Correlation of Gut Microbiota, Vitamin D Status, and Pulmonary Function Tests in Children With Cystic Fibrosis

Hadeel Albedewi¹, Iman Bindayel¹*, Ahmed Albarrag² and Hanaa Banjar³

¹ Department of Community Health Sciences, College of Applied Medical Sciences, King Saud University, Riyadh, Saudi Arabia, ² Department of Pathology, College of Medicine, King Saud University, Riyadh, Saudi Arabia, ³ Department of Pediatrics, King Faisal Specialist Hospital and Research Center, Riyadh, Saudi Arabia

Background: Children with cystic fibrosis (CF) are expected to have suboptimal serum vitamin D status and altered gut microbiota. The altered gut microbiota is hypothesized to have a pro-inflammatory effect that further complicates the existing respiratory inflammation. Emerging evidence suggests an association between vitamin D and gut microbiota. The aim of this study was to assess the relationships between 25-hydroxyvitamin D [25(OH)D] status, pulmonary function, and fecal bacteria in children with CF.

Methods: In this cross-sectional study, a total of 35 children with CF (8.7 ± 2.83 years) and 24 controls without CF (9 ± 2.7 years) were included in this study. Serum 25(OH)D status was measured using the Elecsys vitamin D total II assay. In the CF group, gut microbiota composition was assessed using real-time PCR analysis. Pulmonary function tests (PFTs) were measured using spirometry. Comparisons between the CF and non-CF controls were conducted using the independent sample t-test. In the CF group, one-way analysis of variance (ANOVA) was used to assess differences in PFTs and gut microbiota composition across the three vitamin D subgroups. The correlations between 25(OH)D status and PFTs, or gut microbiota composition, and PFTs with gut microbiota composition were analyzed using the Pearson's correlation coefficient test.

Results: Children with CF had significantly lower serum 25(OH)D levels compared with children without CF (44.3 ± 22.4 vs. 59 ± 25.5, respectively, \(P = 0.026\)). Children with CF with optimal serum 25(OH)D level had significantly higher levels of Bacteroidetes, Firmicutes, and total bacteria (\(P = 0.007\), \(P = 0.007\), and \(P = 0.022\), respectively). The level of Firmicutes was found to be significantly higher in mild forced expiratory volume in 1 s (FEV1) compared with moderate FEV1 (\(P = 0.032\)), whereas the level of the other bacteria species was comparable across FEV1 severity groups.

Conclusion: Our findings may encourage studies that target and modify gut microbiota to potentially achieve better outcomes in terms of respiratory function in CF.

Keywords: cystic fibrosis, 25-hydroxyvitamin D, gut microbiota, real-time PCR, pulmonary function tests
INTRODUCTION

Cystic fibrosis (CF) is an autosomal recessive disorder caused by a mutation in CF transmembrane conductance regulator (CFTR) gene (1). There are several manifestations of CF, including chronic lung inflammation and repeated infections, and gastrointestinal abnormalities such as exocrine pancreatic insufficiency and fat malabsorption that leads to malnutrition. These characteristics of the disease impose gastrointestinal microbiota alteration and vitamin D deficiency or insufficiency that is often observed in individuals with CF (2). Vitamin D deficiency or insufficiency is common in patients with CF due to pancreatic insufficiency, fat malabsorption, low vitamin D binding protein level, corticosteroid medication use, and inadequate exposure to sunlight (3, 4). In addition to maintaining skeletal strength and bone mineralization, vitamin D has the ability to decrease colonization of the airways through stimulating antimicrobial peptides LL-37 that promote bronchial cell reaction to Pseudomonas aeruginosa, a pathogen related to CF (5). Furthermore, vitamin D was found to be an inhibitor of airway pro-inflammatory cytokines indicated by a significant reduction in interleukin 17A and interleukin 23 levels after vitamin D supplementation (6). It has also been observed that polymorphisms in vitamin D binding protein correlate with 25(OH)D levels and decreased pulmonary function measured by forced expiratory volume in 1 s (FEV1) in patients with CF (7).

Individuals with CF are expected to have gastrointestinal microbiota alteration due to changes in gastrointestinal mucus, pH, fat malabsorption, and antibiotic use (8). Moreover, the severity of CFTR mutation was found to be correlated with gut microbial imbalance, namely, an increase in pathogenic bacteria, such as Escherichia coli (E. coli), and a reduction in Bifidobacteria (9). The E. coli species produces pro-inflammatory mediators that initiate a low-grade inflammation environment, which may trigger systematic inflammation and eventually chronic lung inflammatory disease (8). Furthermore, a study showed a disease-specific pattern of gut microbiota alteration in CF, regardless of patients’ age, antibiotic use, or pancreatic function (10). In fact, the intestine of patients with CF produces suboptimal bicarbonate levels, causing a lower pH level and thicker mucus and resulting in intestinal transit interruption (11). Therefore, the lower secretion of chloride and bicarbonate and higher uptake of water intestinal lumen causes dehydration, increases the lumen content acidity, and eventually compromises nutrient absorption (2). Individuals with CF manifest an impairment in mucus production and transport, creating a suitable environment for small intestine bacterial overgrowth. This was also related to a reduced diversity in commensal gut microbiota. As a result, the formerly mentioned factors act collectively in the progression of chronic low-grade inflammation and impaired immunity in the gut of individuals with CF (2). In fact, a study conducted on infants with CF has linked the alteration of the gut microbial community in CF to pulmonary exacerbation during the first year of life (12). Furthermore, a longitudinal study on 13 children with CF revealed a significant correlation between gut inhibiting bacteria and respiratory symptoms rather than respiratory inhabiting bacteria, namely; Bacteroides and Bifidobacterium (13). Another longitudinal study conducted on 7 CF neonates has demonstrated a matched variation and richness of several bacterial taxa inhabiting the gut and respiratory tract (14). Several bacterial species that were elevated in the gut were also found to be elevated in the respiratory tract. Interestingly, the gut microbiome colonization was found to precede that of the respiratory tract; for instance, the gut colonization of Enterococcus was followed by a cluster formation of these potentially pathogenic bacteria in the respiratory tract (14). The suggested pathways for the gut-lung axis include endothelium signal absorption by epithelial cells from local to distal sites, mesenteric lymphatic system, and local and long-reaching immune reactions (15, 16). Intestinal epithelial cells are directly stimulated by gut microbiota, which regulates immune cell release. Inflammation is linked to bacterial translocation, which involves the migration of microbes or their metabolites from the gastrointestinal system to the circulation across the mucosal barrier (17).

Recently, emerging evidence suggests an association between vitamin D and the composition of gut microbiota (18). A cross-sectional study has reported a negative correlation of vitamin D status with Coprococcus and Bifidobacteria in a healthy population (r = −0.22 and r = −0.27, P < 0.005, respectively). In addition, the study had shown the subgroup with higher dietary vitamin D intake to have a significantly higher abundance of Prevotella and lower abundance of Haemophilus and Veillonella compared with the lower dietary vitamin D intake subgroups (19). In CF, supplementation of vitamin D3 (50,000 IU per week) for 3 months among 23 adults with CF resulted in a significant increase in Lactococcus and a decrease in Veillonella and Erysipelotrichaceae when compared with placebo (20). The pathways in which vitamin D modulates the gut microbiota composition include reducing intestinal inflammation, decreasing gastrointestinal permeability, increasing intestinal barrier function, and stimulating antimicrobial peptides (18).

Therefore, due to the low number of studies investigating the association between vitamin D and gut microbiota in CF (20, 21), there is a need for more studies that compare vitamin D level in children with CF and children without CF and then examine its association with gut residents and pulmonary function outcomes. Therefore, this study aimed to assess 25-hydroxyvitamin D [25(OH)D] levels in patients with CF and controls without CF, then measure the fecal abundance of five of the most relevant bacteria species in patients with CF, and investigate their association with 25(OH)D status. In addition, relationships between 25(OH)D status and pulmonary function and fecal bacteria and pulmonary function were examined in the children with CF.

Abbreviations: 25(OH)D, 25-hydroxyvitamin D; ANOVA, analysis of variance; BMI, body mass index; CDC, Center for Disease Control; CF, cystic fibrosis; CFTR, CF transmembrane conductance regulator; FEV1, forced expiratory volume in 1 s; FFQ, food frequency questionnaire; FVC, forced vital capacity; KFSH&RC, King Faisal Specialist Hospital and Research Center; PFT, pulmonary function tests; RAC, Research Advisory Committee.
MATERIALS AND METHODS

Participant Recruitment and Ethics Statement

In this cross-sectional study, children with CF who were stable with no acute pulmonary exacerbation, gastrointestinal symptoms, or antibiotic use for at least 2 weeks were recruited from the Cystic Fibrosis Clinic at King Faisal Specialist Hospital and Research Center (KFSH&RC), Riyadh, Saudi Arabia, during their regular clinical visits between July 2019 and December 2019. Moreover, children without CF were recruited from a pediatric clinic and classified as the non-CF control group. Parents were contacted before their appointments by their nurse coordinator to inform them about the study. Patients were screened by their pediatrician for inclusion and exclusion criteria. Anthropometrical measurements were performed by a nurse to assess for growth and malnutrition, and then pulmonary function testing was performed by a respiratory therapist. Blood samples routinely collected were used to assess serum 25(OH)D levels. Parents were then interviewed by the dietician to estimate their dietary intake of vitamin D. The protocol was approved by the institutional review board of KFSH&RC in the Office of Research Affairs at the Research Advisory Committee (RAC) with the project number RAC# 2191068. Written informed consent was obtained from all parents or caregivers of involved study children. All clinical investigations were performed in accordance with the Declaration of Helsinki. The inclusion criteria were as follows: (1) CF diagnosis based on clinical symptoms, high sweat chloride testing (>60 mmol/L), and CFTR mutational analysis, (2) stable condition (assessed by the pediatrician), and (3) age >5 years. The exclusion criteria were as follows: (1) antibiotic use within 3 weeks prior to fecal sample collection and (2) intellectual disability that affects the ability to perform pulmonary function test (PFT) accurately.

Anthropometric Measurements

Anthropometric measurements including weight, height, and body mass index (BMI) were obtained from study subjects during their clinical visits by a registered nurse. BMI z-score and BMI percentile were calculated using Center for Disease Control (CDC) growth charts. The CDC cutoffs for BMI-for-age were used as follows: <5th percentile is considered underweight, 5th to <85th percentile is healthy weight, 85th to <95th percentile is overweight, and >95th percentile is obese (22).

Total 25(OH)D Status

The measurement of total serum 25(OH)D levels was conducted at the laboratory of KFSH&RC using the Elecsys vitamin D total II assay on Cobas 801 immunoassay auto-analyzer system (Roche Diagnostics GmbH, Mannheim, Germany). The cutoff point for vitamin D deficiency and insufficiency was obtained from the Institute of Medicine where a 25(OH)D level of <30 nmol/L is considered vitamin D deficiency, a level between 30 and 50 nmol/L is considered vitamin D insufficiency, and the optimal vitamin D level was set at >50 nmol/L (23).

Dietary Vitamin D Intake

The assessment of dietary vitamin D intake among study subjects was conducted through a validated food frequency questionnaire (FFQ) specific for calcium and vitamin D obtained from a previous study conducted on Saudi children (24). The amounts of vitamin D in each food item included in the questionnaire were adapted from the U.S. Department of Agriculture food composition database (25).

Pulmonary Function Tests

Pulmonary function tests, including FEV1, forced vital capacity (FVC), FEV1/FVC ratio, and forced expiratory flow at 25 and 75% of the pulmonary volume (FEF25−75%), were performed using spirometry. All PFTs were expressed as percentages of predicted values. The cutoff points for the severity of airflow obstruction based on the percentage of the predicted value of FEV1 were as follows: FEV1% predicted >80% was considered as mild lung disease, while FEV1% predicted between 50% and 80% was moderate lung disease, and FEV1% predicted <50% was considered as severe lung disease (26).

Fecal Microbiota Analysis

Collection and Storage

A freshly voided stool sample from each participant in the CF group was collected by their parents or caregivers in sterilized sample containers and kept at 4°C for a maximum of 3 h. Later, two aliquots weighing 250 mg were obtained from each fecal sample and then stored at −80°C for DNA extraction.

DNA Extraction

DNA extraction was performed on a single aliquot for each sample using the commercially available kit (QIAamp DNA stool mini kit; Qiagen, Hilden, Germany) following the manufacturer’s instructions. The concentration and purity of extracted DNA were measured using the Nanodrop 2000 Spectrophotometer (Thermo Fisher Scientific, MA, USA) at 260 nm and 260/280 ratio, respectively. Extracted DNA samples were adjusted to a concentration of 10 ng/µl and stored at −20°C until they are analyzed for bacterial quantification by real-time PCR.

Standard Curve

Positive controls were used to construct a standard curve for total and target bacteria. Bacterial strains used as positive controls are shown in Table 1. Genomic DNA for each bacterial strain was extracted from pure cultures using the QIAamp DNA mini kit (Qiagen, Hilden, Germany) following the manufacturer’s instructions. To generate the standard curves, we used ten-fold serial dilutions of genomic DNA extracted from pure cultures with known concentrations ranging from 1 × 10^8 to 1 × 10^2 colony-forming units per milliliter (CFU/ml) of each reference bacterium. We performed real-time PCR reactions and plotted the CT values against the log concentration of the reference bacterium copy number to create a linear regression.

Real-Time PCR

PCR primers were selected to determine the abundance of the gut microbiota strains of interest (Table 1). Real-time PCR was performed in a 50 µl reaction run in triplicate. Each
The primers and positive control used in this study.

| Primer       | Sequence                              | Target               | Positive control        | References |
|--------------|---------------------------------------|----------------------|-------------------------|------------|
| Bact934F     | GGACATGTGGTTATTCGATGAT                | Bacteroidetes        | B. fragilis             | (27)       |
| Bact1060R    | AGCTGAGCACAACACTGCA                   | Enterobacteriaceae   | E. coli                 | (28)       |
| Eco1457F     | CATTGAGTTACCGCGAAGAAGG               | Bifidobacterium spp. | B. longum               | (29)       |
| Eco1652R     | CTCTACGAGACTCAAGGTGCC                | Firmicutes           | C. perfringens          | (27)       |
| F-bifido     | CGCGTCYGGTGTTAAG                    | Lactobacillus spp.   | L. rhamnosus            | (30)       |
| R-bifido     | CCCACATCGAACATGCA                   | Lactobacillus spp.   | L. rhamnosus            | (30)       |
| Firm934F     | GGAGYATGGTGGTTAATCGAAGGA             | Bacteroidetes        | B. fragilis             | (27)       |
| Firm1060R    | AGCTGAAGCAGACACATGCA                | Enterobacteriaceae   | E. coli                 | (28)       |
| Lacto-F      | ACGATAGGGAATCTTCCCA                  | Firmicutes           | C. perfringens          | (27)       |
| Lacto-R      | CACGCTACACATGGAG                    | Lactobacillus spp.   | L. rhamnosus            | (30)       |
| UnrF         | TCCTACGGGAGCAGCAAGT                  | Total                | L. rhamnosus            | (30)       |
| UnrR         | GGACTACCCAGGATCCTATCTCGTT           |                      |                         |            |

reaction consists of 25 μl QuantiTect SYBR Green PCR Master Mix (Qiagen, Hilden, Germany), 7.5 μl of each forward and reverse primer, and 10 μl of DNA. Amplification was performed using Rotor-Gene Q (Qiagen, Hilden, Germany) starting with denaturation at 95°C for 15 min, followed by 40 cycles of denaturation at 94°C for 15 s, annealing for 30 s, and extension at 72°C for 40 s. The annealing temperature was 61.5°C for E. coli, 59°C for Bifidobacteria, 60.44°C for Firmicutes, and 60°C for Bacteroidetes, Lactobacillus, and total bacteria. The genome copy number of each reaction was calculated using the standard curves. Additionally, to calculate the total genome copy number per gram of wet stool, we used the equation (QM × C × DV)/(S × V) adapted from Metzler-Zebeli et al. (31) where QM is the copy number quantitative mean, C is the sample DNA concentration, DV is the dilution volume of extracted DNA, S is the amount of DNA used in the reaction, and V is the initial sample weight that was used for DNA extraction (31).

Statistical Analysis

All data analyses were performed using Statistical Package for Social Science Statistics IBM SPSS for Macintosh, version 25.0 (IBM Corp., Armonk, N.Y., USA). Descriptive data are expressed as mean and standard deviation. Comparisons between the CF and non-CF control were conducted using the independent sample t-test. In the CF group, one-way analysis of variance (ANOVA) was used to assess differences in PFTs and gut microbiota composition across the three vitamin D subgroups (i.e., deficiency, insufficiency, and optimal). The correlations between 25(OH)D status and PFTs, or gut microbiota composition, and PFTs with gut microbiota composition were analyzed using the Pearson’s correlation coefficient (two-tailed) test. A P-value < 0.05 was considered significant in all tests.

RESULTS

In this cross-sectional study, a total of 35 children (16 males; 19 females, mean age, 8.7 ± 2.83 years) with CF and 24 non-CF control (14 males; 10 females, mean age, 9 ± 2.7 years) were included. The general characteristics of participants are shown in Table 2. The CF group had significantly lower BMI and BMI z-score compared with the non-CF group (P = 0.002, P = 0.016, respectively). In addition, there was a significant difference in regional distribution between the groups, with the majority of the CF children (54%) living in the Eastern region of Saudi Arabia (P = 0.001) (Table 2).

Vitamin D in Serum and Dietary Intake

The total serum 25(OH)D level of CF children was significantly lower compared with non-CF children (independent sample t-test, P = 0.026, Table 2). There was no significant difference in age, sex, and BMI across the vitamin D subgroups in both children groups. The median estimated dietary intake of vitamin D and calcium for the CF group was 60% and 69.8% of the Recommended Dietary Allowance, respectively (Table 2).

Pulmonary Function Tests

The PFTs were not different among the vitamin D subgroups in children with CF. Similarly, there was no significant correlation between PFT and vitamin D serum level. However, there was a significant positive correlation between BMI z-score and each one of the PFTs: FEV1, FVC, and FEF 25–75% (r = 0.515, P = 0.002; r = 0.472, P = 0.005; and r = 0.358, P = 0.041, respectively, data not shown).

Gut Microbiota

The bacterial copy number in feces for target and total bacteria is shown in Figure 1. Data are shown as the log10 of the number of genome copies/g of a fecal sample. Children with CF with optimal serum vitamin D had significantly higher levels of Bacteroidetes, Firmicutes, and total bacteria (P = 0.007, P = 0.007, and P = 0.022, respectively, Figure 2). Correspondingly, Bacteroidetes, Firmicutes, and total bacteria were found to be positively correlated with serum 25(OH)D level (Pearson’s correlation, r = 0.385 P = 0.027; r = 0.461, P = 0.007; and r = 0.410, P = 0.018, respectively, Figure 3). However, there was no difference in Bifidobacteria, Lactobacillus spp., E. coli, and Firmicutes/Bacteroidetes ratio across vitamin D groups in...
TABLE 2 | Study participant characteristics.

| Variable                  | CF N = 35 | Non-CF N = 24 | P-value |
|---------------------------|-----------|---------------|---------|
| Age, years, mean (SD)     | 8.7 (2.83)| 9 (2.7)       | 0.601   |
| Sex (male/female)         | 16/19     | 14/10         | 0.341   |
| Region, n (%)             |           |               | 0.001   |
| Central                   | 7 (20)    | 15 (63)       |         |
| East                      | 19 (54)   | 2 (8)         |         |
| West                      | 1 (3)     | 1 (4)         |         |
| North                     | 5 (14)    | 2 (8)         |         |
| South                     | 3 (9)     | 4 (17)        |         |
| Height, cm, mean (SD)     | 124 (15.57)| 123 (18.3)    | 0.805   |
| Weight, kg, mean (SD)     | 22.9 (7.11)| 27 (11.1)     | 0.107   |
| BMI, kg/m², mean (SD)     | 14.6 (1.4)| 17.3 (3.8)    | 0.002   |
| BMI z-score, mean (SD)    | −1.3 (1.4)| −0.23 (1.8)   | 0.016   |
| BMI-for-age, n (%)        |           |               | 0.148   |
| Underweight               | 13 (38)   | 6 (25)        |         |
| Normal                    | 20 (59)   | 14 (58)       |         |
| Overweight                | 1 (3)     | 4 (17)        |         |
| Serum 25(OH)D (nmol/L), mean (SD) | 44.3 (22.4) | 59 (25.5) | 0.026 |
| Serum 25(OH)D status (n/%) |           |               | 0.083   |
| Deficiency                | 8 (24.2)  | 1 (4)         |         |
| Insufficiency             | 14 (42.4) | 10 (42)       |         |
| Optimal                   | 11 (33.3) | 13 (54)       |         |
| Vitamin D (supplements) (IU/day), mean (SD) | 2,912 (1,083) | – | – |
| Vitamin D dietary intake (FFQ) (IU/day), mean (SD) | 365 (141) | – | – |
| Vitamin D total intake (IU/day), mean (SD) | 3,090 (1,288) | – | – |
| Pulmonary function testing|           |               |         |
| Fev1 (%), mean (SD)       | 71 (27)   | –             | –       |
| FVC (%), mean (SD)        | 72 (27)   | –             | –       |
| FEV1/FVC (%), mean (SD)   | 97 (13)   | –             | –       |
| FEF 25–75 (%), mean (SD)  | 74 (35)   | –             | –       |

The independent sample t-test was used to determine a significant difference in continuous variables between the CF and non-CF groups. Pearson chi-square test was used to determine a significant difference in the categorical variables between the CF and non-CF groups.

25(OH)D, 25-hydroxyvitamin D; SD, standard deviation; CD, cystic fibrosis.

DISCUSSION

This study was conducted to investigate vitamin D hypovitaminosis and its association with PFTs and gut microbiota in children with CF. Our study showed a significantly lower serum 25(OH)D level in children with CF than in children without CF, and this is in accordance with previous observations suggesting a lower serum vitamin D level in children with CF (32). Reasons for the development of suboptimal vitamin D status in children with CF were attributed to the nature of CF disease that results in pancreatic insufficiency, reduced vitamin D binding protein levels, and fat malabsorption, therefore, vitamin D malabsorption (3, 4). Moreover, the clinical management of CF often necessitates the use of corticosteroids that may increase the expression of vitamin D catabolizing enzyme CYP24, in addition to some antibiotics that cause photosensitivity and lead to lower exposure to sunlight (3, 4). The mean BMI-for-age for children with CF in our sample was between the 5th and 15th percentile, with 40% of the sample being at <5th percentile, which indicates that they either are nutritionally at risk or have a nutritional failure (33). Children with CF are known to have lower than normal body weight due to energy loss that is caused by maldigestion and malabsorption, increased energy expenditure, and decreased energy intake, resulting from anorexia and gastrointestinal symptoms (34). In fact, our study has shown significantly lower BMI and BMI z-score in children with CF compared with children without CF.

Previous studies had shown a significant positive association between vitamin D status and PFTs (35–37); however, this study was unable to demonstrate a significant association between 25(OH)D and PFTs. A possible explanation might be the fact that the majority of our sample (69%) had mild to moderate airflow obstruction with no pulmonary exacerbations. In fact, few previous studies were unable to demonstrate any significant association between vitamin D status and PFTs;
FIGURE 2 | Difference in fecal bacteria strains among deficiency, insufficiency, and optimal serum vitamin D in children with CF using ANOVA. (A) Bacteroidetes. (B) Firmicutes. (C) Total bacteria in children with CF. Values are shown as mean and SD of log-transformed genome copies/g of fecal sample. *P < 0.05; **P < 0.005.
FIGURE 3 | Correlation between fecal bacteria strains and serum 25-hydroxyvitamin D level (nmol/L) in children with CF. (A) Bacteroidetes, (B) Firmicutes, (C) Total bacteria. $r$ denotes Pearson’s correlation coefficient.
rather, a significant association with pulmonary exacerbation was demonstrated (38, 39).

It was hypothesized that gut microbiota composition is positively correlated with serum vitamin D status. In this study, gut microbiota composition was found to be significantly different across serum vitamin D subgroups. Particularly, the levels of Bacteroidetes, Firmicutes, and total bacteria were significantly higher in the optimal vitamin D group. These results are consistent with those obtained from Kanhere et al. (20), who found that the level of Bacteroidia, a bacterial class belonging to Bacteroidetes, was higher in adults with optimal vitamin D. Another important finding in this study was the negative correlation between vitamin D status and E. coli. A significant association was found between suboptimal vitamin D levels and a higher abundance of Gammaproteobacteria, a class of bacteria that includes E. coli (40). This link between vitamin D status and microbial colonization can be explained by the significant role vitamin D plays in modulating the immune system in the intestine (41). The absence of the anticipated association between beneficial species such as Bifidobacteria and Lactobacillus spp. and vitamin D level in this study is perhaps related to the redundant effect of vitamin D at its optimal level to modify the existing dysbiosis related to the pro-inflammatory environment in children with CF. In fact, this may highlight the need to increase the currently prescribed regimen for vitamin D in children with CF to reach visible changes in gut dysbiosis. Clinical studies that assess the effect of vitamin D supplementation on gut dysbiosis in children with CF are warranted.

With respect to the association between gastrointestinal bacterial species and PFTs, our study showed higher levels of Firmicutes in children with mild airflow obstruction compared with those with moderate obstruction. Similarly, a previous study showed a significantly higher level of Firmicutes in children with CF with mild lung disease than those with moderate lung disease (42). This can be explained by the role Firmicutes play in producing short-chain fatty acids (SCFAs), particularly butyrate, which has been shown to downregulate pro-inflammatory factors, such as IL-6 and IL-12 (43). SCFAs produced by Bacteroides, Bifidobacterium, Fecalibacterium, and Enterobacteria are also known for promoting the release of anti-inflammatory cytokines (IL-10 and IL-21) and the inhibition of lipopolysaccharide (LPS)-induced pro-inflammatory cytokines (44, 45). As a result, SCFA plays a major role in the modulation of intestinal regulatory T cells which are essential in which they maintain the host immunological homeostasis (44). In CF, the increase in pro-inflammatory factors, attributed to CFTR deficiency in the epithelial cells of the airways, has been linked to the exacerbation of lung function (46, 47).

We acknowledge that our study has certain limitations. The major limitation of this study is the absence of gut microbiota composition assessment of children without CF, due to lack of access to their samples. Furthermore, since the study excluded unstable children with CF, it was not possible to evaluate the effect of vitamin D status and gut microbiota composition on pulmonary exacerbations. Finally, this study was cross-sectional in nature, which neglects the potential prospective impact. Despite these limitations, this study has some strengths as it is a nutrition-inclusive study that considered...
CONCLUSION

The most significant finding to emerge from this study is the difference in serum 25(OH)D levels between the CF and non-CF groups. Another important finding is the significant difference in gut microbiota composition in relation to vitamin D status and PFTs in children with CF. This study identified two phyla (Firmicutes and Bacteroidetes) to be significantly associated with optimal serum vitamin D levels and one species that has a negative correlation with serum vitamin D, which is E. coli. Moreover, this study has provided a deeper insight into the relationship between gut microbiota and pulmonary outcome as it demonstrated a positive correlation of PFTs with Firmicutes and the ratio of Firmicutes to Bacteroidetes. Our findings strengthen the idea of modifying gut microbiota to achieve better outcomes in CF in terms of pulmonary function. Future studies need to be conducted to investigate the benefit of probiotics use as a part of clinical management for CF and measuring its impact on clinical outcomes, such as pulmonary function and pulmonary exacerbations, and vitamin D status.

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