Efficacy and Safety of Elexacaftor/Tezacaftor/Ivacaftor in Children 6 Through 11 Years of Age with Cystic Fibrosis Heterozygous for F508del and a Minimal Function Mutation
A Phase 3b, Randomized, Placebo-controlled Study

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Abstract

Rationale: The triple-combination regimen elexacaftor/tezacaftor/ivacaftor (ELX/TEZ/IVA) was shown to be safe and efficacious in children aged 6 through 11 years with cystic fibrosis and at least one F508del-CFTR allele in a phase 3, open-label, single-arm study.

Objectives: To further evaluate the efficacy and safety of ELX/TEZ/IVA in children 6 through 11 years of age with cystic fibrosis heterozygous for F508del and a minimal function CFTR mutation (F/MF genotypes) in a randomized, double-blind, placebo-controlled phase 3b trial.

Methods: Children were randomized to receive either ELX/TEZ/IVA (n = 60) or placebo (n = 61) during a 24-week treatment period. The dose of ELX/TEZ/IVA administered was based on weight at screening, with children <30 kg receiving ELX 100 mg once daily, TEZ 50 mg once daily, and IVA 75 mg every 12 hours, and children ≥30 kg receiving ELX 200 mg once daily, TEZ 100 mg once daily, and IVA 150 mg every 12 hours (adult dose).

Measurements and Main Results: The primary endpoint was absolute change in lung clearance index from baseline through Week 24. Children given ELX/TEZ/IVA had a mean decrease in lung clearance index2.5 of 2.29 units (95% confidence interval [CI], 1.97–2.60) compared with 0.02 units (95% CI, −0.29 to 0.34) in children given placebo (between-group treatment difference, −2.26 units; 95% CI, −2.71 to −1.81; P < 0.0001). ELX/TEZ/IVA treatment also led to improvements in the secondary endpoint of sweat chloride concentration (between-group treatment difference, −51.2 mmol/L; 95% CI, −55.3 to −47.1) and in the other endpoints of percent predicted FEV1 (between-group treatment difference, 11.0 percentage points; 95% CI, 6.9–15.1) and Cystic Fibrosis Questionnaire-Revised Respiratory domain score (between-group treatment difference, 5.5 points; 95% CI, 1.0–10.0) compared with placebo from baseline through Week 24. The most common adverse events in children receiving ELX/TEZ/IVA were headache and cough (30.0% and 23.3%, respectively); most adverse events were mild or moderate in severity.

Conclusions: In this first randomized, controlled study of a cystic fibrosis transmembrane conductance regulator modulator conducted in children 6 through 11 years of age with F/MF genotypes, ELX/TEZ/IVA treatment led to significant improvements in lung function, as well as robust improvements in respiratory symptoms and cystic fibrosis transmembrane conductance regulator function. ELX/TEZ/IVA was generally safe and well tolerated in this pediatric population with no new safety findings.

Keywords: cystic fibrosis; elexacaftor; tezacaftor; ivacaftor; children
Cystic fibrosis (CF) is an autosomal recessive disease that results from mutations in the CF transmembrane conductance regulator (CFTR) gene (1). More than 1,000 pathogenic CFTR mutations have been described (2, 3); the F508del-CFTR mutation is the most common of these, being present in nearly 90% of patients with CF in some parts of the world (4). Patients with F508del-CFTR mutations have decreases in the quantity and function of the CFTR anion channel present at epithelial cell surfaces, leading to diverse clinical consequences that manifest early in life and include pancreatic insufficiency, growth impairment, and progressive lung disease (5–9).

CFTR modulators are small-molecule therapeutics designed to address the underlying cause of CF (8, 9). CFTR correctors, such as elexacaftor (ELX) and tezacaftor (TEZ), improve CFTR processing and trafficking to epithelial surfaces, whereas CFTR potentiators, such as ivacaftor (IVA), enhance CFTR channel gating (5, 10). In adolescents and adults who are heterozygous for F508del and a minimal function CFTR mutation (F/MF genotypes) or homozygous for F508del (F/F genotype), a triple combination regimen of ELX/TEZ/IVA was shown to be safe and efficacious (11–13). ELX/TEZ/IVA treatment resulted in robust and clinically meaningful improvements in lung function (as assessed by percent predicted FEV1, [ppFEV1]), respiratory symptoms (as assessed by Cystic Fibrosis Questionnaire-Revised [CFQ-R] respiratory domain score), and CFTR function (as assessed by sweat chloride concentration) in these patients and provided greater efficacy than the previously approved dual combination of TEZ/IVA in patients with the F/F genotype. These results established ELX/TEZ/IVA as a highly effective treatment for adolescents and adults with CF who have at least one F508del allele (14).

The progressive lung disease associated with CF develops early in life, with pulmonary infection, inflammation, and structural lung damage occurring frequently in school-aged children with CF; thus, early treatment is critical to improving clinical outcomes and life expectancy (15–19).

Sweat chloride and baseline lung function are considered important efficacy endpoints in clinical trials involving children with CF. The lung clearance index2.5 (LCI2.5) was generally safe and well tolerated, indicating a safety profile in 6–through 11-year-olds consistent with that previously established in adults and adolescents (20). Further studies showed that children obtained similar clinical benefits from ELX/TEZ/IVA treatment as older patients, despite having a lower baseline lung function and CFQ-R respiratory domain scores than adults and adolescents (20). Because the primary objective of the open-label study in children aged 6 through 11 years was to assess safety, a placebo-controlled trial focused on efficacy was performed to better understand the extent to which ELX/TEZ/IVA treatment ameliorates early airway disease and improves lung function in this pediatric population.

Here, we report results from a 24-week placebo-controlled trial designed to quantify the efficacy of ELX/TEZ/IVA in children 6 through 11 years of age with CF with F/MF genotypes. Absolute change in LCI2.5 was designated the primary endpoint because LCI derived from multiple-breath washout testing is considered a highly sensitive measure for small airway disease and lung function change in this age group and has been shown to detect treatment responses in children who have normal spirometry values (ppFEV1 ≥ 80 percentage points) (19, 21–25).

Given the substantial clinical benefits of ELX/TEZ/IVA observed in adults and adolescents with at least one F508del allele (11, 12), an open-label phase 3 study was conducted to assess the safety, pharmacokinetics, and efficacy of ELX/TEZ/IVA in pediatric patients aged 6 through 11 years with either F/MF or F/F genotypes (20). In this trial, ELX/TEZ/IVA was generally safe and well tolerated, indicating a safety profile in 6–through 11-year-olds consistent with that previously established in adults and adolescents (20). Furthermore, ELX/TEZ/IVA treatment led to improvements in ppFEV1, CFQ-R respiratory domain score, lung clearance index2.5 (LCI2.5), and sweat chloride concentration. These results suggest that children obtain similar clinical benefits from ELX/TEZ/IVA treatment as older patients, despite having a lower baseline lung function and CFQ-R respiratory domain scores than adults and adolescents (20). Because the primary objective of the open-label study in children aged 6 through 11 years was to assess safety, a placebo-controlled trial focused on efficacy was performed to better understand the extent to which ELX/TEZ/IVA treatment ameliorates early airway disease and improves lung function in this pediatric population.
Methods

Participants, Trial Design, and Oversight

This phase 3b, randomized, double-blind, placebo-controlled, multicenter trial of ELX/TEZ/IVA enrolled children aged 6 through 11 years with CF and F/MF genotypes and LCI2,5 $>$ 7.5. The CFTR genotype was confirmed as part of screening. Placebo was considered the most appropriate comparator because, at the time the study was conducted, there was no approved CFTR modulator for children 6 through 11 years of age with F/MF genotypes. For additional details on eligibility criteria, including a list of qualifying MF mutations, see Table E1 in the online supplement.

Children were randomized (1:1) to receive either ELX/TEZ/IVA or a placebo over a 24-week treatment period (Figure E1). Randomization was stratified by LCI2,5 at screening (<10 mmol/L vs. $\geq$ 10 mmol/L) and weight at screening (<30 kg vs. $\geq$ 30 kg). Dosing was based on weight at screening: children weighing <30 kg received ELX 100 mg once daily, TEZ 50 mg once daily, and IVA 75 every 12 hours (50% of adult dose), whereas children weighing $\geq$ 30 kg received ELX 200 mg once daily, TEZ 100 mg once daily, and IVA 150 every 12 hours (full adult dose).

The trial was designed by Vertex Pharmaceuticals Incorporated in collaboration with the authors. For each child enrolled in the study, informed consent was provided by a parent or legal guardian; assent was obtained from the participants in accordance with local regulations. Safety was monitored by an independent data monitoring committee. Vertex Pharmaceuticals performed data collection and analysis in collaboration with the authors and the VX19–445–116 Study Group. Authors had full access to trial data after the final database lock, critically reviewed the manuscript, and approved it for final submission. The investigators vouch for the accuracy and completeness of the data generated at their sites, and the investigators and Vertex Pharmaceuticals vouch for the fidelity of the trial to the protocol.

As this study was initiated during the coronavirus disease 2019 (COVID-19) pandemic, a global protocol addendum provided participants with options to minimize the risk of COVID-19 exposure that might occur through travel.

Implemented measures, as permitted by country and local regulations, enabled remote consent, remote monitoring visits, in-home assessments, and shipment of study drugs to the homes of participants.

Outcome Measures

The primary endpoint was absolute change in LCI2,5 from baseline through Week 24. The EcoMedics Exhalyzer-D multiple-breath washout device with Spiroware Version 3.1.6 was used to determine individual LCI results. Secondary endpoints were absolute change in sweat chloride concentration from baseline through Week 24 and safety and tolerability as assessed by adverse events (AEs), clinical laboratory values, electrocardiograms, vital signs, pulse oximetry, and ophthalmologic examinations. Other efficacy endpoints included absolute changes in ppFEV1 and CFQ-R respiratory domain score from baseline through Week 24. A post hoc analysis was conducted to assess the proportion of children achieving sweat chloride concentrations $<$ 60 mmol/L and $<$ 30 mmol/L.

Statistical Analysis

The primary null hypothesis tested was that the mean absolute change in LCI2,5 from baseline to Week 24 was the same for the two treatment groups (ELX/TEZ/IVA and placebo). A sample size of 49 children per group was determined with a 2-sided, 2-sample $t$ test at a significance level of 0.05. The target for enrollment was 108 participants, allowing for 10% dropout during the treatment period. A mixed-effects model for repeated measures was used to analyze absolute changes in LCI2,5, sweat chloride concentration, ppFEV1, and CFQ-R respiratory domain score. The model included treatment group, visit, and treatment-by-visit interaction as fixed effects, with continuous baseline LCI2,5 and weight at screening (<30 kg vs. $\geq$ 30 kg) as covariates. The primary result obtained from the model was the estimated treatment difference through Week 24 (defined as the average of Weeks 4, 8, 16, and 24). The main analyses for safety included all data collected up to Week 24 in the treatment period and included both in-clinic and at-home assessments. Further details on the statistical analyses are provided in the online supplement.

Results

Population

The trial was conducted at 34 sites in Australia, Canada, Denmark, France, Germany, Israel, Netherlands, Spain, Switzerland, and the United Kingdom from 19 June 2020 to 17 May 2021. Overall, 121 children were randomized and received 1 or more doses of either ELX/TEZ/IVA or placebo in the 24-week treatment period: 60 children received ELX/TEZ/IVA, and 61 children received placebo (Figure 1). One child (1.7%) discontinued ELX/TEZ/IVA because of an AE of rash. The baseline demographics and clinical characteristics were similar between the two treatment groups (Tables 1 and E2).

Efficacy

The primary endpoint of this study was absolute change in LCI2,5 from baseline

Figure 1. Participant disposition diagram. AE = adverse event; ELX/TEZ/IVA = elexacaftor/tezacaftor/ivacaftor.
Table 1. Demographics and Clinical Characteristics of the Participants at Baseline

|                | Placebo n = 61 | ELX/TEZ/IVA n = 60 |
|----------------|---------------|-------------------|
| Female sex, n (%) | 35 (57.4) | 35 (58.3) |
| Age at baseline, mean (SD), y | 9.2 (1.7) | 9.1 (1.8) |
| Race, n (%)† |            |                  |
| White | 42 (68.9) | 45 (75.0) |
| Black or African American | 0 (0) | 1 (1.7) |
| Asian | 0 (0) | 1 (1.7) |
| American Indian or Alaska Native | 0 (0) | 1 (1.7) |
| Native Hawaiian or other Pacific Islander | 0 (0) | 0 (0) |
| Other | 1 (1.6) | 0 (0) |
| Not collected per local regulations | 18 (29.5) | 11 (18.3) |
| Ethnicity, n (%) |            |                  |
| Hispanic or Latino | 0 (0) | 1 (1.7) |
| Not Hispanic or Latino | 42 (68.9) | 48 (80.0) |
| Not collected per local regulations | 19 (31.1) | 11 (18.3) |
| Geographic region, n (%) |            |                  |
| Europe | 49 (80.3) | 43 (71.7) |
| Other countries (Australia, Canada, Israel) | 12 (19.7) | 17 (28.3) |
| Weight, mean (SD), kg | 29.8 (8.6) | 29.1 (7.6) |
| Weight distribution, n (%) |      |                  |
| <30 kg | 38 (62.3) | 39 (65.0) |
| ≥30 kg | 23 (37.7) | 21 (35.0) |
| Height-for-age z-score, mean (SD) |      |                  |
| –2.09 (0.96) | –0.27 (0.99) |
| Height, mean (SD), cm | 134.6 (13.3) | 132.3 (11.7) |
| Height-for-age z-score, mean (SD) |      |                  |
| 0.01 (1.26) | –0.17 (1.02) |
| BMI, mean (SD, kg/m²) | 16.11 (2.32) | 16.33 (1.84) |
| BMI-for-age z-score, mean (SD) |      |                  |
| –0.39 (0.92) | –0.17 (0.85) |
| LCI2.5, mean (SD) | 9.75 (1.95) | 10.26 (2.22) |
| Sweat chloride concentration, mean (SD), mmol/L |     |                  |
| 102.6 (8.6) | 102.8 (10.0) |
| ppFEV₁, mean (SD) | 87.2 (15.8) | 91.4 (13.8) |
| ppFEV₁ category, n (%) |      |                  |
| <70 | 10 (16.4) | 4 (6.7) |
| ≥70 to <90 | 23 (37.7) | 20 (33.3) |
| ≥90 | 28 (45.9) | 36 (60.0) |
| CFQ-R respiratory domain score (child’s version), mean (SD) points‡ | 82.7 (14.1) | 85.7 (11.7) |

Definition of abbreviations: BMI = body mass index; CFQ-R = Cystic Fibrosis Questionnaire Revised; ELX/TEZ/IVA = elixacaftor/tezacaftor/ivacaftor; LCI 2.5 = lung clearance index 2.5; ppFEV₁ = percent predicted FEV₁.

*Child’s version, †race categories may sum to more than 100% because each participant could indicate more than one race, ‡child’s version of CFQ-R was used in the assessment. Scores for the CFQ-R respiratory domain range from 0 to 100, with higher scores indicating a higher patient-reported quality of life with regard to respiratory status.

through Week 24. Mean LCI2.5 at baseline was 10.26 units (standard deviation [SD], 2.22) in the ELX/TEZ/IVA group and 9.75 units (SD, 1.95) in the placebo group (Table 1). Children who received ELX/TEZ/IVA had a mean change in LCI2.5 of –2.29 units (95% confidence interval [CI], –3.40 to –1.19), whereas children who received placebo had a mean change of 0.02 units (95% CI, –0.34 to 0.29); the between-group treatment difference was –2.31 units (95% CI, –3.27 to –1.35; P < 0.0001) (Table 2 and Figure 2A).

An ad hoc subgroup analysis showed no evidence of group treatment interaction (Table 3). The majority of AEs that were mild or moderate in severity and generally consistent with manifestations of CF. The most common AEs (≥15% of children) in the ELX/TEZ/IVA group were cough (30%) and headache (23.3%) and in the placebo group were cough (24.6%), abdominal pain (27.9%), and oropharyngeal pain (19.7%). Serious AEs were reported in 6% and 3% of children in the ELX/TEZ/IVA and placebo groups, respectively. The incidence of serious AEs was similar in the ELX/TEZ/IVA and placebo groups (Table 4).

Safety Overall, 48 children (80%) who received ELX/TEZ/IVA and 57 children (93.4%) who received placebo had AEs (Table 3). The majority had AEs that were mild or moderate in severity and generally consistent with manifestations of CF. The most common AEs (≥15% of children) in the ELX/TEZ/IVA group were cough (30%) and headache (23.3%) and in the placebo group were cough (24.6%), abdominal pain (27.9%), and oropharyngeal pain (19.7%). Serious AEs were reported in 6% and 3% of children in the ELX/TEZ/IVA and placebo groups, respectively. The incidence of serious AEs was similar in the ELX/TEZ/IVA and placebo groups (Table 4).
occurred in 4 children (6.7%) receiving ELX/TEZ/IVA and in 9 children (14.8%) receiving a placebo. One child (1.7%) who received ELX/TEZ/IVA had a serious AE of rash that resolved after treatment discontinuation. On the basis of previous experience with ELX/TEZ/IVA, including phase 3 trials in participants 12 years of age or older and in children 6 through 11 years of age (11, 12, 20), data related to aminotransferases, rash events, blood pressure, and creatine kinase were reviewed. Among children who received ELX/TEZ/IVA, elevated concentrations of alanine aminotransferase and/or aspartate aminotransferase more than three times the upper limit of normal (ULN) occurred in eight children (13.6%), with three (5.1%) having concentrations more than five times the ULN and one (1.7%) having concentrations more than eight times the ULN. Among children who received placebo, three (4.9%) had elevated concentrations of alanine aminotransferase and/or aspartate aminotransferase more than three times the ULN, with one child (1.6%) having concentrations more than five times the ULN and no children having concentrations more than eight times the ULN (Table E5). No children had alanine aminotransferase and/or aspartate aminotransferase concentrations more than three times the ULN concurrent with total bilirubin concentrations more than two times ULN. Adverse events of elevated aminotransferases were reported in six children (10.0%) who received ELX/TEZ/IVA and in three children (4.9%) who received placebo, all of which were mild or moderate in severity and none of which were considered serious or led to treatment discontinuation.

Eight children (13.3%) who received ELX/TEZ/IVA and three children (4.9%) who received placebo had rash events (Table E6). Rash events comprised a group AE term that included preferred terms of rash, rash erythematous, rash maculopapular, rash papular, skin exfoliation, and urticaria. Among children who had rash events, most had events that were mild or moderate in severity. One child (1.7%) had a serious AE of rash that developed on Day 8 of ELX/TEZ/IVA treatment. This AE resolved after study drug discontinuation and treatment with antihistamines and topical steroids. No other children discontinued treatment because of rash events.

In children who received ELX/TEZ/IVA, the mean change from baseline in systolic blood pressure (mm Hg) ranged from 0.1 (Day 15) to 2.6 (Week 8), and in diastolic blood pressure ranged from −2.1 (Day 15) to 1.1 (Week 8) (Table E7). In children who received placebo, the mean change from baseline in systolic blood pressure ranged from 0.0 (Week 4) to 2.6 (Week 16), and in diastolic blood pressure ranged from −0.3 (Week 4) to 1.3 (Week 8). No children had AEs of blood pressure increased. No children had creatine kinase concentrations more than five times the ULN (Table E8). There were no notable safety findings in other clinical or laboratory assessments.

**Discussion**

The efficacy and safety of ELX/TEZ/IVA were evaluated in a 24-week randomized, double-blind, placebo-controlled trial in children 6 through 11 years of age with F/MF genotypes. Treatment with ELX/TEZ/IVA resulted in significant improvements in LCI2.5 as well as robust improvements in ppFEV1, CFQ-R respiratory domain score, and sweat chloride concentration compared with placebo. Safety data were consistent with the established safety profile for ELX/TEZ/IVA, with no new safety concerns observed.

Impaired lung function is a hallmark of CF disease progression that begins early in life (17). In adults and adolescents with CF, lung function impairment is typically assessed using spirometry. However, in children with CF, baseline FEV1 is often

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**Table 2. Primary, Secondary, and Other Efficacy Endpoints**

|                          | Placebo n = 61 | ELX/TEZ/IVA n = 60 |
|--------------------------|---------------|-------------------|
| **Primary endpoint**     |               |                   |
| Absolute change          |               |                   |
| in LCI2.5, units         |               |                   |
| Baseline, mean (SD)^*    | 9.75 (1.95)   | 10.26 (2.22)      |
| Absolute change through  | −0.02 (−0.34 to 0.29) | −2.29 (−2.60 to −1.97) |
| Week 24, LS mean (95% CI)| −2.26 (−2.71 to −1.81) | P < 0.0001 |
| Between-group difference |               |                   |
| (95% CI)                 |               |                   |
| **Secondary endpoint**   |               |                   |
| Absolute change in       |               |                   |
| sweat chloride, mmol/L   |               |                   |
| Baseline, mean (SD)^*    | 102.6 (8.6)   | 102.8 (10.0)      |
| Absolute change through  | −0.9 (−3.8 to 2.0) | −52.1 (−55.0 to −49.2) |
| Week 24, LS mean (95% CI)| −51.2 (−55.3 to −47.1) | P < 0.0001† |
| Between-group difference |               |                   |
| (95% CI)                 |               |                   |
| **Other endpoint**       |               |                   |
| Absolute change in       |               |                   |
| ppFEV1, percentage points|               |                   |
| Baseline, mean (SD)^*    | 87.2 (15.8)   | 91.4 (13.8)       |
| Absolute change through  | −1.5 (−4.4 to 1.4) | 9.5 (6.6 to 12.4) |
| Week 24, LS mean (95% CI)| 11.0 (6.9 to 15.1) | P < 0.0001† |
| Between-group difference |               |                   |
| (95% CI)                 |               |                   |
| **Other endpoint**       |               |                   |
| Absolute change in CFQ-R |               |                   |
| respiratory domain score,|               |                   |
| points                   |               |                   |
| Baseline, mean (SD)^*    | 82.7 (14.1)   | 85.7 (11.7)       |
| Absolute change through  | 0.5 (−2.7 to 3.6) | 5.9 (2.8 to 9.1)  |
| Week 24, LS mean (95% CI)| 5.5 (1.0 to 10.0) | P = 0.0174†      |
| Between-group difference |               |                   |
| (95% CI)                 |               |                   |

**Definition of abbreviations:** CFQ-R = Cystic Fibrosis Questionnaire Revised; CI = confidence interval; ELX/TEZ/IVA = elexacaftor/tezacaftor/ivacaftor; LCI2.5 = lung clearance index2.5; LS = least-squares; ppFEV1 = percent predicted FEV1; SD = standard deviation.

^Baseline was defined as the most recent nonmissing measurement before the first dose of the study drug in the treatment period.

†P values are considered to be nominal.
Figure 2. Efficacy results by visit. (A) Absolute change in LCI₂₅ from baseline at each visit. Lower values indicate decreased airway obstruction and improved homogeneity of ventilation. (B) Absolute change in sweat chloride concentration from baseline at each visit; lower values indicate increased CFTR function. (C) Absolute change in ppFEV₁ from baseline at each visit. (D) Absolute change in the respiratory domain score on
within the normal range (26), as was seen in the children enrolled in this study. LCI$_{2.5}$, a measure of ventilation inhomogeneity derived from the multiple-breath washout test, can detect early changes in lung function and small airway disease and is therefore considered a more sensitive predictor of lung disease progression than FEV$_1$ in children with CF (27, 28). A longitudinal natural history study showed that children 6 through 11 years of age with CF who were not treated with a CFTR modulator had an annual increase in LCI$_{2.5}$ of 0.21 units (29).

Treatment with the CFTR modulators TEZ/IVA (within-group change $-0.51$ units at Week 8) and LUM/IVA (within-group change $-0.88$ at Week 24) was associated with improved LCI$_{2.5}$ in children 6 through 11 years (30, 31). In the current study, abnormal LCI$_{2.5}$ ($\geq 7.5$) values were an inclusion criterion, indicating the presence of early small airway disease in these children. In contrast to the natural history data, as well as the TEZ/IVA and LUM/IVA data described above, ELX/TEZ/IVA treatment resulted in a statistically significant improvement in LCI$_{2.5}$ of $-2.26$ units from baseline through Week 24 compared with placebo, which was rapid (occurring by Day 15) and sustained. Although the minimal clinically important difference for LCI$_{2.5}$ has not been defined, our results indicate ELX/TEZ/IVA treatment is associated with a robust and sustained improvement in small airway function and pulmonary ventilation in children with CF.

After the data from this clinical trial had been analyzed, the software for the EcoMedics Exhalyzer-D multiple-breath washout device, used for LCI$_{2.5}$ assessment in the current study, was updated to correct for cross-sensitivity in the device’s oxygen and carbon dioxide sensors that would otherwise result in overestimation of the nitrogen concentration (32). The effect of this software update on the interpretation of LCI results was assessed in a recent report that reanalyzed data sets from six previous studies involving 1,036 multiple-breath washout tests (33). As expected, the correction algorithm resulted in somewhat lower LCI values but did not change their interpretation or the significance of treatment effects (33).

Table 3. Adverse Events*$^*$

|                      | Placebo $n=61$, $n$ (%) | ELX/TEZ/IVA $n=60$, $n$ (%) |
|----------------------|-------------------------|----------------------------|
| Any AE               | 57 (93.4)               | 48 (80.0)                  |
| AE by maximum severity† |                        |                            |
| Mild                 | 26 (42.6)               | 30 (50.0)                  |
| Moderate             | 29 (47.5)               | 16 (26.7)                  |
| Severe               | 2 (3.3)                 | 2 (3.3)                    |
| Serious AE           | 9 (14.8)                | 4 (6.7)                    |
| Serious related AE   | 1 (1.6)                 | 1 (1.7)†                   |
| AE leading to death  | 0 (0)                   | 0 (0)                      |
| AE leading to discontinuation | 0 (0) | 1 (1.7)†                   |
| Most prevalent AEs²  |                        |                            |
| Headache             | 12 (19.7)               | 18 (30.0)                  |
| Cough                | 26 (42.6)               | 14 (23.3)                  |
| Nasopharyngitis      | 9 (14.8)                | 7 (11.7)                   |
| Productive cough     | 6 (9.8)                 | 7 (11.7)                   |
| Rhinorrhea           | 7 (11.5)                | 7 (11.7)                   |
| Rash                 | 3 (4.9)                 | 6 (10.0)                   |
| Abdominal pain       | 17 (27.9)               | 5 (8.3)                    |
| Oropharyngeal pain   | 12 (19.7)               | 3 (5.0)                    |
| Infective pulmonary exacerbation of cystic fibrosis | 16 (26.2) | 1 (1.7) |

Definition of abbreviations: AE = adverse event; ELX/TEZ/IVA = elexacaftor/tezacaftor/ivacaftor.

* A participant with multiple events within a category was counted only once in that category.
† Severity was determined by the investigator observing the event.
‡ One child had a serious AE of rash that was considered possibly related to ELX/TEZ/IVA and resolved after study discontinuation.
§ Only AEs that occurred in $\geq 10\%$ of participants are listed; the listing is according to the preferred term (Medical Dictionary for Regulatory Activities version 24.0).

In light of the robust LCI$_{2.5}$ treatment effect seen with ELX/TEZ/IVA in the current study and the results of the reanalysis performed with updated software in the study above, it is highly unlikely that a reanalysis of the LCI data would alter the interpretation of the results obtained with the prespecified analysis as reported here.

Children in this study also had substantial improvements in both ppFEV$_1$ and CFQ-R respiratory domain score. Decreases in FEV$_1$ are a sensitive indicator of airflow limitation in larger conducting airways that are often seen in adolescents and adults with CF, corresponding to the degree of airway obstruction determined by wall thickening and mucus plugging (34). In this study, despite preserved spirometry at baseline, children given ELX/TEZ/IVA had a mean increase of 11.0 percentage points in ppFEV$_1$ compared with placebo from baseline through Week 24, similar to the improvement observed in older patients with $F/MF$ genotypes treated with ELX/TEZ/IVA who had substantially lower baseline ppFEV$_1$ values as well as in the open-label study of ELX/TEZ/IVA in this age group (11, 20).

This result indicates that, in addition to the substantial improvement in small airway function seen on the basis of the improvement in LCI$_{2.5}$ with ELX/TEZ/IVA treatment, these children also had improved large airway function, further confirming the robust clinical benefit of ELX/TEZ/IVA treatment on lung function in this pediatric population. Furthermore, children given ELX/TEZ/IVA also had a mean increase of 5.5 points in CFQ-R respiratory domain score compared with placebo, reflecting improved respiratory symptoms and exceeding the minimal clinically important difference of four points (35). These findings corroborate the marked improvement in respiratory status indicated by improved LCI$_{2.5}$ as the primary endpoint.

Figure 2. (Continued). the CFQ-R (child’s version) from baseline at each visit; scores normalized to a 100-point range, with higher scores indicating a higher patient-reported quality of life with regard to respiratory symptoms. Data are least-squares means based on a mixed-effects model for repeated measures; l-bars indicate the standard error of the mean, and the dashed horizontal line corresponds to the baseline. The sample size shown below the x-axis is the number of children at the time point with evaluable in-clinic data. CFQ-R = Cystic Fibrosis Questionnaire Revised; ELX/TEZ/IVA = elexacaftor/tezacaftor/ivacaftor; LCI$_{2.5}$ = lung clearance index 2.5; ppFEV$_1$ = percent predicted FEV$_1$. 

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Sweat chloride concentration provides a direct measure of systemic CFTR function (6). A recent observational study in adolescents and adults with CF and at least one F508del allele showed improvements in sweat chloride concentration achieved with ELX/TEZ/IVA treatment were associated with improvements in CFTR function in a clinical trial (726). In addition, 80% of children achieved sweat chloride concentrations <60 mmol/L from baseline through Week 24 compared with placebo. Improvements in respiratory symptoms and CFTR function, similar to those seen in adolescents and adults (11, 12), were observed in children given ELX/TEZ/IVA compared with placebo. ELX/TEZ/IVA was generally safe and well tolerated in this pediatric population with no new safety findings. Taken together, these results demonstrate the ability of ELX/TEZ/IVA treatment to ameliorate early airway disease in CF and alter the natural trajectory of CF disease in children.

Conclusions
In this first randomized controlled study of a CFTR modulator in children 6 through 11 years of age with the F508del genotype, treatment with ELX/TEZ/IVA led to rapid, statistically significant, and clinically meaningful improvements in lung function compared with placebo. Improvements in respiratory symptoms and CFTR function, similar to those seen in adolescents and adults (11, 12), were observed in children given ELX/TEZ/IVA compared with placebo. ELX/TEZ/IVA was generally safe and well tolerated in this pediatric population with no new safety findings. Taken together, these results demonstrate the ability of ELX/TEZ/IVA treatment to ameliorate early airway disease in CF and alter the natural trajectory of CF disease in children.

Author disclosures are available with the text of this article at www.atsjournals.org.

Acknowledgments: The authors thank the patients and their families for participating in this trial; all site investigators and coordinators; the members of the Cystic Fibrosis Foundation Therapeutics Development Network and the European Cystic Fibrosis Society Clinical Trials Network for their support of the trial sites; Swati Thorat, Ph.D., an employee of Vertex Pharmaceuticals, who may own stock or stock options in the company, for providing medical writing and editorial support under the guidance of the authors; and Hostein Heidari Torabadi, Pharm.D., Ph.D., of ArticulateScience, LLC, for providing editorial assistance under the guidance of the authors and with support from Vertex Pharmaceuticals. J.C.D. is supported by the National Institutes of Health Research through a Senior Investigator Award, the Imperial Biomedical Research Centre, and the Brompton Clinical Research Facility.

References
1. Riordan JR, Rommens JM, Kerem B, Alon N, Rozmahel R, Grzelczak Z, et al. Identification of the cystic fibrosis gene: cloning and characterization of complementary DNA. Science 1989;245:1066–1073.
2. Cystic Fibrosis Mutation Database (CFTR1); 2011 [accessed 2022 Jan 13]. Available from: http://www.genet.sickkids.on.ca/.
3. The Clinical and Functional TRanslation of CFTR (CFTR2); 2011 [accessed 2022 Jan 13]. Available from: https://cftr2.org/.
4. Riordan JR. CFTR function and prospects for therapy. Annu Rev Biochem 2008;77:701–726.
5. Bell SC, Mall MA, Gutierrez H, Macek M, Madge S, Davies JC, et al. The future of cystic fibrosis care: a global perspective. Lancet Respir Med 2020;8:65–124.
6. Elborn JS. Cystic fibrosis. Lancet 2016;388:2519–2531.
7. Anderson MP, Gregory RJ, Thompson S, Souza DW, Paul S, Mulligan RC, et al. Demonstration that CFTR is a chloride channel by alteration of its anion selectivity. Science 1991;253:202–205.
8. Ratjen F, Bell SC, Rowe SM, Goss CH, Quittner AL, Bush A. Cystic fibrosis. Nat Rev Dis Primers 2015;1:15010.
9. Mall MA, Mayer-Hamblett N, Rowe SM. Cystic fibrosis: emergence of highly effective targeted therapeutics and potential clinical implications. Am J Respir Crit Care Med 2020;210:1193–1208.
10. Van Goor F, Hadida S, Grootenhuis PD, Burton B, Cao D, Neuberger T, et al. Rescue of CF airway epithelial cell function in vitro by a CFTR potentiator. VX-770. Proc Natl Acad Sci USA 2009;106:18825–18830.
11. Middleton PG, Mall MA, Drevenik P, Lands LC, McKone EF, Polineni D, et al.; VX17-445-102 Study Group. Elexacaftor-tezacaftor-ivacaftor for cystic fibrosis with a single Phen08del allele. N Engl J Med 2019;381:1809–1819.
12. Heijerman HGM, McKone EF, Downey DG, Van Braeckel E, Rowe SM, Tullis E, et al.; VX17-445-103 Trial Group. Efficacy and safety of the elixacaftor plus tezacaftor plus ivacaftor combination regimen in people with cystic fibrosis homozygous for the F508del mutation: a double-blind, randomised, phase 3 trial. Lancet 2019;394:1940–1948.

13. Sutharsan S, McKone EF, Downey DG, Duckers J, MacGregor G, Tullis E, et al.; VX18-445-109 Study Group. Efficacy and safety of elixacaftor plus tezacaftor plus ivacaftor versus tezacaftor plus ivacaftor in people with cystic fibrosis homozygous for F508del-CFTR: a 24-week, multicentre, randomised, double-blind, active-controlled, phase 3b trial. Lancet Respir Med 2022;10:267–277.

14. Middleton PG, Taylor-Cousar JL. Development of elixacaftor - tezacaftor - ivacaftor: highly effective CFTR modulation for the majority of people with cystic fibrosis. Expert Rev Respir Med 2021;15:723–735.

15. Farrell PM, Kosorok MR, Rock MJ, Laxova A, Zeng L, Lai HC, et al. Ventilation inhomogeneities in relation to standard lung function in patients with cystic fibrosis. Am J Respir Crit Care Med 2005;171:371–378.

16. Ranganathan SC, Hall GL, Sly PD, Stick SM, Douglas TA; Australian Respiratory Early Surveillance Team for Cystic Fibrosis (AREST-CF). Early lung disease in infants and preschool children with cystic fibrosis. Thorax 2009;64:1458–1462.

17. Ranganathan SC, Hall GL, Sly PD, Stick SM, Douglas TA. Magnetic resonance imaging detects progression of lung disease and impact of newborn screening in preschool children with cystic fibrosis. Am J Respir Crit Care Med 2021;204:943–953.

18. Stahl M, Steinke E, Graeber SY, Joachim C, Seitz C, Kauczor HU, et al. Comparison of lung clearance index and magnetic resonance imaging for assessment of lung disease in children with cystic fibrosis. Am J Respir Crit Care Med 2017;195:349–359.

19. Stahl M, Wielputz MO, Graeber SY, Joachim C, Sommerburg O, Kauczor HU, et al. Comparison of lung clearance index and magnetic resonance imaging for assessment of lung disease in children with cystic fibrosis. Eur Respir J 2021;58:2000686.

20. Zemanick ET, Taylor-Cousar JL, Davies J, Gibson RL, Mall MA, McKone EF, et al. A phase 3 open-label study of elixacaftor/tezacaftor/ivacaftor in patients aged 6-11 years with cystic fibrosis homozygous for F508del or heterozygous for the F508del-CFTR mutation and a residual function mutation. J Cyst Fibros 2021;20:68–77.

21. Ratjen F, Hug C, Marigowda G, Tian S, Huang X, Stanojevic S, et al. Effect of sensor cross-talk error in Exhalyzer D multiple-breath washout device significantly impacts outcomes in children with cystic fibrosis. J Appl Physiol 2021;131:1148–1156.

22. Robinson PD, Jensen R, Seeto RA, Stanoevsic S, Saunders C, Short C, et al. Impact of cross-sensitivity error correction on representative nitrogen-based multiple breath washout data from clinical trials. J Cyst Fibros 2022;21:e204–e207.

23. Kolodziej M, de Veer MJ, Cholewa M, Egan GF, Thompson BR. Lung function imaging methods in cystic fibrosis pulmonary disease. Respir Res 2017;18:96.

24. Quittner AL, Modi AC, Wainwright C, Otto K, Kirhara J, Montgomery AB. Determination of the minimal clinically important difference scores for the Cystic Fibrosis Questionnaire-Revised respiratory symptom scale in two populations of patients with cystic fibrosis and chronic Pseudomonas aeruginosa airway infection. Chest 2009;135:1610–1618.

25. Graeber SY, Vitzthum C, Pallenberg ST, Naehrlich L, Stahl M, Rohrbach A, et al. Efficacy of elixacaftor/tezacaftor/ivacaftor therapy on CFTR function in patients with cystic fibrosis and one or two F508del alleles. Am J Respir Crit Care Med 2022;205:540–549.

26. Lombardi E, Gambazza S, Pradal U, Braggion C. Lung clearance index in subjects with cystic fibrosis in Italy. Ital J Pediatr 2019;45:56.

27. Kraemer R, Blum A, Schibler A, Ammann RA, Gallati S. Ventilation inhomogeneities in relation to standard lung function in patients with cystic fibrosis. Am J Respir Crit Care Med 2005;171:371–378.

28. Amin R, Subbarao P, Jabar A, Balkovec S, Jensen R, Kerrigan S, et al. Ventilation inhomogeneities in relation to standard lung function in patients with cystic fibrosis. Am J Respir Crit Care Med 2019;195:349–359.

29. Mall, Brugha, Gartner, et al.: ELX/TEZ/IVA in Children with F/MF Genotypes