Behavioural and psychological patterns of patients with idiopathic pulmonary fibrosis: a prospective study

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Abstract

Background: Idiopathic pulmonary fibrosis (IPF) is a chronic and progressive lung condition. Currently, care models predominantly focus on acute medical and pharmacological needs. As a step towards holistic care, the aim of this prospective study was to investigate the psychological and behavioural needs of IPF patients treated with pirfenidone from diagnosis until two years of follow-up.

Methods: The following variables were selected from the literature on patients’ needs and the COM-B model, a theoretical model explaining behaviour: medication adherence, barriers to adherence, importance and intentions of medication adherence, anxiety, depression, health literacy, knowledge, reported side effects, adherence to sun protection recommendations, alcohol use, physical activity, quality of life and health status. Linear and generalised linear models for longitudinal data were used to evaluate the evolution since treatment initiation.

Results: We included 66 outpatients: 72.7% men, mean age of 70.3 years (range 50–87), predicted mean forced vital capacity of 85.8% (SD 17.4) and predicted mean diffusing capacity for monoxide of 56.9% (SD 15.7). The participants placed considerable importance on following the treatment recommendations. We noticed difficulties regarding health literacy, alcohol use, pirfenidone adherence (decline over time) and adherence to sun protection recommendations (early in follow-up care). There were low levels of physical activity (no effect of time), high body mass indices (decline over time) and moderate levels of depression and anxiety.

Conclusion: When providing care to IPF patients, behavioural issues, health literacy and psychological well-being should be taken into consideration. There is a need to further explore interventions and care models to tackle these difficulties.

Trial registration This study was registered in the ClinicalTrials.gov database (identifier NCT03567785) on May 9th, 2018

Keywords: Idiopathic pulmonary fibrosis, Psychological wellbeing, Behavioural lifestyle

Background

Idiopathic pulmonary fibrosis (IPF) is a lung disease characterized by progressive lung fibrosis and results in a prognosis of 2–5 years postdiagnosis [1, 2]. IPF patients experience physical symptoms including cough, fatigue, and exertional dyspnoea as well as an overall decrease in health-related quality of life (HRQoL) [3, 4]. The provision of care for patients has changed because a...
pharmacological antifibrotic treatment is now available. These drugs slow disease progression and have a beneficial effect on prognosis but do not cure the disease [1]. Moreover, they do not seem to have a positive impact on patients’ HRQoL or symptoms and require long-term medication adherence [5]. Patients may also face burdensome treatment side effects, depressive feelings, and a decrease in daily activities, among other problems [6].

Altogether, these medical and nonmedical needs should be targeted when delivering holistic care, yet the IPF literature and European IPF charter highlight several unmet needs patients experience across their disease trajectory [6, 7]. Available evidence focuses on medical and pharmacological points of view and HRQoL as a general term. Additionally, most evidence stems from patient registries or secondary analyses of clinical trials. Most studies use a cross-sectional design, thereby limiting the understanding of the dynamic evolution of persons’ needs for support over time. Building upon this limited available evidence, we conducted a prospective cohort study with follow-up up to two years after treatment initiation to understand the psychological and behavioural needs of IPF patients.

**Methods**

**Study design and sample**

This prospective cohort study followed the principles laid down in the Declaration of Helsinki and was conducted at the ILD/IPF Centre of the University Hospitals Leuven (Belgium), where approximately 50 patients start pirfenidone treatment each year [8].

Between July 2018 and March 2020, we invited participants who were 18 years or older, Dutch- or French-speaking and diagnosed with IPF. Patients had to start pirfenidone, remain in follow-up at UZ Leuven and be able and willing to provide written informed consent. Patients not managing their medications independently (e.g., patients living in a nursing home) were excluded unless they received help from informal caregivers (i.e., family).

UZ Leuven collaborates with six district general hospitals. Initially, IPF patients being followed-up at a collaborating hospital were not included, but to increase the sample size, an amendment was submitted and approved by the Ethical Committee in July 2019, allowing us to enrol these patients also.

Figure 1 provides an overview of the collected variables and the study visits that took place during a scheduled outpatient clinic visit. In routine care, patients received a face-to-face group information session at treatment initiation, after which we enrolled eligible patients (Visit 1). All patients were then followed up every three months, with an additional consultation six weeks after treatment initiation. If a face-to-face study follow-up visit was not possible (e.g., during the COVID-19 pandemic) or when patients were followed-up by a collaborating hospital, we sent the questionnaires by post to the patient’s home at the time of the planned data collection points. Due to the COVID-19 pandemic, data collection was paused between March 2020 and May 2020. Data collection was ended in February 2021; hence, not all enrolled patients reached the two-year follow-up time point.

**Variables and measurements**

Variables were selected based on the existing IPF literature and the COM-B model, a theoretical framework explaining behaviour. The COM-B model states that behaviour (e.g., taking medication, applying sun protection measures) is the result of capability, opportunity and motivation [9]. The questionnaire bundle consisted of following variables and their measurement instrument (Additional file 1: Table S2): medication adherence

| Socio-demographic (Q) | Health literacy (Q) | IPF knowledge (Q) | Adherence to pirfenidone (Q) | Intentions (Q) | Barriers for adherence (Q) | Side effects (Q) | Depression (Q) | Anxiety (Q) | Sun protection (Q) | Physical activity (Q) | Smoking status (Q) | Alcohol intake (Q) | Quality of life/health status (Q) | Clinical variables |
|-----------------------|--------------------|------------------|-----------------------------|---------------|--------------------------|-----------------|----------------|-----------|-------------------|-------------------|------------------|----------------|---------------------|--------------------|
| Visit 1: Inclusion    | X                  |                  | X                           |               | X                        | X               | X              | X         | X                 | X                 | X                | X                  | X              | X                     | X                  |
| Visit 2: Six weeks    | X                  |                  | X                           | X             | X                        | X               | X              | X         | X                 | X                 | X                | X                  | X              | X                     | X                  |
| Visit 3: Three months | X                  |                  | X                           | X             | X                        | X               | X              | X         | X                 | X                 | X                | X                  | X              | X                     | X                  |
| Visit 4: One year     |                    |                  |                             |               |                          |                 | X              | X         | X                 | X                 | X                | X                  | X              | X                     | X                  |
| Visit 5: 1.5 years    |                    |                  |                             |               |                          |                 |                |           | X                 | X                 | X                | X                  | X              | X                     | X                  |
| Visit 6: Two years    |                    |                  |                             |               |                          |                 |                |           |                   | X                 | X                | X                  | X              | X                     | X                  |

**Fig. 1** Study visits and variables
Data collected n=66 Data collected n=62 Data collected n=61 Data collected n=46 Data collected n=27 Data collected n=11

Fig. 2 Study Flowchart. ‘Study discontinuation’ refers to the patients who had a data collection point planned but discontinued the study (e.g., deceased, medication switch). ‘No data collection’ refers to the patients who did not have a new data collection point planned and thus ended the study as anticipated (e.g., due to the prospective inclusion and design of the study).
follow-up point, respectively. A significant decline in DLco was observed between baseline and the one-year timepoint, the 1.5-year timepoint and the two-year timepoint. For the pairwise comparisons of the FVC mean values, no significant differences between time points were noted (see Additional file 1: Table S4).

There was a significant decrease in mean BMI over time \((p=0.006)\) from baseline to year one, as well as from year one to year two. A total of 80.3% of the participants \((n=53)\) were overweight at baseline, of which 40.9% \((n=27)\) were considered obese. At the one-year follow-up, 17 out of 45 participants \((37.8%)\) were considered obese, and 19 \((42.2%)\) were considered overweight. None of our participants were underweight during the study period. The BMI category did not change significantly over time \((p=0.099)\).

### Patient-reported variables

Table 3 and the supplementary material (S) provide additional information on the patient-reported characteristics. The total depression score did not change significantly over time \((p=0.379)\). Eight out of 50 participants \((16%)\) had a moderate level of depression, of whom three were moderately severely depressed at the six-week timepoint. Three months and one year after treatment initiation, we observed moderate levels of depression in 16.3% \((n=8/49)\) and 8.8% \((n=3/34)\) of the participants, respectively.

For the total anxiety score, a significant decline over time was observed \((p<0.0001)\). Between the six-week and one-year follow-up points, the difference was not significant, but a significant decline in anxiety was observed between the six-week and two-year follow-up points \((p<0.0001)\). We observed moderate levels of anxiety in 17.6% \((n=9/51)\) and 8.6% \((n=3/35)\) of the participants at the six-week and one-year follow-ups, respectively.

Table 1

| Sociodemographic characteristics \((n=66)\) |
|---------------------------------------------|
| **Baseline (Visit 1)**                      |
| **Sex, \(n (\%)\)**                         |
| Male                                        | 48 (72.7) |
| Female                                      | 18 (27.3) |
| **Age (years)**                             |
| Mean (SD)                                   | 70.3 (8.4) |
| Range                                       | 50–87 |
| Median (IQR)                                | 72 (11) |
| **Ethnicity**                               |
| Caucasian, \(n (\%)\)                      | 66 (100) |
| **Marital status, \(n (\%)\)**              |
| Partner                                     | 54 (81.8) |
| No partner                                  | 12 (18.2) |
| **Education level, \(n (\%)\)**             |
| Lower education                             | 20 (30.3) |
| Moderate education                          | 32 (48.5) |
| High education                              | 14 (21.2) |
| **Employment status, \(n (\%)\)**           |
| Yes                                         | 9 (13.6) |
| Fulltime                                    | 5 (55.6) |
| Halftime                                    | 4 (44.4) |
| No                                          | 57 (86.4) |
| Retired                                     | 53 (93) |
| Stay-at-home partner                        | 3 (5.3) |
| No further information provided              | 1 (1.7) |
| **One-way distance from the clinic (km)**    |
| Mean (SD)                                   | 71.9 (41.8) |
| Range                                       | 3–166 |
| Median (IQR)                                | 65 (62) |

Overall, there was a significant decrease in pirfenidone adherence rates over time \((p=0.0268)\) with the predicted adherence \((95\% \text{ CI})\) being 90.4% \((77.0; 96.3)\) at baseline and 75.9% \((55.5; 88.8)\) at the one-year time point. Drug holidays were observed in 18.2% \((n=2/11)\) and 23% \((n=3/12)\) of participants at week six and at the three-month follow-up, respectively. Overall, intention to adhere to one’s treatment was high, and participants deemed taking medication important. Barriers to adherence are reported in Additional file 1: Table S3.

A significant effect of time was observed for the total number of self-reported side effects \((p=0.0002)\). More side effects were reported at the three-month time point than at week six and to the one-year time point. The reported side effects can be found in Additional file 1: Table S3. Regarding the use of sunscreen, a significant increase in adherence rates was observed over time \((p=0.0245)\). Participants were more likely to be nonadherent to the use of sunscreen at week six \((42.3\%\) compared to later timepoints.

A total of 20.3% \((n=13/64)\) of participants were classified as having suboptimal health literacy. No overall significant change in mean knowledge scores over time was observed \((p=0.802)\). Participants’ knowledge of the disease and treatment was high, with an estimated mean score \((95\% \text{ CI})\) of 5.7 \((5.4; 5.9)\) and 5.6 \((5.3; 5.9)\) at the 6-week and 1-year follow-ups, respectively. The proportion of patients providing a wrong answer was highest for
|                       | Baseline (Visit 1) | Three months (Visit 3) | One year (Visit 4) | One year and a half (Visit 5) | Two years (Visit 6) | Main effect of time |
|-----------------------|--------------------|------------------------|-------------------|-------------------------------|--------------------|--------------------|
| **Weight (Kg)**       | N 66               | 45                     | 5                  | 12                            |                    |                    |
| Mean (SD)             | 82.5 (16)          | 83 (13.1)              | 82.3 (10.6)        |                               |                    |                    |
| Range                 | 46–129             | 54–112                 | 71–100             |                               |                    |                    |
| Median (IQR)          | 83 (17.6)          | 83 (18)                | 79.5 (20)          |                               |                    |                    |
| Estimated mean (95% CI)* | 82.5 (78.6; 86.4) | 80.9 (77.2; 84.6)      | 78.6 (74.5; 81.7)  |                               |                    | p = 0.0110         |
| **BMI (Kg/m²)**       | N 66               | 45                     | 5                  | 12                            |                    |                    |
| Mean (SD)             | 28.6 (4.4)         | 28.5 (4)               | 27.3 (3.3)         |                               |                    |                    |
| Range                 | 19.8–40.4          | 21–37.8                | 20.8–32.4          |                               |                    |                    |
| Median (IQR)          | 28.1 (5.9)         | 27.6 (6.4)             | 27.1 (3.2)         |                               |                    |                    |
| Estimated mean (95% CI)* | 28.6 (27.5; 29.6) | 28.0 (27.2; 29)        | 27.1 (26; 28.3)    |                               |                    | p = 0.0061         |
| **BMI category**      | N 66               | 45                     | 5                  | 12                            |                    |                    |
| Underweight n (%)     | 0                  | 0                      | 0                  |                               |                    |                    |
| Normal n (%)          | 13 (9.7)           | 9 (20.0)               | 2 (16.7)           |                               |                    |                    |
| Overweight n (%)      | 27 (40.9)          | 19 (42.2)              | 8 (66.7)           |                               |                    |                    |
| Obese n (%)           | 26 (39.4)          | 17 (37.8)              | 2 (16.7)           |                               |                    |                    |
| Main effect of time*  |                    |                        |                    |                               |                    | p = 0.0992         |
| **Smoking status**    | N 66               | 45                     | 5                  | 12                            |                    |                    |
| Never n (%)           | 14 (21.2)          | 7 (15.2)               | 2 (15.4)           |                               |                    |                    |
| Former n (%)          | 52 (78.8)          | 5 (83.3)               | 2 (100)            |                               |                    |                    |
| **Oxygen use**        | N 66               | 46                     | 5                  | 13                            |                    |                    |
| Yes n (%)             | 1 (1.5)            | 7 (15.2)               | 2 (15.4)           |                               |                    |                    |
| Continuous use        | 1 (100)            | 6 (100)                | 2 (100)            |                               |                    |                    |
| **Gastro oesophageal reflux** | N 66 | 45 | 5 | 12 | | |
| Yes n (%)             | 9 (13.6)           | 1 (16.7)               | 0                  |                               |                    |                    |
| **DLco % predicted**  | N 65               | 56                     | 44                 | 21                            | 11                 | p < 0.0001         |
| Mean (SD)             | 56.9 (15.7)        | 57.8 (15)              | 55.1 (15.4)        | 54.2 (17.4)                    | 58.5 (17.6)        |                    |
| Range                 | 24–111             | 21–103                 | 23–99              | 25–102                        | 42–98              |                    |
| Median (IQR)          | 57 (18)            | 56.5 (13)              | 56 (17)            | 51 (19)                       | 51 (22)            |                    |
| Estimated mean (95% CI)* | 56.9 (53; 60.7)   | 56.1 (52.3; 59.8)      | 53.2 (49.1; 57.3)  | 50.9 (46.3; 55.5)              | 49 (44.4; 53.7)    | p < 0.0001         |
| **FVC % predicted**   | N 65               | 58                     | 44                 | 21                            | 12                 |                    |
| Mean (SD)             | 85.8 (17.4)        | 88.9 (21)              | 87 (19.1)          | 85 (23.9)                      | 79.9 (17.8)        |                    |
| Range                 | 50–126             | 42–147                 | 54–120             | 39–116                        | 46–101             |                    |
| Median (IQR)          | 88 (26)            | 89 (29)                | 88.5 (28)          | 89 (28)                       | 84 (23)            |                    |
| Estimated mean (95% CI)* | 85.8 (81.5; 90.1) | 87.7 (82.5; 92.8)      | 86.1 (80.6; 91.5)  | 83.8 (77; 90.7)                | 84.5 (77.4; 91.7)  | p = 0.0007         |
| **GAP index**         | N 65               | 44                     | 24                 | 7                              | 11                 |                    |
| Stage 1 n (%)         | 35 (13.8)          | 24 (54.5)              | 7 (63.6)           |                               |                    |                    |
| Stage 2 n (%)         | 30 (46.2)          | 17 (38.6)              | 4 (36.4)           |                               |                    |                    |
| Stage 3 n (%)         | 0                  | 3 (6.8)                | 0                  |                               |                    |                    |
| **6MWD Meters**       | N 18               | 5                      | 1                  |                               |                    |                    |
| Mean (SD)             | 474.9 (128.3)      | 325.4 (225.5)          | NA                 |                               |                    |                    |
| Range                 | 140–666            | 50–595                 | 300                |                               |                    |                    |
| Median (IQR)          | 461 (164)          | 388 (430)              | NA                 |                               |                    |                    |
the statement ‘pirfenidone repairs damaged lung tissue’ (14%).

No significant effect of time was observed on physical activity (p = 0.227). The predicted percentage (95% CI) of participants being physically inactive was 59.2% (44.1; 72.7) and 42.9% (23; 65.4) at the three-month and 1.5-year follow-up points, respectively. Fifteen patients (30.6%) had at-risk alcohol drinking behaviour at month three, and seven patients (31.8%) had at-risk alcohol drinking behaviour at the 1.5-year time point. Of these seven patients, five showed at-risk behaviour at the three-month follow-up.

**Discussion**

To our knowledge, this is the first prospective study that investigated the prevalence of behavioural and psychological needs of persons with IPF and their evolution over time up to two years after diagnosis. We identified a need for support regarding health literacy, medication adherence, mental health, and lifestyle behaviours. Below, we discuss our results in light of available evidence and the implications for further research and clinical care.

We are the first to document health literacy in IPF patients. A total of 20.3% of the participants had inadequate health literacy skills, which is higher than the 11.6% prevalence reported in the Belgian national health survey, although a different questionnaire was used [22]. Poor health literacy is associated with poorer knowledge regarding disease and treatment, a poorer adherence, and might result in negative health outcomes and higher health care resource use [23, 24].

Overall, our participants overall had high levels of disease- and treatment-related knowledge, which did not significantly change over time. However, patients with poor levels of knowledge, and low health literacy should be targeted for additional support.

Second, participants were highly motivated and deemed taking medication important, confirming available evidence [25]. Only 3% of our participants reported having discontinued pirfenidone based on their own initiative, which is slightly lower than other real-world studies, reporting a 5.5–6% discontinuation rate of pirfenidone [26, 27]. However, we detected problems with adherence already early after treatment initiation (19.6% at week six), and nonadherence increased over time (up to 36.4% at year two). Another prospective study reported a prevalence of self-reported nonadherence of 12% at month six [25]. Our findings presumably underestimate the true issue of nonadherence, given that we used self-report, yet self-report questionnaires are an easy-to-use method to detect at least some of the patients who need support [28]. In our study, we noted several barriers that may affect adherence, such as forgetfulness or the presence of side effects, which might form a good basis for tailored adherence interventions.

Third, we showed high numbers of nonadherence to sun protection (especially at the start of treatment, 42.3%), despite its importance in mitigating the phototoxicity side effect of pirfenidone [15]. These numbers are in line with the high numbers (51.4%) observed in a Belgian heart transplantation population [29]. More research is needed to understand IPF patients’ barriers to using sun protection to develop supportive interventions.

Fourth, shortly after treatment initiation, 16% and 17.6% of our participants had moderate levels of depression and anxiety, respectively. Over time, we found no significant change in levels of depression, but lower levels of anxiety were reported. These levels were also lower than those described in other papers on IPF (24.3–49.2% for depression), but comparisons should be performed carefully as we used different questionnaires (i.e., the validated GAD7 and PHQ9) [30, 31]. Selection bias or
| Table 3  Patient-reported characteristics | Baseline (Visit 1) | Six weeks (Visit 2) | Three months (Visit 3) | One year (Visit 4) | One year and a half (Visit 5) | Two years (Visit 6) | Main effect of time |
|-----------------------------------------|-------------------|---------------------|-----------------------|------------------|-------------------------------|-------------------|-------------------|
| **Health literacy**<sup>a</sup>          | N 64              | Inadequate n (%) 13 (20.3) |
| **Knowledge**                           | N 50             | Median total score (IQR) 6 (0) | 6 (0) | 6 (1) |
|                                         | Knowledge N 50   | Mean (SD) 5.7 (1) | 5.6 (0.9) | 5.6 (0.7) |
|                                         | Range 1–6        | Score lower than 6, n (%) 8 (16) | 8 (23.5) | 3 (33.3) |
|                                         | Estimated mean score (95% CI) 5.7 (5.4; 5.9) | 5.6 (5.3; 5.9) | 5.6 (5.3; 5.9) | p = 0.8021 |
| **Side effects to pirfenidone**         | N 54             | Patients experiencing min 1 side effect, n (%) 38 (70.4) | 40 (78.4) | 24 (63.2) | 15 (68.2) | 9 (81.8) |
|                                         | Number of side effects/patients Median (IQR) 1 (2) | 1 (1) | 1 (3) | 1.5 (3) | 3 (4) |
|                                         | Range 0–5 | 0–7 | 0–5 | 0–4 | 0–6 |
|                                         | Predicted mean count (95% CI) 1.4 (1.1; 1.7) | 2.5 (2; 3) | 1.5 (1.1; 2) | 1.5 (1.1; 2.2) | 2.7 (1.8; 4) | p = 0.0002 |
| **Depression**                           | N 50             | Total score: median (IQR) 3 (6) | 4 (7) | 3 (6) | 2 (4) |
|                                         | Total score: range 0–17 | 0–14 | 0–13 | 0–7 |
|                                         | Moderate depression n (%) 8 (16) | 8 (16.3) | 3 (8.8) | 0 |
|                                         | Estimated mean total score (95% CI) 2.9 (2.1; 3.9) | 2.8 (2; 3.8) | 2.2 (1.5; 3.1) | 2.3 (1.3; 3.9) | p = 0.3785 |
| **Anxiety**                             | N 51             | Total score: median (IQR) 4 (11) | 2 (5) | 3.5 (6) |
|                                         | Total score: range 0–18 | 0–15 | 0–9 |
|                                         | Moderate anxiety n (%) 9 (17.6) | 3 (8.6) | 0 |
|                                         | Estimated mean total score (95% CI) 3 (1.9;4.7) | 1.9 (1.1;3) | 0.8 (0.1;1.9) | p < 0.0001 |
Table 3 (continued)

|                          | Baseline (Visit 1) | Six weeks (Visit 2) | Three months (Visit 3) | One year (Visit 4) | One year and a half (Visit 5) | Two years (Visit 6) | Main effect of time |
|--------------------------|--------------------|---------------------|------------------------|-------------------|-------------------------------|---------------------|---------------------|
| Intention to be adherent to treatment¹ |                    |                     |                        |                   |                               |                     |                     |
| 'I expect to […]'        | N 60               | 51                  | 35                     | 10                |                               |                     |                     |
|                          | Mean (SD)          | 6.8 (0.9)           | 6.8 (0.5)              | 6.7 (1)           | 7 (0)                         |                     |                     |
|                          | Range              | 1–7                 | 5–7                    | 1–7               | 7                             |                     |                     |
| 'I want to […]'          | N 60               | 51                  | 35                     | 10                |                               |                     |                     |
|                          | Mean (SD)          | 6.9 (0.8)           | 6.9 (0.3)              | 6.8 (0.9)         | 7 (0)                         |                     |                     |
|                          | Range              | 1–7                 | 5–7                    | 1–7               | 7                             |                     |                     |
| 'I intend to […]'        | N 60               | 51                  | 35                     | 10                |                               |                     |                     |
|                          | Mean (SD)          | 6.9 (0.8)           | 6.9 (0.3)              | 6.8 (0.9)         | 7 (0)                         |                     |                     |
|                          | Range              | 1–7                 | 5–7                    | 1–7               | 7                             |                     |                     |
| Level of motivation²     | N 59               | 53                  | 35                     | 10                |                               |                     |                     |
|                          | Pre-Contemplation n | 0                   | 0                      | 0                 | 0                             |                     |                     |
|                          | Contemplation n    | 2                   | 0                      | 0                 | 0                             |                     |                     |
|                          | Sufficient motiva- | 57                  | 53                     | 35                | 10                            |                     |                     |
| Pantoprazole medication adherence |                    |                     |                        |                   |                               |                     |                     |
| Taking non-adherence n (%) | 11 (19.6)         | 12 (24.5)           | 12 (32.4)             | 8 (33.3)          | 4 (36.4)                      |                     |                     |
| Drug holiday n (%)       | N 11               | 12                  | 12                     | 8                 | 4                             |                     |                     |
|                          | 2 (18.2)           | 3 (25)              | 0                      | 1 (12.5)          | 0                             |                     |                     |
| Dosing non-adherence n (%) | N 56               | 47                  | 36                     | 24                | 11                            |                     |                     |
|                          | 0                  | 0                   | 0                      | 1 (4.2)           | 0                             |                     |                     |
| Discontinuation n (%)    | N 56               | 8                   | 36                     | 24                | 11                            |                     |                     |
|                          | 0                  | 0                   | 0                      | 0                 | 0                             |                     |                     |
| Omitted to take pirfenidone n (%) | 10                 | 12                  | 12                     | 8                 | 4                             |                     |                     |
|                          | 1 time             | 7 (70)              | 6 (50)                 | 7 (58.3)          | 4 (50)                        | 2 (50)              |                     |
|                          | 2 times            | 1 (10)              | 4 (33.3)               | 2 (16.7)          | 2 (25)                        | 1 (25)              |                     |
|                          | 3 times            | 0                   | 1 (8.3)                | 1 (8.3)           | 1 (12.5)                      | 0                   |                     |
|                          | 4 times            | 1 (10)              | 0                      | 2 (16.7)          | 0                             | 1 (25)              |                     |
|                          | More than 4 times  | 1 (10)              | 1 (8.3)                | 0                 | 1 (12.5)                      | 0                   |                     |
| Predicted % of taking adherence (95%CI) | 90.4 (77.9;96.3)   | 86.7 (71.9;94.5)     | 75.9 (55.5;88.8)     | 67.9 (46.2;83.9)   | 67.9 (46.2;83.9)              | p = 0.0268          |                     |

Pantoprazole medication adherence³

|                          | Baseline (Visit 1) | Six weeks (Visit 2) | Three months (Visit 3) | One year (Visit 4) | One year and a half (Visit 5) | Two years (Visit 6) | Main effect of time |
|--------------------------|--------------------|---------------------|------------------------|-------------------|-------------------------------|---------------------|---------------------|
| Pantoprazole intake yes n (%) | 42 (75)           | 38 (82.6)           | 25 (69.4)              | 18 (75)           | 10 (90.9)                      |                     |                     |
| Taking non-adherence n (%) | 3 (7.5)            | 1 (2.7)             | 1 (4)                  | 0                 | 1 (10.0)                       |                     |                     |
Table 3 (continued)

|                                | Baseline (Visit 1) | Six weeks (Visit 2) | Three months (Visit 3) | One year (Visit 4) | One year and a half (Visit 5) | Two years (Visit 6) | Main effect of time |
|--------------------------------|--------------------|---------------------|------------------------|-------------------|-------------------------------|---------------------|---------------------|
| **Barriers to medication adherence** | N                  | 51                  | 35                     | 8                 |                               |                     |                     |
| Total numbers of barriers     |                    |                     |                        |                   |                               |                     |                     |
| Median (IQR)                  | 1 (4)              | 4 (4)               | 4 (6)                  |                   |                               |                     |                     |
| Mean (SD)                     | 3 (4.2)            | 3.7 (3.4)           | 5.6 (5.9)              |                   |                               |                     |                     |
| Range                         | 0–15               | 0–11                | 19–Jan                 |                   |                               |                     |                     |
| Persons having min 1 barrier n (%) | 35 (68.6)         | 27 (77.1)           | 8 (100)                |                   |                               |                     |                     |
| **Sun protection**            | N                  | 52                  | 48                     | 35                | 23                            | 10                  | p = 0.0245          |
| Inadequate sunscreen use      |                    |                     |                        |                   |                               |                     |                     |
| N                             | 52                 | 46                  | 34                     | 23                | 9                             |                     |                     |
| Mean (SD)                     | 22 (42.3)          | 20 (41.7)           | 6 (17.1)               | 4 (17.4)          | 3 (30.0)                      |                     |                     |
| Inadequate use of protective clothes | 18 (34.6)     | 14 (30.4)           | 7 (20.6)               | 5 (21.7)          | 4 (44.4)                      |                     |                     |
| Not staying in the shadows    |                    |                     |                        |                   |                               |                     |                     |
| Use of sunscreen*             |                    |                     |                        |                   |                               |                     |                     |
| N                             | 52                 | 46                  | 34                     | 23                | 9                             |                     |                     |
| Mean (SD)                     | 14 (26.9)          | 11 (23.9)           | 7 (20.6)               | 5 (21.7)          | 2 (22.2)                      |                     |                     |
| Physical inactivity           | N                  | 49                  | 21                     |                   |                               |                     |                     |
| Not sufficiently active n (%) |                    |                     |                        |                   |                               |                     |                     |
| Predicted timepoint % (95% CI)|                    |                     |                        |                   |                               |                     |                     |
| N                             | 49                 | 28 (57.1)           | 9 (42.9)               |                   |                               |                     |                     |
| Alcohol use*                  | N                  | 49                  | 22                     |                   |                               |                     |                     |
| At-risk drinking behaviour n (%) |                    |                     |                        |                   |                               |                     |                     |
| N                             | 49                 | 15 (30.6)           | 7 (31.8)               |                   |                               |                     |                     |
| Perceived health status EQ-5D-5L | Descriptive health index, N | 62 | 50 | 35 | 22 | 10 | | |
| Median (IQR)                  | 0.840 (0.210)      | 0.863 (0.225)       | 0.857 (0.219)          | 0.753 (0.218)     | 0.824 (0.183)                 |                     |                     |
| Mean (SD)                     | 0.780 (0.189)      | 0.804 (0.193)       | 0.824 (0.161)          | 0.760 (0.174)     | 0.794 (0.205)                 |                     |                     |
| Global health score based on the VAS, N |                  |                     |                        |                   |                               |                     |                     |
| Median (IQR)                  | 62                 | 51                  | 35                     | 22                | 10                            |                     |                     |
| Mean (SD)                     | 70 (23)            | 70 (20)             | 70 (20)                | 66 (21)           | 62.5 (26)                     |                     |                     |
| Range                         | 67.8 (15.8)        | 68.4 (20.3)         | 68 (13)                | 63.6 (15.9)       | 66.8 (17.4)                   |                     |                     |
| Frequencies reported problems, N |                    |                     |                        |                   |                               |                     |                     |
| Mobility, n (%)               | 64                 | 51                  | 35                     | 22                | 10                            |                     |                     |
| No problems                   |                    |                     |                        |                   |                               |                     |                     |
| Slight-extreme problems       | 28 (43.8)          | 24 (47.1)           | 17 (48.6)              | 9 (40.9)          | 7 (70)                        |                     |                     |
Table 3 (continued)

|                          | Baseline (Visit 1) | Six weeks (Visit 2) | Three months (Visit 3) | One year (Visit 4) | One year and a half (Visit 5) | Two years (Visit 6) | Main effect of time |
|--------------------------|--------------------|---------------------|------------------------|-------------------|-----------------------------|-------------------|---------------------|
| **Self-care, n (%)**     |                    |                     |                        |                   |                             |                   |                     |
| No problems              | 36 (56.2)          | 27 (52.9)           | 18 (51.4)              | 13 (59.1)         | 3 (30)                      |                   |                     |
| Slight-extreme problems  | 47 (73.4)          | 36 (72)             | 29 (82.9)              | 16 (72.7)         | 8 (80)                      |                   |                     |
| **Activities, n (%)**    |                    |                     |                        |                   |                             |                   |                     |
| No problems              | 17 (26.6)          | 14 (28)             | 6 (17.1)               | 6 (27.3)          | 2 (20)                      |                   |                     |
| Slight-extreme problems  | 29 (45.3)          | 29 (56.9)           | 17 (48.6)              | 9 (40.9)          | 6 (60)                      |                   |                     |
| **Pain/discomfort, n (%)** |                  |                     |                        |                   |                             |                   |                     |
| No problems              | 35 (54.7)          | 22 (43.1)           | 18 (51.4)              | 13 (59.1)         | 4 (40)                      |                   |                     |
| Slight-extreme problems  | 33 (51.6)          | 23 (45.1)           | 16 (45.7)              | 9 (40.9)          | 3 (30)                      |                   |                     |
| **Anxiety/depression, n (%)** |              |                     |                        |                   |                             |                   |                     |
| No problems              | 31 (48.4)          | 28 (54.9)           | 19 (54.3)              | 13 (59.1)         | 7 (70)                      |                   |                     |
| Slight-extreme problems  | 39 (60.9)          | 35 (68.6)           | 25 (71.4)              | 13 (59.1)         | 5 (50)                      |                   |                     |
| Slight-extreme problems  | 25 (39.1)          | 16 (31.4)           | 10 (28.6)              | 9 (40.9)          | 5 (50)                      |                   |                     |
| Mobility*                |                    |                     |                        |                   |                             |                   |                     |
| Self-care*               |                    |                     |                        |                   |                             |                   |                     |
| Activities*              |                    |                     |                        |                   |                             |                   |                     |
| Pain/discomfort*        |                    |                     |                        |                   |                             |                   |                     |
| Anxiety/depression*     |                    |                     |                        |                   |                             |                   |                     |

Mobility* p = 0.1957
Self-care* p = 0.7186
Activities* p = 0.4568
Pain/discomfort* p = 0.4575
Anxiety/depression* p = 0.2751

Quality of life K-BILD

|                          | Baseline (Visit 1) | Six weeks (Visit 2) | Three months (Visit 3) | One year (Visit 4) | One year and a half (Visit 5) | Two years (Visit 6) | Main effect of time |
|--------------------------|--------------------|---------------------|------------------------|-------------------|-----------------------------|-------------------|---------------------|
| **Total score**          |                    |                     |                        |                   |                             |                   |                     |
| Mean (SD)                | 57.1 (12)          | 60.7 (10.4)         | 61.9 (13)              | 60 (12.3)         | 62.5 (11.4)                 |                   |                     |
| Range                    | 32–100             | 35.5–84.6           | 36.5–90.8              | 36.5–90.8         | 47.8–84.6                   |                   |                     |
| **Breathlessness/activities** |                  |                     |                        |                   |                             |                   |                     |
| Mean (SD)                | 48.4 (19.8)        | 50.5 (20.2)         | 50.9 (17.9)            | 49.3 (17.6)       | 55.7 (13.5)                 |                   |                     |
| Range                    | 0–100              | 0–100               | 0–79.9                 | 0–79.9            | 39.9–79.9                   |                   |                     |
| **Psychological**        |                    |                     |                        |                   |                             |                   |                     |
| Mean (SD)                | 55.1 (14.7)        | 60.2 (15)           | 63.7 (18.4)            | 59 (17.6)         | 61.6 (18.2)                 |                   |                     |
| Range                    | 28–100             | 32.3–100            | 33.9–100               | 25.3–100          | 41.2–100                    |                   |                     |
| **Chest symptoms**       |                    |                     |                        |                   |                             |                   |                     |
| Mean (SD)                | 73.8 (20.4)        | 77.6 (18)           | 75.2 (20)              | 72.9 (16.3)       | 79.8 (18.1)                 |                   |                     |
| Range                    | 17.3–100           | 17.3–100            | 32.1 (100)             | 32.1–100          | 44–100                      |                   |                     |
| Estimated mean (95% CI)  |                    |                     |                        |                   |                             |                   |                     |
| Total score              | 57.5 (54.6;60.4)   | 60.6 (57.7;63.4)    | 59.3 (55.7;62.9)       | 56.8 (53.1;60.5)  | 57.4 (52.9;62)              |                   | p = 0.0397          |
participants discontinuing the study might have influenced our findings. Interestingly, the COVID-19 pandemic did not seem to have inflated anxiety or depression levels. Ample attention to patients’ psychological well-being is needed, given that this might be associated with a poorer HRQoL, respiratory symptoms and nonadherence [32–34].

Given that IPF is a chronic disease, attention should also be given to healthy lifestyle behaviours. A total of 39.4% of our participants had a high BMI reflecting obesity, which is in line with the Belgian population of 65 years or older [35]. Whether BMI is associated with worse outcomes remains the subject of debate, as studies report mixed findings, leaving ample room for further research on IPF patients’ BMI, nutritional status, and dietary habits [36–38].

In our study, approximately 30% of participants showed at-risk alcohol use. This is only 7% in the Belgian population, although the CAGE and not the AUDIT-C was used [35]. Alcohol-related research is an underinvestigated field in IPF, which is surprising, given that at-risk drinking might aggravate the hepatoxicity of antifibrotic drugs and is known to negatively impact health in other disease populations.

Half of our study population was classified as being insufficiently physically active, which is not surprising considering the nature of the disease. However, trying to maintain an active lifestyle is important, as physical

| Table 3 (continued) |
|----------------------|
| Baseline | Six weeks | Three months | One year | One year and a half | Two years | Main effect of time |
| (Visit 1) | (Visit 2) | (Visit 3) | (Visit 4) | (Visit 5) | (Visit 6) | p = 0.2339 |
| Breathlessness/activities | 48.4 (43.6; 53.2) | 49.7 (44.5; 55.1) | 46.3 (40.5; 52) | 43.5 (37.2; 49.8) | 48 (42.5; 54) |
| Psychological | 55.1 (51.6; 58.7) | 60 (55.8; 64.2) | 61.4 (56.3; 66.9) | 56.1 (51.2; 61) | 56.7 (49.6; 64.5) | p = 0.0151 |
| Chest symptoms | 73.8 (68.9; 78.7) | 78.1 (73.4; 82.9) | 73.2 (67.3; 79.1) | 72.1 (66.7; 81.1) | 72.9 (64.2; 81.5) | p = 0.1798 |

Quality of life SGRQ

| N | 39 | 34 | 28 | 20 | 6 |
| Total score | 39 (20.6) | 33.3 (21.1) | 35.3 (17.2) | 35.3 (18.4) | 26.5 (13.6) |
| Range | 0–92.4 | 43–76.3 | 9.6–67.4 | 6.7–85.5 | 10.5–47 |
| Symptom | 42.6 (25.2) | 30.8 (24) | 29.7 (21.7) | 35.8 (23.6) | 22.8 (9.8) |
| Range | 0–97.7 | 0–88 | 0–73 | 2–72.9 | 29–40 |
| Activities | 54.4 (26) | 49.9 (26.7) | 53.3 (23) | 50.5 (23.1) | 45 (20.4) |
| Range | 0–100 | 0–100 | 0–92.5 | 0–100 | 18.1–67.2 |
| Impact | 29.2 (19.3) | 24.4 (19.7) | 25.6 (17.3) | 26.5 (16.9) | 16.7 (12.1) |
| Range | 0–85.3 | 0–70.5 | 0–97.5 | 1.8–82 | 5.5–37.6 |
| Estimated mean (95% CI) | 36.2 (30.4; 42.1) | 33.4 (27.6; 39.2) | 37.8 (31.6; 44.4) | 36.7 (30.2; 43.2) | 35.1 (28.8; 41.4) | p = 0.4953 |
| Symptoms | 37.6 (31.2; 40) | 31 (24.4; 37.3) | 29.4 (22.6; 36.1) | 35.9 (27.9; 43.9) | 35.3 (29; 41.6) | p = 0.0532 |
| Activities | 52.4 (45.8; 59.1) | 52 (44.7; 59.3) | 55.2 (47.9; 62.6) | 53.8 (46.5; 61.1) | 51.9 (43.5; 60.2) | p = 0.8818 |
| Impact | 25.2 (20.3; 30.2) | 23.3 (18.3; 28.3) | 28.8 (22.3; 35.3) | 27 (21; 33) | 33.8 (22.3; 45.3) | p = 0.1039 |

Significant p-values are indicated in bold
N refers to the number of participants who filled in the questionnaire or for whom the variable was applicable
*Variables of which the evolution over time is not assessed
*For the ‘adherence to sunscreen use’ and the ‘EQ-5D’, we only report the main effect of time (p-value) in this table
We refer the readers to Additional file 1: Table S4 for the pairwise comparisons between timepoints
Abbreviations: K-BILD (The King’s Brief Interstitial Lung Disease questionnaire), SGRQ (The St. George’s Respiratory Questionnaire), EQ-5D (EuroQol 5D)
inactivity is known to be associated with a range of negative outcomes, including mortality and cardiovascular risks [39]. Pulmonary rehabilitation programs for IPF patients do exist and have a positive short-term effect on QoL, fatigue and exercise tolerance [40]. However, referral of all patients to such programs is not part of routine practice and patients might face practical challenges to attend programs (e.g., mobility issues, low self-efficacy). Further research is needed on how physical activity in patients with IPF can be improved should rehabilitation programs not be feasible.

**Strengths and limitations**

This study was conducted at a large ILD centre of expertise where information sessions and long-term follow-up consultations are implemented. The study provides unique insights; however, there are some limitations to consider.

First, we did not measure the prevalence of all potential comorbidities. Because comprehensive evidence on non-medical needs was limited, we decided to assess those needs in depth only.

Second, we used validated questionnaires when available, yet comparing our findings with other studies should be performed cautiously, given that often different instruments were often used.

Additionally, due to the COVID-19 pandemic, we were not able to conduct all study visits face-to-face. The pandemic might have influenced our observations, yet patients did not indicate specific concerns, and our findings that depression and anxiety decreased over time suggest otherwise.

Selection bias might have occurred. However, the sociodemographic characteristics of our sample are comparable to those reported in other IPF studies. Refusal to participate was mainly due to a lack of time or because participants felt too overwhelmed early after diagnosis. Reasons for study discontinuation were mainly due to death or switching to nintedanib.

Regarding the statistical analysis, we realized that we assessed many variables. Given our study’s exploratory nature, no corrections for multiple testing over all these variables were applied. Therefore, caution is warranted when interpreting a single p value. Additionally, due to the small study sample (especially at visit 5 and visit 6) and the high numbers of missing values, we consider the data sparse, which was challenging for binary and ordinal outcomes but nevertheless has high clinical relevance. When the data were too sparse, no formal comparisons were possible for these outcomes. Note that the longitudinal analyses used all available information, i.e., were not restricted to complete cases. Finally, our study contains descriptive data only and was not designed to predict how patients might evolve based on their initial needs profile, yet this could be an interesting area for further research.

**Conclusion**

Conclusively, patients with IPF face issues that go beyond their medical needs. We call for the management of IPF as a chronic disease, thereby focusing on behavioural issues, health literacy and psychological well-being.

**Abbreviations**

AUDIT: Alcohol use disorder identification test; BAASIS: Basel assessment of adherence to immunosuppressive medication scale; BMI: Body mass index; DLco: Diffusion capacity for monoxide; EQ: EuroQol Research Foundation questionnaire; FVC: Forced vital capacity; GAD: Generalized anxiety disorder; HRQoL: Health-related Quality of Life; ILD: Interstitial lung disease; IMAB: Inventory of medication adherence barriers; IPF: Idiopathic pulmonary fibrosis; K-BILD: King’s Brief interstitial lung disease questionnaire; PHQ: Patient health questionnaire; SGRQ: St. George respiratory questionnaire; SHLS: Subjective health literacy screener.

**Supplementary Information**

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**Author contributions**

Credit Author Statement (CRediT). AD: conceptualization, methodology, investigation, formal analysis, writing (original draft). FD: conceptualization, methodology, investigation, validation, writing (review and editing). SF: formal validation, writing (review and editing). KL: investigation, writing (review and editing). SV: investigation, writing (review and editing). WW: conceptualization, validation, writing (review and editing). All authors read and approved the final manuscript.

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**Availability of data and materials**

The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

**Declarations**

**Ethics approval and consent to participate**

The Ethical Committee Research UZ/KU Leuven granted approval (ref. S61427). All participants provided their written informed consent. The study was conducted according to the principles of the Declaration of Helsinki. All data was coded in line with the General Data Protection Regulation (GDPR).

**Consent for publication**

All participants provided their written informed consent.

**Competing interests**

All authors report having no conflict of interest related to this study.
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