Comorbidity present | 72% (n = 26) | 82% (n = 27) | 67% (n = 12) | ns (p = 0.58) | V = 0.14 |
| Metabolic disease | 39% (n = 14) | 58% (n = 19) | 28% (n = 5) | ns (p = 0.18) | V = 0.23 |
| Diabetes | 14% (n = 5) | 15% (n = 5) | 17% (n = 3) | ns (p = 0.96) | V = 0.029 |
| Hypercholesterolemia | 31% (n = 11) | 30% (n = 10) | 0% (n = 0) | ns (p = 0.058) | V = 0.29 |
| CVDb | 36% (n = 13) | 61% (n = 20) | 22% (n = 4) | p = 0.042 | V = 0.3 |
| Pulmonary disease | 11% (n = 4) | 24% (n = 8) | 17% (n = 3) | ns (p = 0.51) | V = 0.15 |
| Malignancy | 8.3% (n = 3) | 15% (n = 5) | 0% (n = 0) | ns (p = 0.34) | V = 0.19 |
| Immune deficiency | 2.8% (n = 1) | 3% (n = 1) | 11% (n = 2) | ns (p = 0.49) | V = 0.16 |
| CKDc | 5.6% (n = 2) | 12% (n = 4) | 5.6% (n = 1) | ns (p = 0.68) | V = 0.12 |
| GIDd | 11% (n = 4) | 15% (n = 5) | 11% (n = 2) | ns (p = 0.9) | V = 0.059 |
| Rehabilitation | no: 86% (n = 31) inpatient: 11% (n = 4) outpatient: 2.8% (n = 1) | no: 58% (n = 19) inpatient: 30% (n = 10) outpatient: 12% (n = 4) | no: 44% (n = 8) inpatient: 50% (n = 9) outpatient: 5.6% (n = 1) | p = 0.025 | V = 0.28 |
| COVID-19 severity | Ambulatory: 33% (n = 12) Moderate: 50% (n = 18) Severe: 17% (n = 6) | Ambulatory: 12% (n = 4) Moderate: 67% (n = 22) Severe: 21% (n = 7) | Ambulatory: 33% (n = 6) Moderate: 17% (n = 3) Severe: 50% (n = 9) | p = 0.012 | V = 0.3 |
| aCategorical variables: χ² test with Cramer V effect size statistic; numeric variables: Kruskal-Vallis test with η² effect size statistic. P values corrected for multiple testing with Benjamini-Hochberg method. | | | | | |
| bCVD: Cardiovascular disease. | | | | | |
| cCKD: Chronic kidney disease | | | | | |
| dGID: Gastrointestinal disease | | | | | |

**Supplementary** **Table S4:** Symptoms and cardiopulmonary abnormalities at the 1-year follow-up in the COVID-19 recovery clusters.

| **Variable** | **Cluster #1** | **Cluster #2** | **Cluster #3** | **Significancea** | **Effect sizea** |
| --- | --- | --- | --- | --- | --- |
| N number | 36 | 33 | 18 |  |  |
| Symptoms present | 28% (n = 10) | 85% (n = 28) | 100% (n = 18) | p < 0.001 | V = 0.65 |
| # symptoms | median: 0 [IQR: 0 - 1] range: 0 - 4 | median: 1 [IQR: 1 - 2] range: 0 - 5 | median: 4 [IQR: 3.2 - 4.8] range: 1 - 9 | p < 0.001 | η² = 0.49 |
| Sleep problems | 11% (n = 4) | 24% (n = 8) | 67% (n = 12) | p < 0.001 | V = 0.47 |
| Dyspnea | 5.6% (n = 2) | 18% (n = 6) | 72% (n = 13) | p < 0.001 | V = 0.59 |
| Cough | 5.6% (n = 2) | 9.1% (n = 3) | 56% (n = 10) | p < 0.001 | V = 0.52 |
| Night sweat | 11% (n = 4) | 21% (n = 7) | 28% (n = 5) | ns (p = 0.43) | V = 0.17 |
| Gastrointestinal | 2.8% (n = 1) | 0% (n = 0) | 22% (n = 4) | p = 0.0091 | V = 0.37 |
| Hypo/anosmia | 14% (n = 5) | 15% (n = 5) | 11% (n = 2) | ns (p = 0.95) | V = 0.043 |
| Reduced performance | 5.6% (n = 2) | 48% (n = 16) | 83% (n = 15) | p < 0.001 | V = 0.62 |
| Hair loss | 0% (n = 0) | 3% (n = 1) | 33% (n = 6) | p < 0.001 | V = 0.48 |
| Dermatological | 2.8% (n = 1) | 21% (n = 7) | 28% (n = 5) | ns (p = 0.052) | V = 0.29 |
| LFT abnormalityb | 25% (n = 9) | 42% (n = 14) | 44% (n = 8) | ns (p = 0.37) | V = 0.19 |
| CT abnormality (CT score ≥ 1)c | 25% (n = 9) | 85% (n = 28) | 44% (n = 8) | p < 0.001 | V = 0.54 |
| Diastolic dysfunction | 58% (n = 21) | 70% (n = 23) | 56% (n = 10) | ns (p = 0.65) | V = 0.12 |
| aCategorical variables: χ² test with Cramer V effect size statistic; numeric variables: Kruskal-Vallis test with η² effect size statistic. P values corrected for multiple testing with Benjamini-Hochberg method. | | | | | |
| bLFT abnormality: abnormality in lung function testing, > 80% predicted value (FEV1: forced expiratory volume in 1 second; FVC: forced vital capacity; DLCO: diffusion lung capacity for carbon monoxide; TLC: total lung capacity) or > 70% predicted value cutoffs (FEV1:FVC: FEV1 to FVC ratio.) | | | | | |
| cCT abnormality: any abnormality in chest computed tomography, CT severity score ≥ 1. | | | | | |

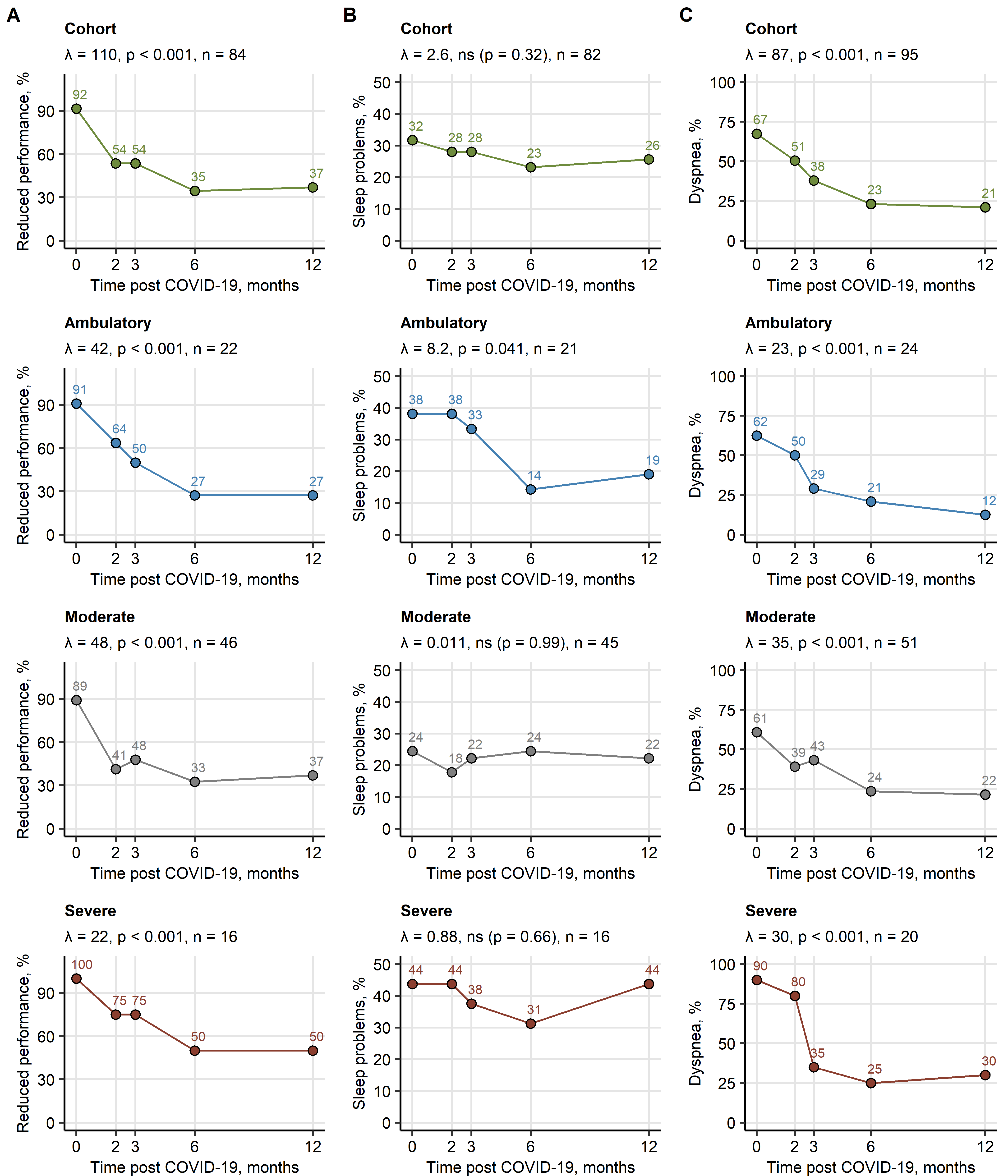
**Supplementary** **Table S5:** Mobility, physical performance and psychosocial rating at the 1-year follow-up in the COVID-19 recovery clusters.

| **Variable** | **Cluster #1** | **Cluster #2** | **Cluster #3** | **Significancea** | **Effect sizea** |
| --- | --- | --- | --- | --- | --- |
| N number | 36 | 33 | 18 |  |  |
| SMWD, mb | median: 580 [IQR: 530 - 640] range: 460 - 740 | median: 520 [IQR: 450 - 580] range: 270 - 760 | median: 530 [IQR: 420 - 620] range: 310 - 710 | p = 0.025 | η² = 0.085 |
| SMWD vs ref., mc | median: 38 [IQR: 5.1 - 62] range: -220 - 130 | median: -44 [IQR: -79 - 19] range: -220 - 120 | median: -72 [IQR: -100 - -9.3] range: -230 - 140 | p = 0.0022 | η² = 0.15 |
| SMWD < ref.d | 19% (n = 7) | 73% (n = 24) | 78% (n = 14) | p < 0.001 | V = 0.54 |
| Fatigue score (likert CFS)e | median: 11 [IQR: 11 - 12] range: 1 - 24 | median: 13 [IQR: 10 - 15] range: 1 - 25 | median: 20 [IQR: 16 - 26] range: 0 - 32 | p < 0.001 | η² = 0.24 |
| Fatigue (bimodal CFS ≥ 4)e | 22% (n = 8) | 36% (n = 12) | 72% (n = 13) | p = 0.0056 | V = 0.38 |
| General health score (EQ5D5L VAS)f | median: 90 [IQR: 80 - 95] range: 70 - 100 | median: 80 [IQR: 80 - 90] range: 60 - 100 | median: 70 [IQR: 61 - 75] range: 40 - 90 | p < 0.001 | η² = 0.26 |
| Imp. general health (VAS < 73, EQ5D5L)f | 8.3% (n = 3) | 9.1% (n = 3) | 61% (n = 11) | p < 0.001 | V = 0.54 |
| Mobility impairment score (EQ5D5L) | median: 1 [IQR: 1 - 1] range: 1 - 2 | median: 1 [IQR: 1 - 1] range: 1 - 2 | median: 1 [IQR: 1 - 2] range: 1 - 3 | p = 0.02 | η² = 0.093 |
| Imp. mobility (score > 1, EQ5D5L)g | 2.8% (n = 1) | 15% (n = 5) | 33% (n = 6) | p = 0.023 | V = 0.33 |
| Self-care impairment score (EQ5D5L) | median: 1 [IQR: 1 - 1] range: 1 - 1 | median: 1 [IQR: 1 - 1] range: 1 - 1 | median: 1 [IQR: 1 - 1] range: 1 - 2 | p = 0.0087 | η² = 0.12 |
| Imp. self-care (score > 1, EQ5D5L)g | 0% (n = 0) | 0% (n = 0) | 17% (n = 3) | p = 0.0084 | V = 0.37 |
| Activity impairment score (EQ5D5L) | median: 1 [IQR: 1 - 1] range: 1 - 1 | median: 1 [IQR: 1 - 1] range: 1 - 2 | median: 2 [IQR: 2 - 3] range: 1 - 3 | p < 0.001 | η² = 0.73 |
| Imp. usual activity (score > 1, EQ5D5L)g | 0% (n = 0) | 3% (n = 1) | 83% (n = 15) | p < 0.001 | V = 0.86 |
| Pain/discomfort score (EQ5D5L) | median: 1 [IQR: 1 - 1] range: 1 - 2 | median: 1 [IQR: 1 - 1] range: 1 - 3 | median: 2.5 [IQR: 2 - 3] range: 1 - 4 | p < 0.001 | η² = 0.39 |
| Pain/discomfort (score > 1, EQ5D5L)g | 22% (n = 8) | 21% (n = 7) | 89% (n = 16) | p < 0.001 | V = 0.57 |
| Anxiety/depression score (EQ5D5L) | median: 1 [IQR: 1 - 1] range: 1 - 2 | median: 1 [IQR: 1 - 1] range: 1 - 2 | median: 2 [IQR: 2 - 3.8] range: 1 - 5 | p < 0.001 | η² = 0.5 |
| Anxiety/depression (score > 1, EQ5D5L)g | 5.6% (n = 2) | 15% (n = 5) | 83% (n = 15) | p < 0.001 | V = 0.69 |
| Stress score (PSS)h | median: 5 [IQR: 1 - 8] range: 0 - 10 | median: 5 [IQR: 3 - 8] range: 0 - 11 | median: 9 [IQR: 8.2 - 10] range: 5 - 11 | p < 0.001 | η² = 0.27 |
| Elevated stress (PSS > 5)h | 39% (n = 14) | 42% (n = 14) | 94% (n = 17) | p < 0.001 | V = 0.44 |
| Somatic symptom disorder score (SSD-12)i | median: 4 [IQR: 1 - 7.5] range: 0 - 24 | median: 5 [IQR: 2 - 10] range: 0 - 26 | median: 20 [IQR: 12 - 24] range: 7 - 30 | p < 0.001 | η² = 0.29 |
| Resilience score (BRCS)j | median: 16 [IQR: 12 - 19] range: 4 - 20 | median: 16 [IQR: 12 - 18] range: 4 - 20 | median: 16 [IQR: 14 - 18] range: 9 - 19 | ns (p = 0.96) | η² = -0.022 |
| Resilience (BRCS)k | low: 35% (n = 12) medium: 24% (n = 8) high: 41% (n = 14) n = 34 | low: 31% (n = 10) medium: 19% (n = 6) high: 50% (n = 16) n = 32 | low: 17% (n = 3) medium: 39% (n = 7) high: 44% (n = 8) n = 18 | ns (p = 0.6) | V = 0.15 |
| aCategorical variables: χ² test with Cramer V effect size statistic; numeric variables: Kruskal-Vallis test with η² effect size statistic. P values corrected for multiple testing with Benjamini-Hochberg method. | | | | | |
| bSMWD: six-minute walking distance, meters. | | | | | |
| cSMWD vs ref.: difference between the reference and observed SMWD value, meters. | | | | | |
| dSMWD < ref.: SMWD below the reference value. | | | | | |
| eCFS: 11-item Chalder fatigue score, incr.: increased. | | | | | |
| fVAS: visual analogue scale, imp.: impaired. | | | | | |
| gEQ5D5L subscore > 1, imp.: impaired. | | | | | |
| hPSS: 4-item perceived stress scale. | | | | | |
| iSSD-12: 12-item somatic syndrome disorder – B criteria scale. | | | | | |
| jBRCS: brief resilient coping score. | | | | | |
| klow: 4 - 13 points, medium: 14 - 16 points, high: 17 - 21 points in the BRCS scale. | | | | | |

**Supplementary** **Table S6:** Significant results of univariable modeling of the risk of symptom presence, lung function abnormalities, radiological chest abnormalities and diastolic dysfunction at the 1-year follow-up.

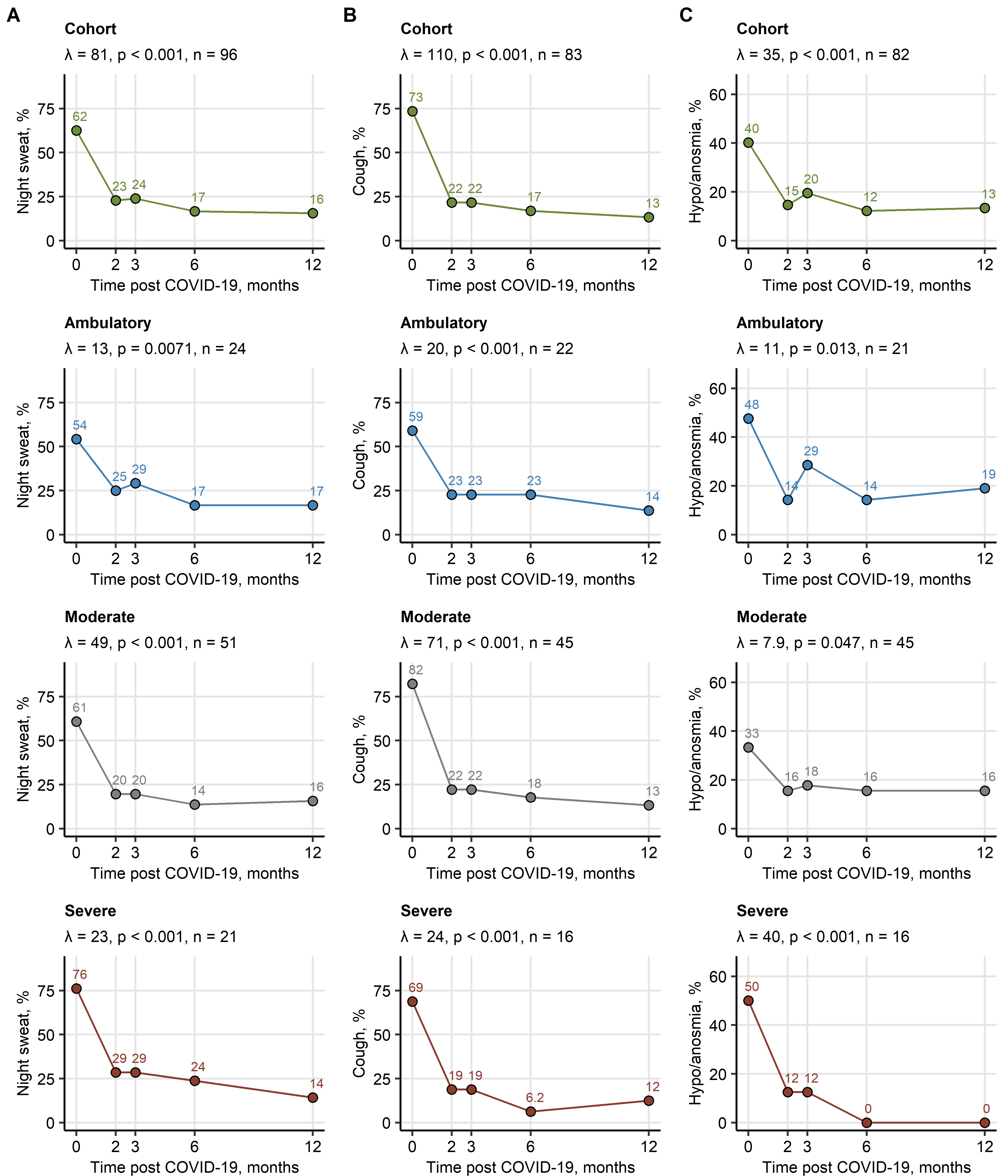
| **Response** | **Independent variable** | **Reference** | **Level** | **N level** | **N total** | **OR, 95% CI1** | **Significance2** |
| --- | --- | --- | --- | --- | --- | --- | --- |
| CT abnormality, 1 year FUP | COVID-19 severity, acute COVID-19 | Ambulatory | Moderate | 52 | 101 | 7.2 [2.1 - 33] | p = 0.027 |
| Ambulatory | Severe | 26 | 101 | 37 [8.3 - 220] | p < 0.001 |
| Sex, acute COVID-19 | female | male | 62 | 101 | 6.1 [2.6 - 15] | p = 0.0019 |
| Age, acute COVID-19 | up to 50 | 51 - 65 | 45 | 101 | 3 [1.1 - 8.5] | ns (p = 0.13) |
| up to 50 | over 65 | 29 | 101 | 4.5 [1.5 - 15] | p = 0.043 |
| # comorbidities, acute COVID-19 |  | per item | 101 | 101 | 1.5 [1.2 - 2] | p = 0.0045 |
| Diabetes, acute COVID-19 | no | yes | 16 | 101 | 8.7 [2.2 - 57] | p = 0.032 |
| Hypertension, acute COVID-19 | no | yes | 28 | 101 | 4.1 [1.6 - 11] | p = 0.027 |
| Anemia, 60-day FUP | no | yes | 25 | 101 | 5.5 [2 - 18] | p = 0.023 |
| anti-S1/S2 IgG, 60-day FUP | Q1 | Q2 | 26 | 93 | 22 [3.7 - 420] | p = 0.027 |
| Q1 | Q3 | 21 | 93 | 36 [5.8 - 710] | p = 0.02 |
| Q1 | Q4 | 23 | 93 | 230 [29 - 5600] | p < 0.001 |
| LFT abnormality, 1 year FUP | # comorbidities, acute COVID-19 |  | per item | 106 | 106 | 1.5 [1.2 - 2] | p = 0.0013 |
| elevated IL6, 60-day FUP | no | yes | 10 | 106 | 10 [2.4 - 71] | p = 0.015 |
| elevated D-dimer, 60-day FUP | no | yes | 38 | 106 | 3.2 [1.4 - 7.7] | p = 0.019 |
| Diastolic dysfunction, 1 year FUP | COVID-19 severity, acute COVID-19 | Ambulatory | Moderate | 54 | 107 | 5.2 [1.9 - 15] | p = 0.02 |
| Ambulatory | Severe | 26 | 107 | 13 [3.7 - 57] | p = 0.0046 |
| Age, acute COVID-19 | up to 50 | 51 - 65 | 46 | 107 | 11 [3.9 - 33] | p < 0.001 |
| up to 50 | over 65 | 29 | 107 | 11 [3.7 - 41] | p = 0.0026 |
| # comorbidities, acute COVID-19 |  | per item | 107 | 107 | 1.5 [1.2 - 1.9] | p = 0.014 |
| anti-S1/S2 IgG, 60-day FUP | Q1 | Q2 | 28 | 99 | 2.9 [0.98 - 9.2] | ns (p = 0.14) |
| Q1 | Q3 | 22 | 99 | 8.5 [2.4 - 37] | p = 0.024 |
| Q1 | Q4 | 23 | 99 | 6.8 [2 - 27] | p = 0.035 |
| Symptoms present, 1 year FUP | # symptoms, acute COVID-19 |  | per item | 104 | 104 | 1.4 [1.1 - 1.8] | p = 0.041 |
| Sleep problems, acute COVID-19 | no | yes | 40 | 104 | 4.2 [1.7 - 11] | p = 0.041 |
| elevated NT-proBNP, 60-day FUP | no | yes | 38 | 105 | 3.6 [1.4 - 9.9] | p = 0.043 |
| 1Odds ratio with 95% confidence intervals. | | | | | | | |
| 2Wald Z test. P values corrected for multiple testing with Benjamini-Hochberg method. | | | | | | | |

# Supplementary Figures



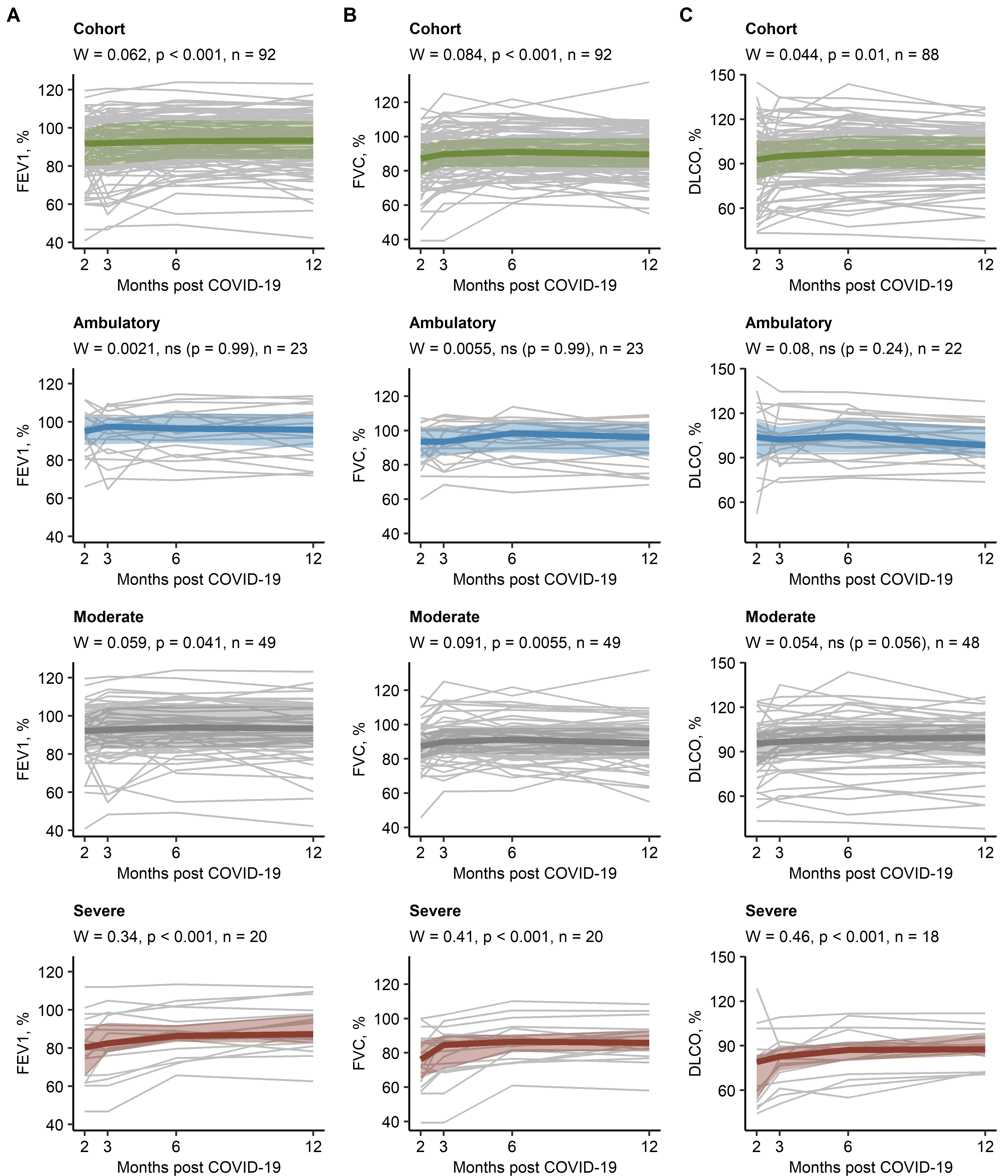
**Supplementary Figure S1. Recovery of fatigue, sleep problems and dyspnea.**

Frequencies of self-reported fatigue (**A**), sleep problems (**B**) and dyspnea (**C**) during acute COVID-19 and at the 2-, 3-, 6-month and 1-year follow-up were investigated in the entire study collective and in ambulatory, moderate and severe COVID-19 survivors. Participants with the complete longitudinal data set were included in the analysis. The recovery was modeled by second-order mixed-effect logistic modeling and likelihood ratio test (full vs null model). P values were corrected for multiple testing with the Benjamini-Hochberg method. Likelihood ratio (), p values and numbers of participants with the complete longitudinal data set are presented in the plot captions.



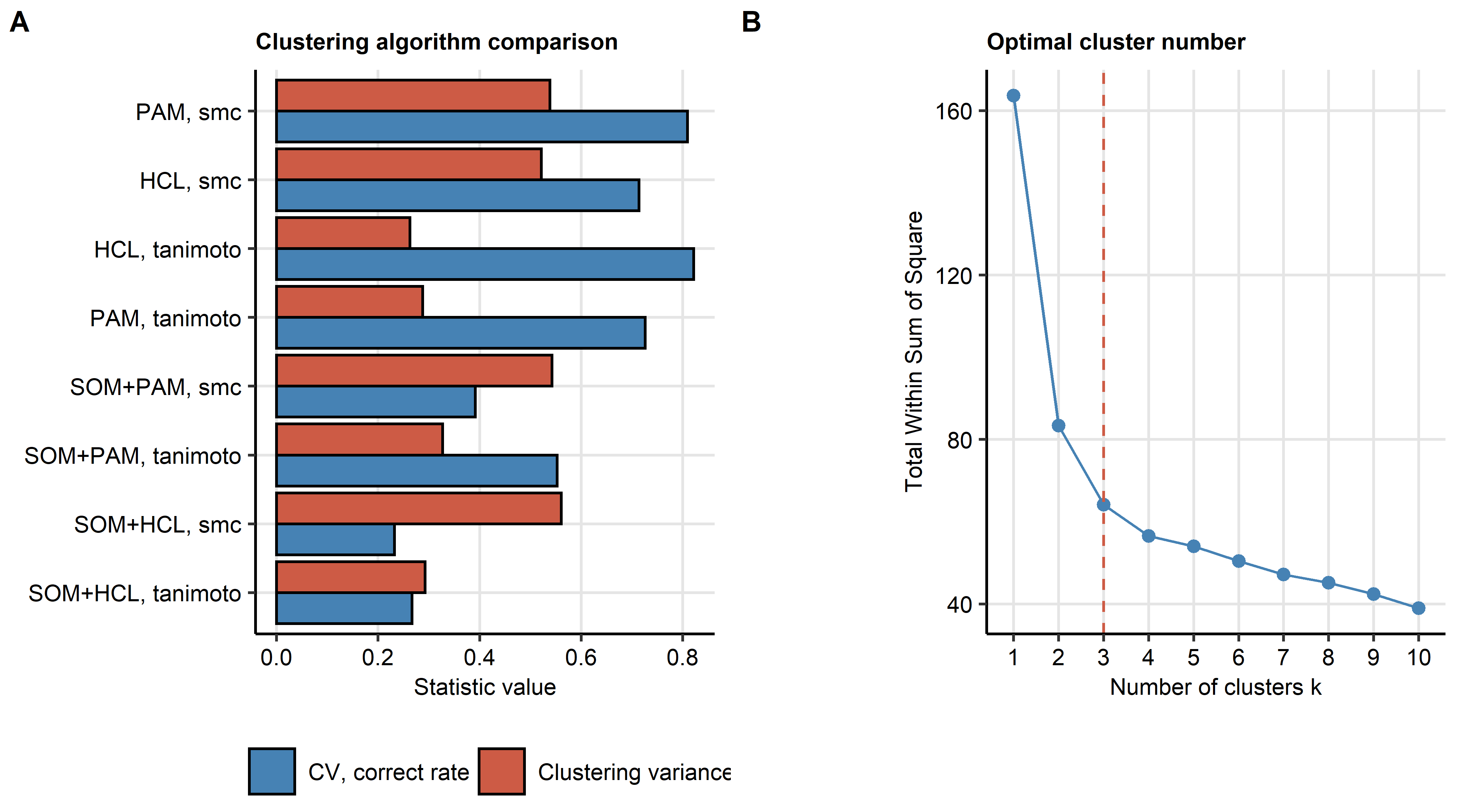
**Supplementary Figure S2. Recovery from night sweating, cough and smell disorders.**

Frequencies of self-reported night sweating (**A**), cough (**B**) and hypo- or anosmia (**C**) during acute COVID-19 and at the 2-, 3-, 6-month and 1-year follow-up were investigated in the entire study collective and in ambulatory, moderate and severe COVID-19 survivors. Participants with the complete longitudinal data set were included in the analysis. The recovery was modeled by second-order mixed-effect logistic modeling and likelihood ratio test (full vs null model). P values were corrected for multiple testing with the Benjamini-Hochberg method. Likelihood ratio (), p values and numbers of participants with the complete longitudinal data set are presented in the plot captions.



**Supplementary Figure S3. Changes in FEV1, FVC and DLCO during COVID-19 convalescence.**

FEV1 (forced expiratory volume in 1 second, **A**), FVC (forced vital capacity, **B**) and DLCO (diffusion lung capacity for carbon monoxide, **C**) were analyzed at the 2-, 3-, 6-month and 1-year follow-up in the entire study collective and in ambulatory, moderate and severe COVID-19 survivors. Participants with the complete longitudinal data set were included in the analysis. The recovery was assessed with Friedman test with Kendall’s W effect size statistic. P values were corrected for multiple testing with the Benjamini-Hochberg method. W, p values and numbers of participants with the complete longitudinal data set are presented in the plot captions. Gray lines indicate parameter values for particular individuals, thick colored lines represented parameter medians, tinted regions indicate interquartile ranges.

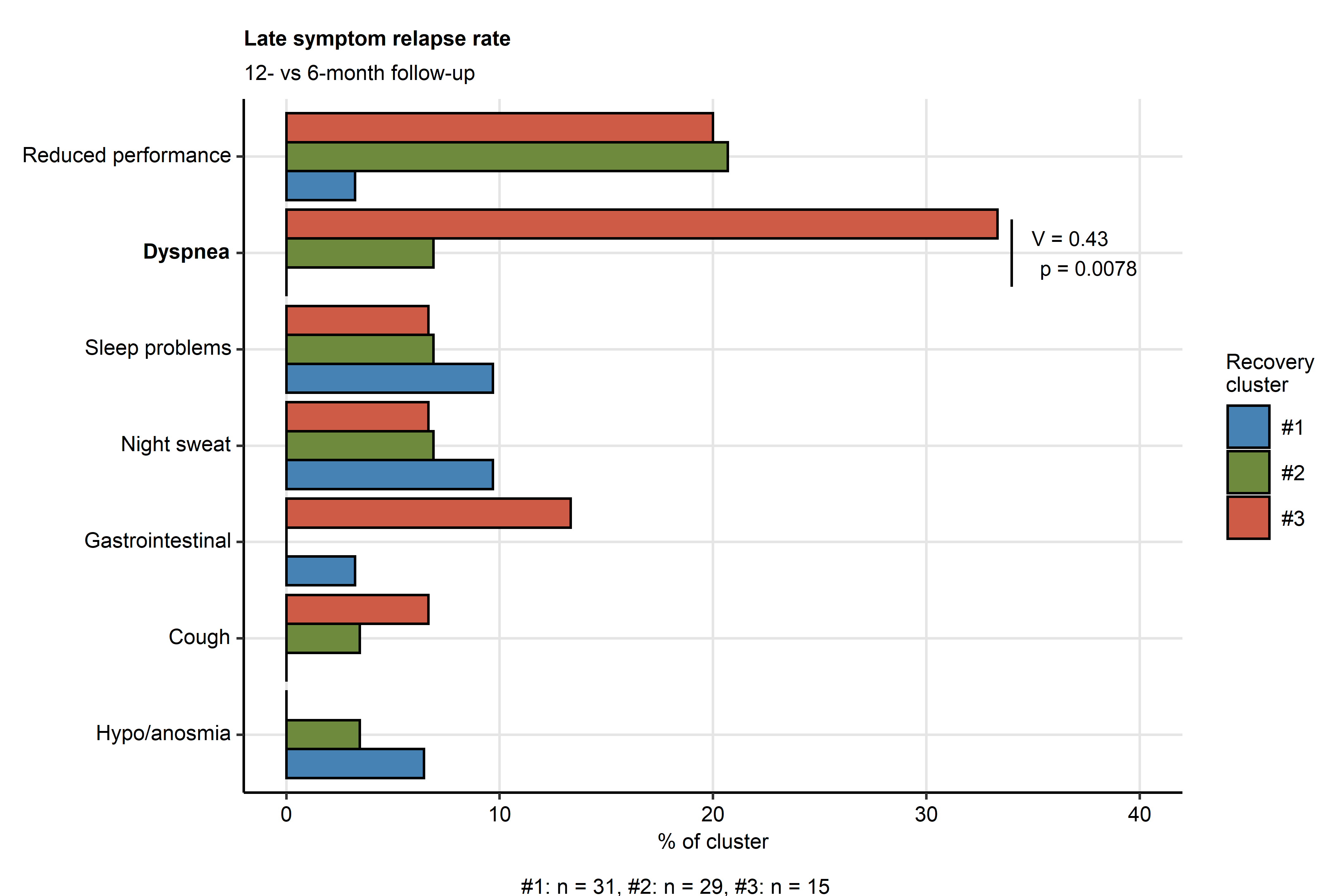


**Supplementary Figure S4. Development of COVID-19 recovery clusters.**

Clustering of the study participants in respect to 19 binary symptom, cardiopulmonary and psychosocial features recorded at the 1-year follow-up (**Supplementary Tables S1** and **S2**) was investigated with the PAM (partitioning around medoids) algorithm with simple matching distance (SMD) statistic. Study participants with the complete clustering data set were included in the analysis (n = 87).

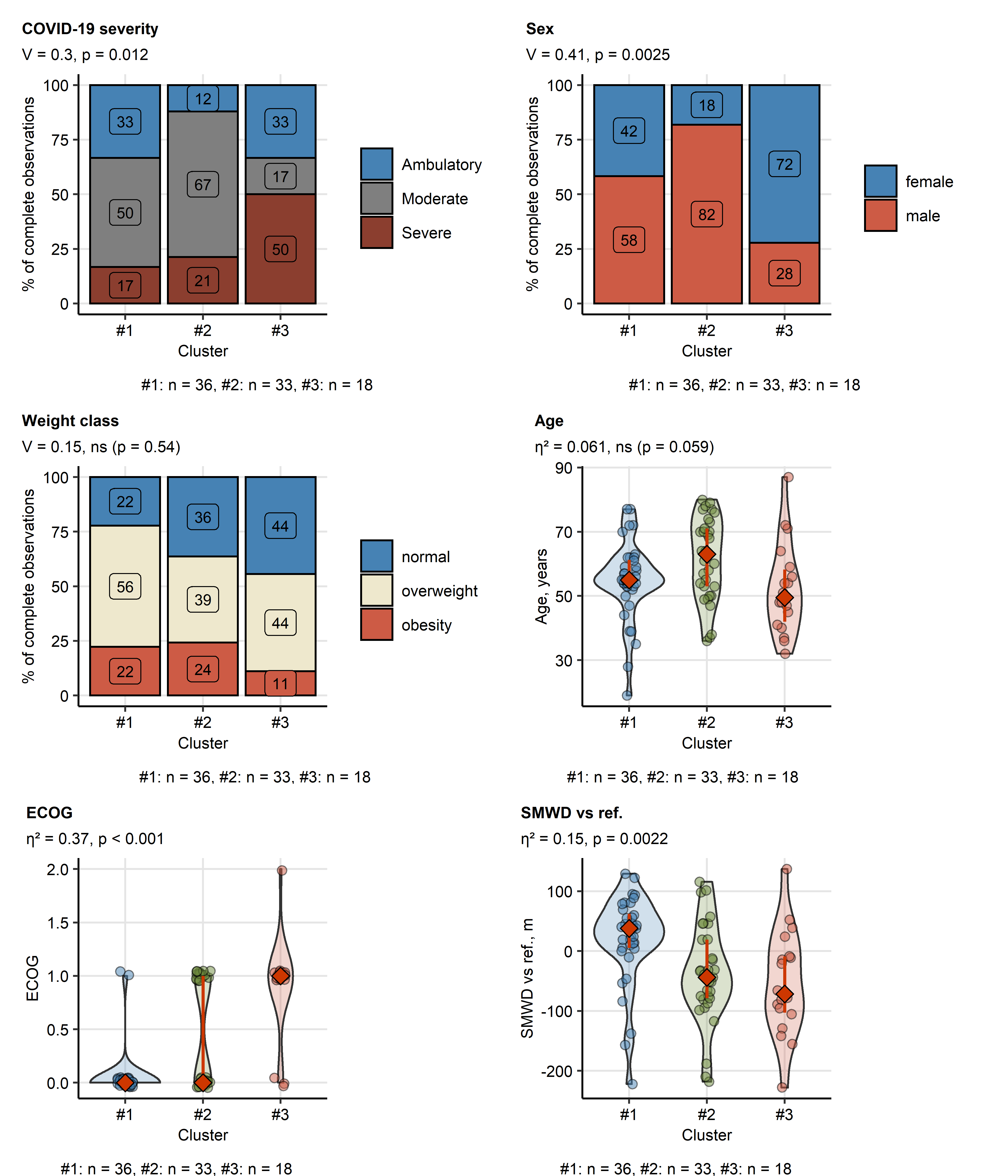
**(A)** Performance comparison of various clustering algorithms and distance measures. Clustering algorithm stability was assessed by the rate of correct cluster assignment in 10-fold cross-validation (CV). Clustering variance is expressed as a ratio of the total between-cluster sum of squares to total sum of squares. Based on the optimal stability, PAM (partitioning around medoids) algorithm with the Manhattan distance statistic was chosen for further analyses.

**(B)** The total within-cluster sum sum of squares as a function of the cluster number for the PAM/SMD clustering procedure. The optimal cluster number (dashed red line) was set at the bend of the curve.



**Supplementary Figure S5. Rates of symptom relapse at the 12-month follow-up in the COVID-19 recovery clusters.**

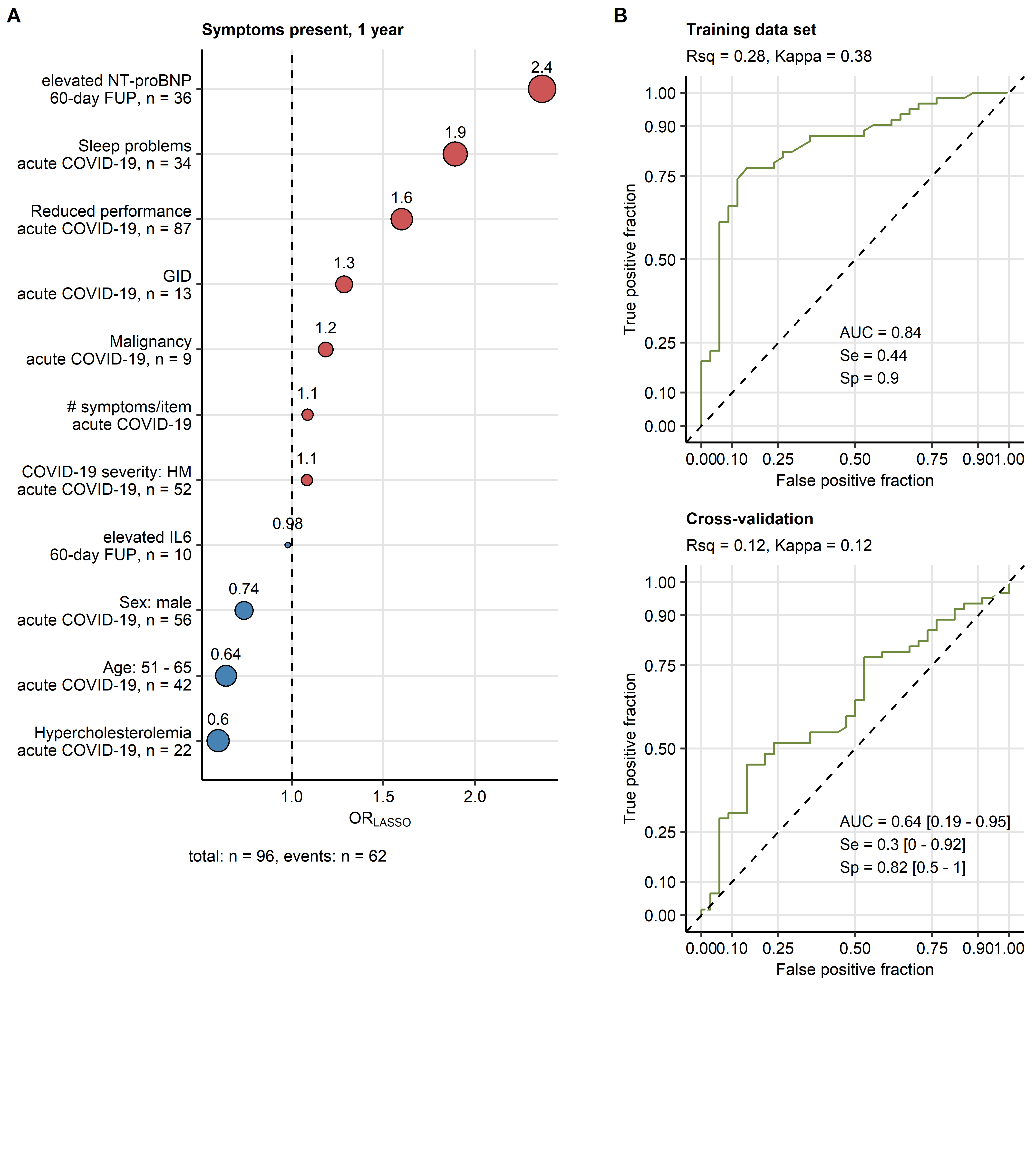
Rates of relapse for particular COVID-19-related symptoms at the 12-month follow-up as compared with the 6-month follow-up were compared between the COVID-19 recovery clusters (**Figure 7**, **Supplementary Figure S4**) with test with Cramer V effect size statistic. Individuals with the complete symptom record for the investigated follow-up were included in the analysis. P values were corrected for multiple testing with Benjamini-Hochberg method. Percentages of the relapsing individuals are presented as a bar plot. Significant differences are highlighted in bold. Effect size statistics and p values for the significant comparisons are indicated in the plot. Numbers of participants assigned to the clusters are presented under the plots.



**Supplementary Figure S6. COVID-19 severity, demographic features, physical performance and mobility in the COVID-19 recovery clusters..**

Acute COVID-19 severity, sex and weight class distribution, age at COVID-19 diagnosis as well as physical performance (ECOG: Eastern Cooperative Oncology Group performance status) and six-minute walking distance (SMWD, difference between the observed and reference value at the 1-year follow-up were compared between the COVID-19 recovery clusters (**Figure 7**, **Supplementary Figure S4**). Statistical significance for numeric values was assessed by Kruskal-Wallis test with effect size statistic or by test with categorical variables with Cramer V effect size statistic. P values were corrected for multiple testing with Benjamini-Hochberg method. Effect size statistic and p values are presented in plot captions. Numbers of participants assigned to the clusters are presented under the plots.

SMWD vs ref.: six-minute walking distance, observed versus reference value.



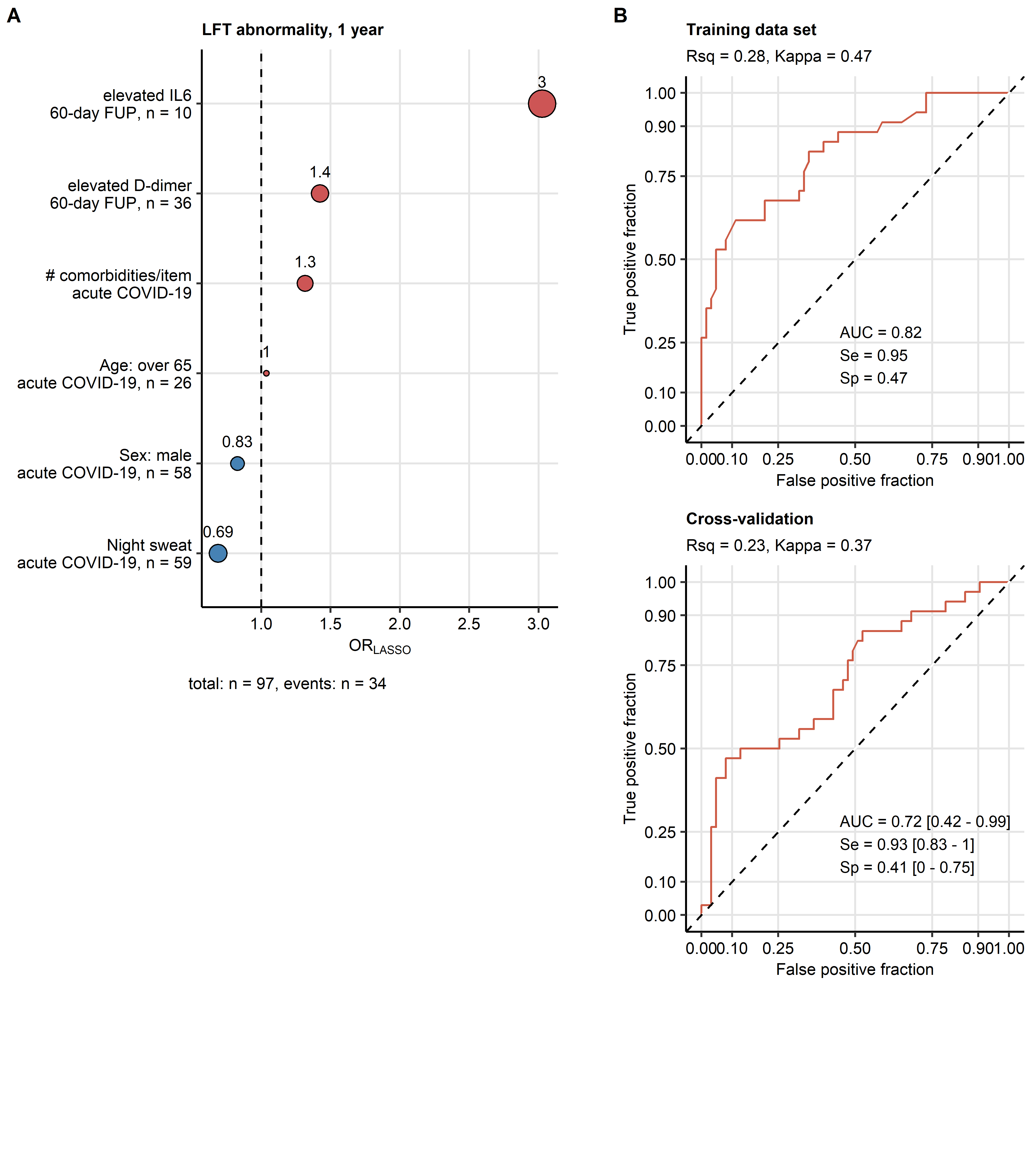
**Supplementary Figure S7. Modeling of the persistent symptom risk at the 1-year post-COVID-19 follow-up.**

The risk of persistent COVID-19 symptoms at the 1-year follow-up was modeled with multi-parameter logistic LASSO (least absolute shrinkage and selection operator) regression and 34 independent variables recorded during acute COVID-19 and at the 60-day follow-up (**Supplementary Table S1**). Study participants with the complete independent variable set were included in the analysis, numbers of complete observations and participants with symptoms are presented in **A**.

**(A)** Non-zero model coefficient values presented as odds ratios (OR). Point size codes for the absolute OR value, point color codes for the correlation with the risk (blue: favorable, red: unfavorable factor).

**(B)** Performance of the LASSO model at predicting the presence of symptoms at the 1-year follow-up in the training data set and 10-fold cross-validation (CV) assessed by receiver-operating characteristic (ROC). Area under the ROC curve (AUC), sensitivity (Se) and specificity (Sp) values with 95 confidence intervals (for CV) are presented in the plots. R-squared (Rsq) and Cohen’s kappa statistic values are shown in the plot captions.

NT-proBNP: N-terminal pro - brain natriuretic peptide; GID: gastrointestinal disease; #: number of; HM: hospitalized moderate COVID-19; IL6: interleukin 6.



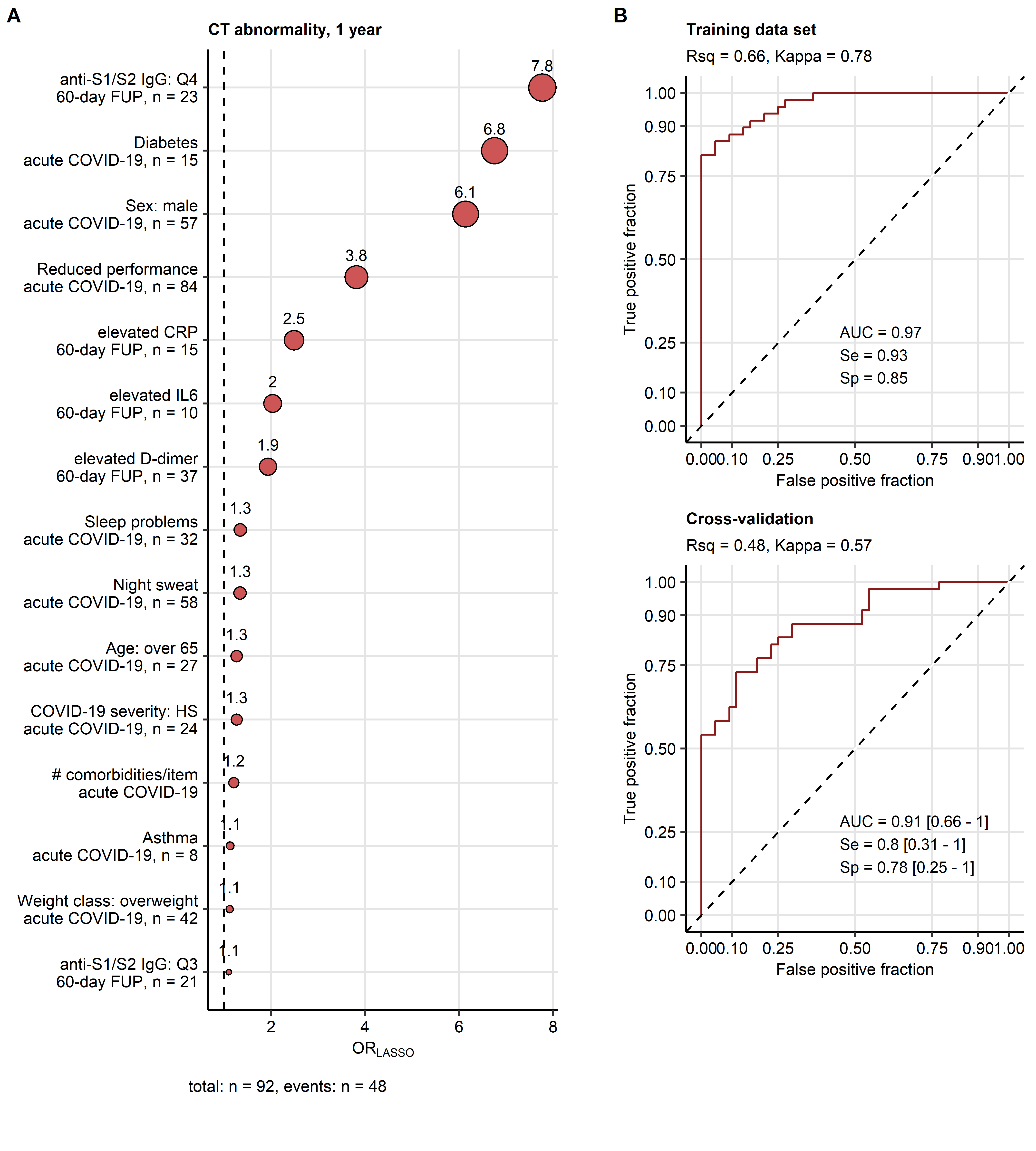
**Supplementary Figure S8. Modeling of the persistent functional lung abnormality at the 1-year post-COVID-19 follow-up.**

The risk of any lung function testing (LFT) abnormality at the 1-year follow-up was modeled with multi-parameter logistic LASSO (least absolute shrinkage and selection operator) regression and 34 independent variables recorded during acute COVID-19 and at the 60-day follow-up (**Supplementary Table S1**). Study participants with the complete independent variable set were included in the analysis, numbers of complete observations and participants with LFT abnormalities are presented in **A**.

**(A)** Non-zero model coefficient values presented as odds ratios (OR). Point size codes for the absolute OR value, point color codes for the correlation with the risk (blue: favorable, red: unfavorable factor).

**(B)** Performance of the LASSO model at predicting the lung function abnormalities at the 1-year follow-up in the training data set and 10-fold cross-validation (CV) assessed by receiver-operating characteristic (ROC). Area under the ROC curve (AUC), sensitivity (Se) and specificity (Sp) values with 95 confidence intervals (for CV) are presented in the plots. R-squared (Rsq) and Cohen’s kappa statistic values are shown in the plot captions.

IL6: interleukin 6; #: number of.



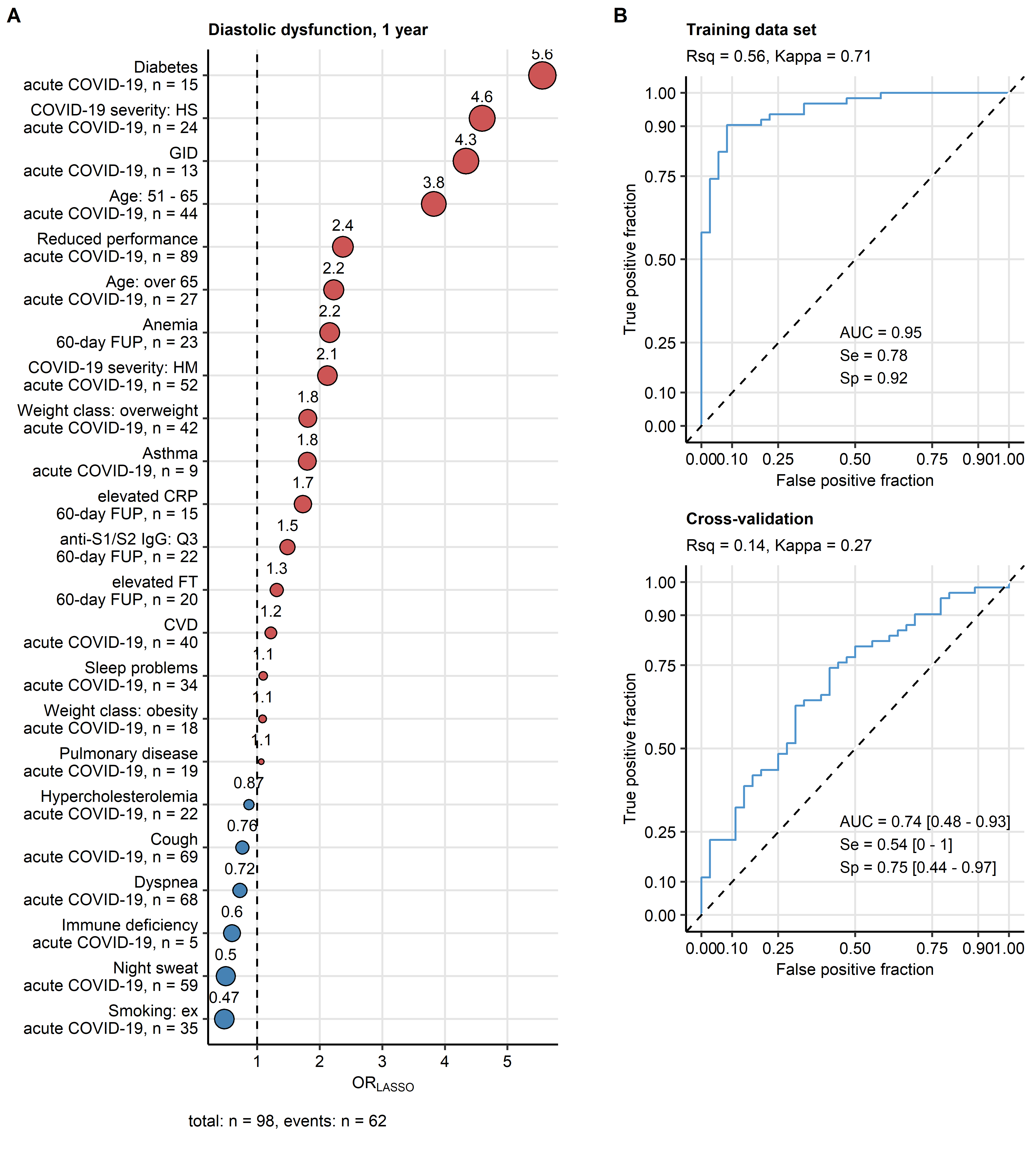
**Supplementary Figure S9. Modeling of the persistent radiological lung abnormality at the 1-year post-COVID-19 follow-up.**

The risk of any chest computed tomography (CT) abnormality at the 1-year follow-up was modeled with multi-parameter logistic LASSO (least absolute shrinkage and selection operator) regression and 34 independent variables recorded during acute COVID-19 and at the 60-day follow-up (**Supplementary Table S1**). Study participants with the complete independent variable set were included in the analysis, numbers of complete observations and participants with CT abnormalities are presented in **A**.

**(A)** Non-zero model coefficient values presented as odds ratios (OR). Point size codes for the absolute OR value, point color codes for the correlation with the risk (blue: favorable, red: unfavorable factor).

**(B)** Performance of the LASSO model at predicting the CT abnormalities at the 1-year follow-up in the training data set and 10-fold cross-validation (CV) assessed by receiver-operating characteristic (ROC). Area under the ROC curve (AUC), sensitivity (Se) and specificity (Sp) values with 95 confidence intervals (for CV) are presented in the plots. R-squared (Rsq) and Cohen’s kappa statistic values are shown in the plot captions.

anti-S1/S2 IgG: anti-S1/S2 SARS-CoV-2 immunoglobulin; Q3, Q4: 3rd and 4th quartile; CRP: C-reactive protein; IL6: interleukin 6; HS: hospitalized severe COVID-19; #: number of.



**Supplementary Figure S10. Modeling of the persistent diastolic dysfunction at the 1-year post-COVID-19 follow-up.**

The risk of diastolic dysfunction at the 1-year follow-up was modeled with multi-parameter logistic LASSO (least absolute shrinkage and selection operator) regression and 34 independent variables recorded during acute COVID-19 and at the 60-day follow-up (**Supplementary Table S1**). Study participants with the complete independent variable set were included in the analysis, numbers of complete observations and participants with diastolic dysfunction are presented in **A**.

**(A)** Non-zero model coefficient values presented as odds ratios (OR). Point size codes for the absolute OR value, point color codes for the correlation with the risk (blue: favorable, red: unfavorable factor).

**(B)** Performance of the LASSO model at predicting diastolic dysfunction at the 1-year follow-up in the training data set and 10-fold cross-validation (CV) assessed by receiver-operating characteristic (ROC). Area under the ROC curve (AUC), sensitivity (Se) and specificity (Sp) values with 95 confidence intervals (for CV) are presented in the plots. R-squared (Rsq) and Cohen’s kappa statistic values are shown in the plot captions.

GID: gastrointestinal disease; HS: hospitalized severe COVID-19; CRP: C-reactive protein; HM: hospitalized moderate COVID-19; FT: ferritin; CVD: cardiovascular disease; anti-S1/S2 IgG: anti-S1/S2 SARS-CoV-2 immunoglobulin; Q4: 4th quartile.

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