Development and validation of the Cystic Fibrosis Decisional Balance for Physical Activity scale (CF-DB-PA)

Valentine Filleul1*, Raphaëlle Ladune1, Mathieu Gruet2, Charlène Falzon1, Amélie Fuchs3, Laurent Mély4, Meggy Hayotte1, Jean-Marc Vallier2, Philippe Giovannetti5, Sophie Ramel6, Anne Vuillemin1, Karine Corrion1 and Fabienne d’Arripe-Longueville1

Abstract

Background: People with cystic fibrosis (pwCF) derive several physiological and psychological benefits from regular physical activity (PA), but the practice is lower than recommended. Knowledge about the facilitators of and barriers to PA at the individual level is important to act positively on PA behaviors. This study validated the Cystic Fibrosis Decisional Balance for Physical Activity scale (CF-DB-PA) for adults with CF.

Methods: French adults with CF were recruited in several specialist centres in France. The CF-DB-PA scale was validated following a quantitative study protocol comprising four stages: (1) tests of the clarity and relevance of a preliminary 44-item version and reduction analysis, (2) confirmatory factor analysis and tests of dimensionality through equation modelling analysis, (3) tests of reliability with Cronbach alphas for the internal consistency and a test–retest with a 2-to-3 week interval for temporal stability, and 4) tests of construct validity with Spearman correlations to measure the associations between each subscale and the theoretically related constructs (i.e., quality of life, PA and exercise tolerance).

Results: A total of 201 French adults with CF participated in the validation study. The CF-DB-PA comprises 23 items divided into two factors: facilitators of and barriers to PA. Each factor is divided into three subscales: physical, psychological and environmental. The factors (facilitators and barriers) can be used independently or combined as a whole. A general score of decisional balance for PA can also be calculated. The bi-factor model presented satisfactory adjustment indexes: χ² (194) = 362.33; p < .001; TLI = .87; CFI = .90; RMSEA = .067. The scale showed satisfactory internal consistency (Cronbach’s α = .77). The test–retest reliability was not significant for either subscale, indicating stability over time. The facilitators subscale correlated significantly with the self-reported score of PA (r = .33, p < .01) and quality of life (r = .24, p < .05). The barriers subscale correlated significantly with the self-reported scores of PA (r = −.42, p < .01), quality of life (r = −.44, p < .01), exercise tolerance (r = −.34, p < .01) and spirometry tests (r = −.30, p < .05).

Conclusions: The CF-DB-PA is a reliable and valid questionnaire assessing the decisional balance for PA, the facilitators of and the barriers to PA for adults with CF in French-speaking samples.

Keywords: Cystic Fibrosis, Physical activity, Exercise, Decisional balance, Barriers and facilitators, Questionnaire

Introduction

Cystic Fibrosis (CF) is a recessive genetic disease that is life-shortening and affects multiple organs of the body. Currently, there is no cure, but advances in treatment...
offer pwCF greater life expectancy than in previous generations [1]. However, the treatments are complex and time consuming, involving daily medications, physiotherapy such as airway clearance, a strict diet, and physical activity (PA) [2]. PA, including sports, exercise, and recreational activities, is widely recommended as part of CF therapy due to its beneficial effects on physiological factors (e.g., improved anaerobic capacity) [3, 4], cardiovascular endurance [5], muscular strength [6], mucus clearance [7], psychological health (i.e., related to quality of life) [4], fatigue [8], and well-being [9].

Although the benefits of PA have been widely demonstrated, it appears that pwCF remain below the recommendations, with levels declining further throughout adolescence [10]. The literature reveals extensive explorations of the facilitators of and barriers to PA in children with CF, although only a few studies have focused on adults. However, with treatments improving, and according to recent annual data from the Cystic Fibrosis Foundation, more adults than children currently have CF, and a child born with CF today has a life expectancy of more than 45 years [11]. From a demographic perspective, the CF population is therefore older and is showing new comorbidities and specific adult constraints (e.g., more pwCF entering the workforce; a high prevalence of diabetes, which is not the case in children and could be a barrier to PA). For these reasons, a specific assessment of the facilitators of and barriers to PA in adults with CF is necessary. Also, the studies in the literature have only identified the facilitators and barriers at the group level in children (e.g., [12–14]), but no tool exists to identify them at the individual level in adults. Yet, clinically, this is essential for two main reasons: (a) time, as these people already have a heavy treatment load and thus may not be readily available for qualitative interviews, which can be lengthy and require expertise, and (b) the need for individualized care, as, given the wide genotypic and phenotypic variability in CF, pwCF are all very different and thus the facilitators of and barriers to PA are likely to vary from one individual to another.

In children with CF, fatigue, negative perceptions of PA, lack of motivation for and interest in PA, a perceived lack of PA, and lack of time, social support or available infrastructures appear to be the main barriers reported [10]. Concerning the facilitators in these patients, the same team of researchers identified the following: improved respiratory capacity and general health, positive perceptions of PA and peer and family social support [10]. A preliminary qualitative study conducted on adults with CF [15], highlighted relationships between physical barriers (i.e., fatigue, respiratory difficulties) and psychological barriers (i.e., lack of perceived physical ability; perceived risk of contamination), which were accentuated by environmental barriers (i.e., lack of time or social support). This study also identified: (a) physical benefits (e.g., improved respiratory capacity), (b) psychological benefits (e.g., well-being), and (c) social support from the environment and time-saving as the main facilitators of PA in adults with CF [15].

In order to successfully change the PA behavior of pwCF, it is necessary to identify and target its determinants. The Transtheoretical Model (TTM), initially created by Prochaska [16], is an integrative model conceptualizing intentional behavior change. The TTM is based on stages that explain when and how people might change their behavior. The Decisional Balance (DB), a key component of the TTM, reflects the relative weight of the pros and cons of changing for an individual. In order to assess the facilitators of (pros) and barriers to (cons) PA, Marcus et al. [17] developed the Decisional Balance Scale for Exercise (DBSE), which was later adapted in French by Eeckhout et al. [18]. This scale was developed for the general population and therefore was not designed to capture the specificities of vulnerable populations, especially the facilitators of and barriers to PA for adults with CF.

Therefore, the purpose of this study was to develop and to validate a DB scale to be used as a PA tool for adults with CF in French-speaking samples. This Cystic Fibrosis Decisional Balance for Physical Activity scale (CF-DB-PA) improves the assessment of the barriers to and facilitators of PA in adults with CF and, consequently, enhances the quality of the support than can be offered to help them adopt an active lifestyle.

Materials and methods
Procedure and participants
We developed and validated the scale through successive steps according to contemporary methodological recommendations [19, 20]: tests for clarity and relevance, dimensionality, reliability tests, and construct validity. Participants were recruited in several specialized CF centers in France. They were all adults with CF (age >18 years). The exclusion criteria were people: (a) under guardianship or trusteeship; (b) in acute exacerbation phase; and (c) with cognitive impairment. The participants’ sociodemographic data are presented in Table 1. They were divided into four samples for the different steps of validation. The questionnaire was administered in paper form with relay persons (nurses and trainees) or online using LimeSurvey CE, version 2.06 + (LimeSurvey CE). This study was approved by the ethics committee from the French National Commission for Information Technology and Civil Liberties (CNIL T 39–2017) and all participants gave their informed consent before
participation. All methods were carried out in accordance with relevant guidelines and regulations.

Measures

**CF-DB-PA**

Items were developed based on existing scales in the general population (e.g., the Decisional Balance Scale for PA) [17], or other vulnerable populations (e.g., the Cancer Exercise Stereotypes Scale) [21] and on the results of a qualitative study of our research team [15]. The Wideband Delphi method, which entails successive rounds of expert input, was employed until consensus was reached [22]. A panel of nine experts was constituted (i.e., 5 researchers in Sport Sciences and Social Psychology, 1 nurse working with pwCF, 1 student in a Master’s program). The experts were identified according to (a) their previous research skills related to exercise in pwCF or to psychological factors of engagement in physical activity in vulnerable populations (5 researchers; 1 master student); (b) recommendations of a physician specialized in CF and part of our research team (2 physicians, 1 nurse). Initially, a list of 69 items was generated, including 40 items related to facilitators and 29 items related to barriers. After two rounds of the Delphi method, several items were deleted due to redundant or ambiguous features to CF. A preliminary version of the CF-DB-PA with 44 items was formulated, including 23 items related to facilitators and 21 items related to barriers. Items were divided into three subscales of facilitators: (a) physical (PHYF, n = 8), (b) psychological (PSYF, n = 5), (c) environmental (ENVF, n = 10); and three subscales of barriers: (a) physical (PHYB, n = 4), (b) psychological (PSYB, n = 7) and (c) environmental (ENVB, n = 10). Participants of sample 1 (see Table 1) were asked to answer questions on a 6-point Likert scale ranging from (1) "Do not at all agree" to (6) "Totally agree".

### Quality of Life

Quality of life (QOL) was assessed using the Cystic Fibrosis Questionnaire for patients over 14 years of age (CFQ14+) [23]. This 49-item questionnaire is composed of nine QOL dimensions: physical functioning, energy/well-being, emotions, social limitations, role, embarrassment, body image, eating disturbances, and treatment burden. QOL was expected to be related to the CF-DB-PA dimensions in the convergent validity step.

### Table 1

Sociodemographic variables and clinical characteristics of study subjects. Descriptive statistics for each sample

| Variable                             | Sample 1   | Sample 2   | Sample 3   | Sample 4   |
|--------------------------------------|------------|------------|------------|------------|
|                                     | n = 9      | n = 192    | n = 73     | n = 53     |
| Age (years)                          | –          | 33.0 (10.5)| 30.1 (9.2) | 32 (9.7)   |
| Age at diagnosis (years)             | –          | 6.1 (12.3) | 3.1 (5.9)  | 3.5 (6.8)  |
| Sex                                  | –          | 111 (57.8) | 38 (52.1)  | 28 (52.8)  |
| Female                               | 5 (55.6)   | 81 (42.2)  | 35 (47.9)  | 25 (47.2)  |
| Male                                 | 4 (44.4)   | 117 (60.9) | 40 (54.8)  | 25 (47.2)  |
| Further education* n (%)             | –          | 153 (79.7) | 52 (71.2)  | 40 (75.5)  |
| Professional status n (%)            | –          | 117 (60.9) | 40 (54.8)  | 25 (47.2)  |
| Workers                              | –          | 4 (2.1)    | 0 (0)      | 0 (0)      |
| Pensioners                           | –          | 71 (37.0)  | 33 (45.2)  | 28 (52.8)  |
| Other people without activity and students | –      | 71 (37.0)  | 33 (45.2)  | 28 (52.8)  |
| Weight (kg)                          | –          | –          | 57.3 (11.6)| –          |
| Height (m)                           | –          | –          | 164.9 (9.1)| –          |
| Body Mass Index (kg/m²)              | –          | –          | 20.9 (3.2) | –          |
| Related disease (n, %)               | –          | 52 (27.1)  | 25 (34.2)  | 20 (37.7)  |
|                                      | –          | 30 (15.6)  | 10 (13.7)  | 6 (11.3)   |
| Distance 6MWT (meters)               | –          | –          | 607.2 (114.7)| –          |
| FEV1 (%)                             | –          | –          | 60.8 (23.7)| –          |
| FEV1 (L)                             | –          | –          | 2.1 (1.0)  | –          |
| FVC (%)                              | –          | –          | 83.6 (21.2)| –          |

* Including university or high school
Level of PA
Self-reported level of PA was assessed with the French version [24] of the Baecke Questionnaire [25], which has been recommended to assess PA in adults with CF [26]. This 16-item instrument with Likert-type responses ranging from 1 for “Never” to 5 for “Always”. The series of questions is divided in three habitual PA scores: occupational; physical exercise in leisure; and leisure and locomotion PA. The PA level was expected to be related to the CF-DB-PA dimensions in the convergent validity step.

Lung function and physical fitness
Forced Vital Capacity (FVC) and forced expiratory volume in one second (FEV₁) were assessed by spirometry testing according to the ATS/ERS TASK FORCE recommendations [27]. Physical fitness was assessed with the six minute Walk Test (6MWT) [28]. The 6MWT was performed according to the recommendations of the ATS guideline [29]. Participants were instructed to walk back and forth in a 30-m length for 6 min. The total covered distance was recorded, rounded off to the nearest meter. These measures were expected to be related to the CF-DB-PA dimensions in the convergent validity step.

Sociodemographic
Sociodemographic information was requested of all participants at the end of the questionnaire and included gender, age, year of diagnosis, related diseases, education level and professional status. Also, lung function and physical fitness were evaluated.

Statistical analyses
All statistical analyses were performed with SPSS (IBM Corporation, version 25) and AMOS (IBM Corporation, version 25) softwares.

Assessment of clarity and relevance
Clarity and relevance were assessed by a large panel of CF professionals and specialists (n = 10; 1 physician, 5 nurses, 1 psychologist, 1 dietician, 1 physical therapist and 1 adapted PA professor) and pwCF (n = 9). They answered on a 6-point Likert scale, ranging from (1) “Totally unclear” to (6) “Totally clear” for the clarity test, and from (1) “Completely irrelevant” to (6) “Highly relevant” for the relevance test, and could comment on their answers. Problematic items were discussed between the members of the expert panel and modified until satisfactory scores were obtained.

Tests of dimensionality
To test the dimensionality of the scale, we ran several structural equation modeling analyses [30]. Based in the recommendations [31], the fit indexes were: chi-square (χ²; significant values p ≤ 0.05), χ² over degrees of liberty (significant values ≤ 3.00), Comparative Fit Index (CFI; value > 0.90), Tucker-Lewis fit index (TLI; value > 0.90), the root mean square error of approximation (RMSEA; value < 0.08) and the 90% confidence interval of RMSEA (ranging from 0.00 to 0.08).

Tests of reliability
Internal consistency of each subscale was assessed using Cronbach’s alpha; a value > 0.70 is considered satisfactory and a value > 0.60 is considered marginally acceptable [32]. The test–retest reliability was calculated twice with a Student t test for paired samples on a reasonable interval of 2 to 3 weeks [33] and a minimum sample size of 50, as recommended [34]. Differences for t were considered statistically significant if p < 0.05.

Tests of convergent validity
Spearman correlation coefficients were used to measure associations between the subscales of the CF-DB-PA and theoretically related constructs (i.e., QOL; PA and exercise tolerance).

Results
Study population
To conduct the successive stages of validation, we divided the participants into four samples: samples 1 (n = 9), 2 (n = 192), 3 (n = 73) and 4 (n = 53). Sample 1 was an independent group. Sample 3 was a subgroup of sample 2 and sample 4 was a subgroup of samples 2 and 3. Assessments of clarity and relevance were conducted on sample 1, tests of dimensionality were conducted on sample 2, test re-test reliability was conducted on sample 4, and the tests of construct validity were conducted on sample 3.

The global sample included 201 participants from ten CF centers of France (i.e., Nice, Giens, Roscoff, Montpellier, Grenoble, Marseille, Caen, Tours, Toulouse and Dunkerque).

Tests of clarity and relevance
The clarity and relevance assessments of the preliminary 44-item version of the CF-DB-PA revealed an acceptable clarity score (M = 5.60, SD = 0.26) and an acceptable relevance score (M = 4.59, SD = 0.66) on a 6-point Likert scale. Three items obtained relevance scores below 3.00 and were removed. Six items with an ambiguous meaning were reworded. Then, redundant items were eliminated after discussion within the expert panel (n = 9) and after the item reduction analysis. Items with the higher clarity and relevance scores were saved, resulting in a new version with 23 items (see Table 2).
Tests of dimensionality

The first maximum likelihood confirmatory factor analysis (CFA) was conducted with the 23-item and six-factor model; four models were examined to assess the dimensionality of the scale [30]. The results of the model fit indexes are presented in Table 3.

Initially, the unidimensional model (model a) did not show satisfactory adjustment indexes. The following analysis examined a first-order model with two factors correlated (facilitators and barriers; model b) and a second-order hierarchical model (model c). At that time, adjustment indexes were not all satisfactory. Finally, analysis using a bi-factor model (model d) offered the most satisfactory adjustment indexes: χ² (194) = 362.33; p < 0.001; TLI = 0.87; CFI = 0.90; RMSEA = 0.067; RMSEA LO/HI = 0.06/0.08 (Fig. 1).

### Table 2 Items of the 23-item version of the CF-DB-PA

| Category | Items |
|----------|-------|
| PHYF | Cela développe mes muscles respiratoires et réduit mon essoufflement. [It develops my respiratory muscles and reduces my shortness of breath.] |
| PHYF | Cela améliore mon endurance. [It improves my endurance.] |
| PHYF | Cela améliore ma force et ma masse musculaire. [It improves my strength and my muscle mass.] |
| PHYF | Une bonne condition physique favorise la réussite de la greffe. [A good physical condition promotes transplant success.] |
| PSYF | C'est l'occasion de penser à autre chose. [This is an opportunity to think about something else.] |
| PSYF | Je me fais plaisir. [I am pleased to do it.] |
| PSYF | Je me permet de rencontrer d'autres personnes. [It allows me to meet other people.] |
| ENVF | Je bénéficie d'un encadrement compétent pour ma pratique. [I benefit from competent supervision for my PA.] |
| ENVF | Je bénéficie de lieux adaptés à ma pratique. [I benefit from adapted places to do my PA.] |
| ENVF | J'ai une offre de pratique près de chez moi. [I have a PA offer in my immediate area.] |
| ENVF | Cela me fatigue trop. [It fatigues me too much.] |
| ENVF | Je supporte mal l'effort physique. [I have trouble tolerating physical effort.] |
| ENVF | J'ai peur d'être trop essoufflé.e. [I worry about getting short of breath.] |
| ENVF | Je n'ai pas l'encadrement adapté à mes besoins. [I don't have supervision that is adapted to my needs.] |
| PSYCH | Je ne pense pas en étant capable physiquement. [I don't think I am physically able to do it.] |
| PSYCH | Je n'arrive pas à suivre le rythme. [I can't follow the rhythm.] |
| PSYCH | Je n'ai pas le temps à cause de mes contraintes familiales. [I don't have time because of my family obligations.] |
| PSYCH | Je n'ai pas l'offre qui me convienne près de chez moi. [I don't have a PA offer that works for me in my immediate area.] |
| PSYCH | Je n'ai pas le temps à cause de mes contraintes familiales. [I don't have time because of my family obligations.] |
| PSYCH | Je n'ai pas l'encadrement adapté à mes besoins. [I don't have supervision that is adapted to my needs.] |

### Table 3 Fit indexes of the different models of the CFA (N= 192)

| Model | χ² (df) | p | RMSEA | CI of RMSEA 90% | TLI | CFI | Δχ²(df) | Δp |
|-------|---------|---|-------|-----------------|-----|-----|---------|-----|
| Model a | 955.89 (223) | < .001 | .13 | .12–.14 | 52 | .57 |
| Model b | 715.44 (222) | < .001 | .11 | .10–.12 | 67 | .71 | 240.45 (1) | < .001 |
| Model c | 544.41 (227) | < .001 | .09 | .08–.10 | 80 | .82 | 171.03 (5) | < .001 |
| Model d | 362.33 (194) | < .001 | .07 | .06–.08 | 87 | .90 | 182.08 (33) | < .001 |

Model a: one-dimensional; Model b: first-order two-factor correlated; Model c: second-order hierarchical order; Model d: confirmatory bi-factor. χ²: Chi²; df: degrees of freedom; RMSEA: root mean square error of approximation; 90% CI: confidence interval of RMSEA 90; CFI: comparative fit index; TLI: Tucker-Lewis index of adjustment.
Tests of reliability
Cronbach alphas ranged from 0.65 to 0.88 (n = 192; i.e., $\alpha_{\text{Facilitators}} = 0.79$; $\alpha_{\text{PHYF}} = 0.70$; $\alpha_{\text{PSYF}} = 0.74$; $\alpha_{\text{ENVF}} = 0.74$; $\alpha_{\text{Barriers}} = 0.88$; $\alpha_{\text{PHYB}} = 0.78$; $\alpha_{\text{PSYB}} = 0.82$; $\alpha_{\text{ENVB}} = 0.65$) and 0.77 for the overall scale, demonstrating satisfactory internal consistency.

The test–retest reliability was not significant for either the barrier or facilitator scales, indicating stability over time for the questionnaire as a whole. At the subscale level, the Student t test was not significant for any subscales except the ENVF subscale. Table 4 presents the results of the t tests.

Tests of construct validity
Correlation analyses showed significant relationships in line with our expectations. Significant positive relationships were observed between the facilitators subscale and (a) the self-reported score of PA ($r = 0.33, p < 0.01$), and (b) the CFQ14+$ (r = 0.24, p < 0.05). The barriers subscale was negatively related to (a) self-reported score of PA ($r = -0.42, p < 0.01$), (b) quality of life ($r = -0.44, p < 0.01$), (c) exercise tolerance ($r = -0.34, p < 0.01$) and (d) spirometry tests ($r = -0.30, p < 0.05$).
### Table 4: Descriptive statistics for the test–retest reliability (n = 53)

| Time 1 M (SD) | Time 2 M (SD) | t tests |
|--------------|--------------|---------|
| Facilitators | PHYF | PSYF | ENVF | Barriers | PHYB | PSYB | ENVB |
| 4.71 (80) | 5.23 (61) | 4.70 (122) | 4.21 (1.13) | 2.76 (90) | 3.06 (1.17) | 2.63 (1.15) | 2.58 (1.16) |
| .76 | .60 | .74 | .42 | .86 | .78 | .83 | .64 |
| 4.83 (91) | 5.23 (61) | 4.78 (1.28) | 4.48 (1.28) | 2.68 (96) | 2.91 (1.12) | 2.58 (1.14) | 2.55 (1.19) |
| .83 | .61 | .72 | .70 | .88 | .76 | .82 | .62 |
| t(52) = −1.70, p = .09 | t(52) = −.74, p = .46 | t(52) = −.2.2, p = .03 | t(52) = −2.2, p = .03 | t(52) = −1.18, p = .24 | t(52) = −1.75, p = .09 | t(52) = 48, p = .63 | t(52) = 31, p = .76 |

PHYF: physical facilitator; PSYF: psychological facilitator; ENVF: environmental facilitator; PHYB: physical barrier; PSYB: psychological barrier; ENVB: environmental barrier

### Discussion

The purpose of this study was to develop and validate a scale to measure the Decisional Balance of PA tailored for a French-speaking adult CF population. Most of the studies focused on PA for pwCF have been conducted in children, and thus knowledge in adults remains scarce. Also, from a clinical point of view, the need for a fast and easy tool to assess PA modifiers at an individual level in adults with CF emerged in parallel to the evolution of the CF population (e.g., longer life expectancy, bigger proportion of adults). The objective of this study was to fill this void. The development of the CF-DB-PA respected the stages of Vallerand’s [20] and Boateng’s [19] methodologies. The preliminary version was based on the findings of a qualitative study that identified the specific characteristics of facilitators of and barriers to PA in adults with CF [15, 16], and the existing scales on the Decisional Balance for PA in the general population [18] and other vulnerable populations [21]. The CF-DB-PA comprises 23 items divided into six subscales: physical facilitators (4 items), psychological facilitators (3 items), environmental facilitators (3 items), physical barriers (4 items), psychological barriers (6 items), and environmental barriers (3 items). The dimensionality tests showed that the bi-factor confirmatory model had the best fit indexes. Thus, the three subscales of facilitators and the three subscales of barriers can be used independently or as a whole. In accordance with Eeckhout et al. [18], a general score of decisional balance for PA can be calculated.

The CF-DB-PA also demonstrated adequate-to-good internal consistency scores for all factors. There were no significant differences in the Student- $t$ test over a 2- to 3-week interval for any subscale except ENVF. This difference can be explained by a change in the participant situation between T1 and T2 (e.g., in care at the CF center for the first time of measure and at home for the second). Thus, these results demonstrated the overall temporal stability of the CF-DB-PA. As predicted, several subscales of the instrument correlated in the expected direction with psychosocial measures such as quality of life from the CFQ14+ [23], PA levels from the Baecke questionnaire [24, 25] and exercise tolerance with the 6MWT [27] and spirometry tests [28], indicating strong convergent validity. Another strength of this scale validation comes from the representativeness of the sample, with participants recruited from several CF centers in France.

Some limitations must nevertheless be acknowledged. First, as for all rating scale, the self-reported nature of the responses may have been biased due to social desirability [35]. Also, this scale was validated for adults and may not be appropriate for children, younger adolescents, or their parents. Our samples did not allow testing for age and gender invariance. Complementary studies could thus be conducted to investigate invariance of this scale. The development of specific versions for children, adolescents and their parents would also be useful. Although the items are also presented in English in the table, the present validation only concerns the French-speaking samples. Future studies on the English validation of the tool or its translation into other languages would be interesting to internationalize its use and allow comparisons between countries.

### Conclusion

This study developed and validated the CF-DB-PA, which is the first measure assessing decisional balance for PA in adults with CF. The psychometric qualities of this scale were demonstrated. The CF-DB-PA offers new possibilities to better measure facilitators of and barriers to PA in adults with CF. The decisional balance score provides useful information on the patient’s stage of change and should help health professionals support and counsel adults with CF to better engage in PA. This scale may also contribute to the development of studies on the determinants of PA adherence in adults with CF. For future research and clinical practice, it might be useful to validate the scale in other languages and develop a digital version of this scale to obtain results more easily and quickly.

### Abbreviations

ATS: American Thoracic Society; CF: Cystic Fibrosis; CF-DB-PA: Cystic Fibrosis Decisional Balance for Physical Activity scale; CFA: Confirmatory factor analysis; CFI: Comparative Fit Index; CFQ14+: Cystic Fibrosis Questionnaire for patients over 14 years of age; CNIL: Commission for Information Technology and Civil Liberties; DF: Degrees of freedom; ENVB: Environmental barriers; ENVF: Environmental facilitators; FEV1: Forced expiratory volume in one second; FVC: Forced vital capacity; M: Mean; PA: Physical activity; PHYB: Physical barriers; PHYF: Physical facilitators; PSYB: Psychological barriers; PSYF: Psychological facilitators; pwCF: People with Cystic Fibrosis; QOL: Quality of life; SD: Standard
deviation; RMSEA: Root mean square error of approximation; TLI: Tucker–Lewis Index of adjustment; 6MWT: Six minute Walk test; 90% CI: Confidence interval of RMSEA 90, χ2: Chi².

Acknowledgements
We thank all the participants for the time they dedicated to our study. We thank the Association Grégory Lemarchal which has supported this study by a subvention.

Authors’ contributions
VF: provided contributions to the conception and design of the work (including design of the questionnaire), to the acquisition, analysis and interpretation of data. Drafted the work. RL: provided contribution to the acquisition of data and drafted and revised the paper. MG and JMV: provided important contributions to the design of the work and the acquisition of data. Revised the paper critically for important scientific and clinical content: CF and AV: revised the paper.
AF, LM, SR and PG: provided contribution to the acquisition of data and implemented the data collection in their own medical structure; revised the paper.
MH: provided contribution to the analysis of data and revised the paper. KC: provided contribution to the design of the work, was the main responsible for the statistical analysis and thus provided substantial contributions to the conception of the statistical analysis plan, conduct of the analysis and interpretation of results. Revised the paper. FAL: was responsible for the scientific project and oversaw funding obtained, oversaw the collection of data, contributed to the interpretation of the results, and helped write the manuscript. All authors have contributed to the drafting, reviewing and approval of the final manuscript of this protocol paper.

Funding
This work was funded by Association Grégory Lemarchal, France.

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
This study was approved by the ethics committee: the French National Committee for Information Technology and Civil Liberties (CNIL T 39-2017) and all participants gave their informed consent before participation.

Consent for publication
Not applicable.

Competing interests
The authors declare that they have no conflict of interest.

Author details
1 Université Côte-d'Azur, LAMHESS, Nice, France. 2 Université de Toulon, Unité de Recherche Impact de l’Activité Physique sur la Santé, Toulon, France. 3 Laboratoire de thérapeutiques non médicamenteuses innovantes Mooven, Montpellier, France. 4 Hôpital Renée Sabran, Centre de Ressources et de Compétences de la Mucoviscidose (CRCM), Giens, France. 5 Service de Pneumologie, Centre de Ressource et de Compétence de la Mucoviscidose (CRCM) adulte et pôle d’Activités Médicales Intersite de Médecine Physique et de Réadaptation APHM-CHU Hôpital Nord, Marseille, France. 6 Fondation Il dys, site de Perpignan, Roscoff, France.

Received: 15 December 2020 Accepted: 15 March 2021 Published online: 14 April 2021

References
1. Burgel PR, Bellis G, Olesen HV, Viviani L, Zolin A, Blasi F, Elborn JS. Future trends in cystic fibrosis demography in 34 European countries. Eur Respir J. 2015;46(1):133–41.
2. Sawicki GS, Sellers DE, Robinson WM. High treatment burden in adults with cystic fibrosis: challenges to disease self-management. J Cyst Fibros. 2009;8:91–6.
3. Klijn PH, Oudshoorn A, van der Ent CK, van der Net J, Kimpen JL, Holders PJ. Effects of anabolic training in children with cystic fibrosis: a randomised controlled study. Chest. 2004;125(4):1299–305.
4. Radtke T, Nevitt SJ, Hebestreit H, Kriemler S. Physical exercise training for cystic fibrosis. Cochrane Database Syst Rev. 2017;11:CD002768.
5. Guilmans VAM, de Meer K, Brackett HJL, Faber JA, Berger R, Holders PJM (1999) Outpatient exercise training in children with cystic fibrosis: physiological effects, perceived competence, and acceptability. Pediatr Pulmonol. 1999;28(1):39–46.
6. Gruet M, Troosters T, Vergees S. Peripheral muscle abnormalities in cystic fibrosis: etiology, clinical implications and response to therapeutic interventions. J Cyst Fibros. 2017;16:538–52.
7. Owrey TJ, Zainulidin R, Davskas E, Bye PT, Alison JA. Effects of treadmill exercise versus flutter® on respiratory flow and sputum properties in adults with cystic fibrosis: a randomised, controlled, cross-over trial. BMC Pulm Med. 2017;17(1):1–8.
8. Orava C, Fitzgerald J, Figliomeni S, Lam D, Naccarato A, Szego E, et al. Relationship between physical activity and fatigue in adults with cystic fibrosis. Physiother Can. 2018;70(1):42–8.
9. Backstrom-Eriksson L, Bergsten-Brueckers A, Hjelte L, Melin B, Sorjonen K. Associations between genetics, medical status, physical exercise and psychological well-being in adults with cystic fibrosis. BMJ Open Resp Res. 2016;3(1):e000141.
10. Jantzen A, Opoku-Pare M, Ruf K, Hebestreit H, Möller A. (2014). Cystic fibrosis and physical activity: Is there a significant difference to healthy individuals?. Eur Respir J 2014;44(Suppl 58).
11. Cystic Fibrosis Foundation Patient Registry. 2018 Annual Data Report Bethesda, Maryland 2019 Cystic Fibrosis Foundation.
12. Swisher AK, Erickson M. Perceptions of physical activity in a group of adolescents with cystic fibrosis. Cardiopulm Phys Ther J. 2008;19:107–13.
13. Moola FJ, Faulkner GE, Schneiderman JE. “No time to play”: perceptions toward physical activity in youth with cystic fibrosis. Adapt Phys Activ Q. 2012;29:44–62.
14. Happ MB, Hoffman LA, Higgins LW, et al. Parent and child perceptions of a self-regulated, home-based exercise program for children with cystic fibrosis. Nurs Res. 2013;62:305–14.
15. Filleul V, Falzon C, Fuchs A, Ladune R, Mely L, Vallier JM, d’Arripe-Longueville F. L’activité physique adaptée chez les patients atteints de mucoviscidose : étude qualitative des barrières et facilitateurs. Sci Sports. 2018;33:529.
16. Prochaska JO. Systems of psychotherapy: a transtheoretical analysis. Homewood, III. Dorsey Press 1979.
17. Marcus BH, Rakowski W, Rossi JS. Assessing motivational readiness and decision making for exercise. Health Psychol. 1992;11(4):257–61.
18. Eckhout C, Francaux M, Heeren A, Philipott P. Mesure de la balance décisionnelle en vue de pratiquer une activité physique régulière (Bdap): Adaptation et validation francophone de l’échelle Decisional Balance for Exercise. Eur Rev Soc Psychol. 2013;15(3):185–91.
19. Falzon C, Sabiston C, Bergamaschi A, Corrion K, Chalabaev A, d’Arripe-Longueville F. Development and validation of the cancer exercise stereo-types scale. J Psychosoc Oncol. 2014;32(6):708–26.
20. Falzon C, Sabiston C, Bergamaschi A, Corrion K, Chalabaev A, d’Arripe-Longueville F. Development and validation of the cancer exercise stereotypes scale. J Psychosoc Oncol. 2014;32(6):708–26.
21. Dalkey NG, Rostand F, Kaufmann A, Brown B, Cochran SW. La prévision à long terme par la méthode Delphi. 1972 Dunod.
26. Karila C, Ravilly S, Gauthier R, Tardif C, Neveu H, Maire J, et al. Activité physique et réentraînement à l’effort du patient atteint de mucoviscidose. Rev Mal Respir. 2010;27(4):301–13.
27. Miller MR, Hankinson J, Brusasco V, Burgos F, Casaburi R, Coates A, et al. ATS/ERS Task Force. Standardisation of spirometry. Eur Respir J. 2005;26(2):319–38.
28. Guyatt GH, Sullivan MJ, Thompson PJ, Fallen EL, Pugsley SO, Taylor DW, Berman LB. The 6-minute walk: a new measure of exercise capacity in patients with chronic heart failure. Can Med Assoc J. 1985;132(8):919–23.
29. Statement ATS. Guidelines for the six-minute walk test. Am J Respir Crit Care Med. 2002;166(1):111–7.
30. Myers ND. Coaching competency and (Exploratory) structural equation modeling: a substantive-methodological synergy. Psychol Sport Exerc. 2013;14(5):709–18.
31. Byrne BM. Structural equation modeling with mplus: basic concepts, applications, and programming. 2nd ed. New York: Routledge, Taylor & Francis Group, 2013.
32. Taber KS. The use of cronbach’s alpha when developing and reporting research instruments in science education. Res Sci Technol Educ. 2018;48(6):1273–96.
33. Marx RG, Menezes A, Horovitz L, Jones EC, Warren RF. A comparison of two time intervals for test-retest reliability of health status instruments. J Clin Epidemiol. 2003;56(8):730–5.
34. Altman DG. Practical statistics for medical research. 1st ed. London: Chapman & Hall/CRC Press; 1990. p. 455–60.
35. Phillips DL, Clancy KJ. Some effects of social desirability in survey studies. Am J Sociol. 1972;77(5):921–40.

Publisher’s Note
Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.