Cystic fibrosis

ORIGINAL ARTICLE

Results of a phase IIa study of VX-809, an investigational CFTR corrector compound, in subjects with cystic fibrosis homozygous for the F508del-CFTR mutation


ABSTRACT

Background VX-809, a cystic fibrosis transmembrane conductance regulator (CFTR) modulator, has been shown to increase the cell surface density of functional F508del-CFTR in vitro.

Methods A randomised, double-blind, placebo-controlled study evaluated the safety, tolerability and pharmacodynamics of VX-809 in adult patients with cystic fibrosis (n=89) who were homozygous for the F508del-CFTR mutation. Subjects were randomised to one of four VX-809 28 day dose groups (25, 50, 100 and 200 mg) or matching placebo.

Results The type and incidence of adverse events were similar among VX-809- and placebo-treated subjects. Respiratory events were the most commonly reported and led to discontinuation by one subject in each active treatment arm. Pharmacokinetic data supported a once-daily oral dosing regimen. Pharmacodynamic data suggested that VX-809 improved CFTR function in at least one organ (sweat gland). VX-809 reduced elevated sweat chloride values in a dose dependent manner (p=0.0013) that was statistically significant in the 100 and 200 mg dose groups. There was no statistically significant improvement in CFTR function in the nasal epithelium as measured by nasal potential difference, nor were there statistically significant changes in lung function or patient-reported outcomes. No maturation of immature F508del-CFTR was detected in the subgroup that provided rectal biopsy specimens.

Conclusions In this study, VX-809 had a similar adverse event profile similar to placebo in F508del-CFTR homozygous CF patients, and reduced sweat chloride values in a dose dependent manner. The findings indicate that F508del-CFTR is a viable target for drug development.

Key messages

F508del-CFTR is the most common cystic fibrosis-causing mutation. Restoring function to F508del-CFTR offers a novel treatment strategy for the majority of patients with cystic fibrosis. VX-809 had an adverse event profile similar to placebo in F508del-CFTR homozygous CF patients, and reduced sweat chloride values in a dose dependent manner. The findings indicate that F508del-CFTR is a viable target for drug development.

INTRODUCTION

Cystic fibrosis (CF) is the most common autosomal recessive lethal genetic disease in the Caucasian population, with an incidence of 1:3500 births in the USA. The most common cause of morbidity and mortality is lung disease, which is characterised by infection, inflammation and airway damage that leads to respiratory failure.1

CF is caused by mutations in the CF transmembrane conductance regulator (CFTR) gene,2 3 which encodes the CFTR protein. CFTR is a member of the ATP-binding cassette protein family and functions as a chloride ion (Cl\(^-\)) channel and a key regulator of salt and water transport across a variety of epithelia.4–8 The CFTR gene has >1600 reported disease-associated mutations, with F508del-CFTR being the most common. F508del-CFTR results from a 3 bp deletion that leads to the omission of phenylalanine at position 508 of the full-length protein.10 The resulting F508del protein product is unstable and susceptible to rapid degradation in the 26S proteasome, with little if any F508del-CFTR at the plasma membrane.11 12 The F508del-CFTR mutation is found in the majority of patients with CI,13 and therefore the consequences...
of this mutation on the CFTR protein are important to address in therapeutic development for CF.

CFTR ‘correctors’ aim to increase the cell surface density of functional CFTR protein, resulting in improved chloride transport and decreased sodium reabsorption.\(^1\)\(^2\)\(^6\) VX-809 restores F508del-CFTR processing and plasma membrane localisation in primary human bronchial epithelial (HBE) airway cells isolated from patients homozygous for the F508del-CFTR mutation, achieving \(\sim 15\%\) of wild-type CFTR levels as measured by the amount of chloride channel function and the quantity of fully mature, C-Band CFTR.\(^1\)\(^7\)

In this report, we describe the safety, tolerability, pharmacodynamics and pharmacokinetics (PK) of escalating doses of VX-809 in patients with CF homozygous for the F508del-CFTR mutation compared with placebo.

**METHODS**

This study was a randomised, double-blind, placebo-controlled, multiple-dose, multicentre, phase IIa study. Institutional Review Board approvals and informed consents were obtained for all study subjects.

**Subjects**

Subjects enrolled in the study had a confirmed diagnosis of CF, accompanied by a sweat chloride value of \(\geq 60\) mmol/l. All subjects were \(\geq 18\) years of age, and were required to have the F508del-CFTR mutation on both alleles. At screening, subjects were required to have a forced expiratory volume in \(1\) s (FEV\(_1\)) of at least \(40\%\) of predicted normal for age, gender and height (Knudson standards).\(^1\)\(^8\)

**Study design**

Subjects were enrolled into two cohorts, group A and group B. Group A subjects were randomised to receive VX-809 once daily at doses of 25 mg or 50 mg, or placebo in a 2:2:1 randomisation ratio. After 15 group A subjects completed 28 days of therapy, a safety review was conducted by an independent Data Monitoring Committee. Enrolment in group B began following the Data Monitoring Committee review. In group B, subjects were randomised to receive VX-809 at doses of 100 mg or 200 mg, or placebo in a 2:2:1 ratio for 28 days. All study sites, the patients, and the sponsor remained blinded to treatment assignment throughout the study.

**End points**

The primary objective, evaluation of safety and tolerability of VX-809, was assessed by adverse events (AEs), haematology, clinical chemistry, urinalysis, ECG, vital signs and physical examinations. Secondary objectives included evaluation of the pharmacodynamic impact of VX-809 on CFTR function. Measures of CFTR activity included sweat chloride and nasal potential difference (NPD).\(^1\)\(^9\) The latter was considered optional. Other secondary end points included spirometry to measure pulmonary function (ie, FEV\(_1\), forced expiratory flow \(25\%-75\%\) (FEF\(_{25\%-75\%}\)) and forced vital capacity (FVC)) and the CF Questionnaire-Revised (CFQ-R), a disease-specific patient-reported outcome.\(^2\)\(^0\) The minimal clinically important difference in respiratory domain (MCID) is improvement \(\pm 4.2\). The details of CFTR biomarkers and pharmacokinetic analysis are included in the Supplemental Methods section.

**Statistical analyses**

Safety data were analysed primarily using descriptive statistics, and efficacy data were analysed primarily using analysis of covariance (ANCOVA) models. The planned total sample size of 90 subjects provided a study power of \(\geq 97\%\) to detect a 20 mmol/l reduction in sweat chloride concentration from baseline, and a probability of \(99\%\) to observe at least 1 adverse event in the study. All subjects who received at least one dose of study drug were included in the analyses.

**RESULTS**

**Subjects**

A total of 109 adult subjects with CF who were homozygous for the F508del-CFTR mutation (based on screening medical history) were screened and 89 were randomised to one of four VX-809 dose groups (25 mg (\(n = 18\)), 50 mg (\(n = 18\)), 100 mg (\(n = 17\)), 200 mg (\(n = 19\)) or to placebo (\(n = 17\)). Confirmatory genotyping identified one subject randomised to the 50 mg VX-809 group who was heterozygous for F508del-CFTR despite a medical history indicating homozygosity. Safety and efficacy data obtained from this subject were included in the final analysis. The identity of this subject’s non-F508del allele was not available (based on parameters of the informed consent document for that study site). Baseline characteristics are provided in table 1. Median sweat chloride values and FEV\(_1\) percentage predicted were similar between the different groups and consistent with the values reported in the literature for patients homozygous for the F508del-CFTR mutation. The baseline median sweat chloride was 103.5 mmol/l and the median baseline FEV\(_1\) was 71% predicted. The study groups were well matched except for trends towards less severe lung disease in the placebo and 25 mg dose groups.

**Safety and AE profile**

The incidence of AEs was similar between dose groups (table 2). Respiratory events were the most commonly reported type of

<table>
<thead>
<tr>
<th>Table 1</th>
<th>Baseline demographic and clinical characteristics of the study subjects</th>
</tr>
</thead>
<tbody>
<tr>
<td>Characteristic</td>
<td>Placebo n = 17</td>
</tr>
<tr>
<td>Male, n (%)</td>
<td>11 (65)</td>
</tr>
<tr>
<td>Caucasian, n (%)</td>
<td>17 (100)</td>
</tr>
<tr>
<td>Age, years, median (range)</td>
<td>28 (19–49)</td>
</tr>
<tr>
<td>BMI, kg/m(^2), median (range)</td>
<td>23 (19–31)</td>
</tr>
<tr>
<td>FEV(_1), % predicted, median (range)</td>
<td>78.4 (48.8–124.9)</td>
</tr>
<tr>
<td>Sweat chloride, mmol/l, median (range)</td>
<td>106.5 (80.0–125.5)</td>
</tr>
<tr>
<td>NPD—chloride-free iso, mV, median (range)</td>
<td>1.55 (–10.6* to 8.6)</td>
</tr>
</tbody>
</table>

\*High confidence analysis removed two outliers but did not change efficacy analysis.

BMI, body mass index; FEV\(_1\), forced expiratory volume in \(1\) s; NPD, nasal potential difference.
AE, with cough occurring in 46% of VX-809-treated subjects and 41% of placebo-treated subjects. There was no difference in the incidence of physician-diagnosed pulmonary exacerbations between VX-809- and placebo-treated subjects (17% of VX-809 subjects compared with 12% of placebo subjects; \( p = 0.62 \)).

AEs that occurred in more than one subject in any VX-809 treatment arm are included in table 2. Eight AEs were considered severe, including fatigue, sinus congestion, musculoskeletal discomfort, two events of cough and three events of acute pulmonary exacerbation. All pulmonary exacerbations were considered serious.

Four of 89 (5%) subjects discontinued study drug during the study, one subject in each of the VX-809 dose groups. No placebo-treated subjects withdrew from treatment.

### Table 2

Frequency of occurrence of adverse events occurring in more than one subject in any VX-809 treatment group (listed alphabetically by MedRA term)

<table>
<thead>
<tr>
<th>Adverse event</th>
<th>Placebo (n = 17)</th>
<th>VX-809 25 mg (n = 18)</th>
<th>VX-809 50 mg (n = 18)</th>
<th>VX-809 100 mg (n = 17)</th>
<th>VX-809 200 mg (n = 19)</th>
<th>Total (n = 45)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cough</td>
<td>7 (41.2)</td>
<td>10 (55.6)</td>
<td>6 (33.3)</td>
<td>7 (41.2)</td>
<td>10 (52.6)</td>
<td>40 (88.9)</td>
</tr>
<tr>
<td>Headache</td>
<td>3 (17.6)</td>
<td>4 (22.2)</td>
<td>5 (27.8)</td>
<td>2 (11.8)</td>
<td>5 (26.3)</td>
<td>19 (42.2)</td>
</tr>
<tr>
<td>Rales</td>
<td>1 (5.9)</td>
<td>6 (33.3)</td>
<td>2 (11.1)</td>
<td>3 (17.6)</td>
<td>3 (15.8)</td>
<td>15 (33.3)</td>
</tr>
<tr>
<td>Productive cough</td>
<td>3 (17.6)</td>
<td>2 (11.1)</td>
<td>0</td>
<td>4 (23.5)</td>
<td>6 (31.6)</td>
<td>15 (17.8)</td>
</tr>
<tr>
<td>Dyspnea</td>
<td>1 (5.9)</td>
<td>5 (27.8)</td>
<td>3 (16.7)</td>
<td>2 (11.8)</td>
<td>4 (21.1)</td>
<td>15 (33.3)</td>
</tr>
<tr>
<td>Pulmonary exacerbation*</td>
<td>2 (11.8)</td>
<td>4 (22.2)</td>
<td>2 (11.1)</td>
<td>2 (11.8)</td>
<td>4 (21.1)</td>
<td>14 (31.1)</td>
</tr>
<tr>
<td>Fatigue</td>
<td>2 (11.8)</td>
<td>3 (16.7)</td>
<td>3 (16.7)</td>
<td>2 (11.8)</td>
<td>3 (15.8)</td>
<td>13 (28.9)</td>
</tr>
<tr>
<td>Fever</td>
<td>2 (11.8)</td>
<td>2 (11.1)</td>
<td>1 (5.6)</td>
<td>1 (5.9)</td>
<td>5 (26.3)</td>
<td>11 (24.4)</td>
</tr>
<tr>
<td>Nasal congestion</td>
<td>3 (17.6)</td>
<td>2 (11.1)</td>
<td>1 (5.6)</td>
<td>2 (11.8)</td>
<td>2 (10.5)</td>
<td>10 (22.2)</td>
</tr>
<tr>
<td>Wheezing</td>
<td>3 (17.6)</td>
<td>1 (5.6)</td>
<td>4 (22.2)</td>
<td>1 (5.9)</td>
<td>0</td>
<td>9 (20)</td>
</tr>
<tr>
<td>Diahorea</td>
<td>3 (17.6)</td>
<td>3 (16.7)</td>
<td>1 (5.6)</td>
<td>2 (11.8)</td>
<td>0</td>
<td>9 (20)</td>
</tr>
<tr>
<td>Oropharyngeal pain</td>
<td>3 (17.6)</td>
<td>0</td>
<td>3 (16.7)</td>
<td>0</td>
<td>2 (10.5)</td>
<td>8 (17.8)</td>
</tr>
<tr>
<td>Upper respiratory tract infection</td>
<td>1 (5.9)</td>
<td>2 (11.1)</td>
<td>1 (5.6)</td>
<td>3 (17.6)</td>
<td>0</td>
<td>7 (15.6)</td>
</tr>
<tr>
<td>Sinus congestion</td>
<td>2 (11.8)</td>
<td>1 (5.6)</td>
<td>2 (11.1)</td>
<td>0</td>
<td>1 (5.3)</td>
<td>6 (13.3)</td>
</tr>
<tr>
<td>Respiration abnormal</td>
<td>0</td>
<td>1 (5.6)</td>
<td>1 (5.6)</td>
<td>0</td>
<td>4 (21.1)</td>
<td>6 (13.3)</td>
</tr>
<tr>
<td>Haemoptysis</td>
<td>2 (11.8)</td>
<td>1 (5.6)</td>
<td>1 (5.6)</td>
<td>0</td>
<td>2 (10.5)</td>
<td>6 (13.3)</td>
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<tr>
<td>Constipation</td>
<td>0</td>
<td>2 (11.1)</td>
<td>2 (11.1)</td>
<td>1 (5.9)</td>
<td>1 (5.3)</td>
<td>6 (13.3)</td>
</tr>
<tr>
<td>Abdominal pain</td>
<td>1 (5.9)</td>
<td>3 (16.7)</td>
<td>1 (5.6)</td>
<td>0</td>
<td>1 (5.3)</td>
<td>6 (13.3)</td>
</tr>
<tr>
<td>Myalgia</td>
<td>1 (5.9)</td>
<td>0</td>
<td>3 (16.7)</td>
<td>0</td>
<td>1 (5.3)</td>
<td>5 (11.1)</td>
</tr>
<tr>
<td>Post-tussive vomiting</td>
<td>0</td>
<td>0</td>
<td>2 (11.1)</td>
<td>1 (5.9)</td>
<td>1 (5.3)</td>
<td>4 (8.9)</td>
</tr>
<tr>
<td>Nausea</td>
<td>0</td>
<td>3 (16.7)</td>
<td>0</td>
<td>0</td>
<td>1 (5.3)</td>
<td>4 (8.9)</td>
</tr>
<tr>
<td>Nasopharyngitis</td>
<td>0</td>
<td>1 (5.6)</td>
<td>0</td>
<td>1 (5.9)</td>
<td>2 (10.5)</td>
<td>4 (8.9)</td>
</tr>
<tr>
<td>Dizziness</td>
<td>0</td>
<td>1 (5.6)</td>
<td>0</td>
<td>2 (11.8)</td>
<td>1 (5.3)</td>
<td>6 (13.3)</td>
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<tr>
<td>Back pain</td>
<td>0</td>
<td>2 (11.1)</td>
<td>1 (5.6)</td>
<td>0</td>
<td>1 (5.3)</td>
<td>4 (8.9)</td>
</tr>
<tr>
<td>Abdominal pain upper</td>
<td>1 (5.9)</td>
<td>0</td>
<td>0</td>
<td>1 (5.9)</td>
<td>2 (10.5)</td>
<td>4 (8.9)</td>
</tr>
<tr>
<td>Sputum abnormal</td>
<td>0</td>
<td>2 (11.1)</td>
<td>0</td>
<td>0</td>
<td>1 (5.3)</td>
<td>3 (6.7)</td>
</tr>
<tr>
<td>Epistaxis</td>
<td>1 (5.9)</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>2 (10.5)</td>
<td>3 (6.7)</td>
</tr>
<tr>
<td>C-reactive protein increased</td>
<td>0</td>
<td>1 (5.6)</td>
<td>0</td>
<td>2 (11.8)</td>
<td>0</td>
<td>3 (6.7)</td>
</tr>
<tr>
<td>Paranasal sinus hypersecretion</td>
<td>0</td>
<td>2 (11.1)</td>
<td>0</td>
<td>0</td>
<td>2 (4.4)</td>
<td></td>
</tr>
<tr>
<td>Lung hyperinflation</td>
<td>0</td>
<td>0</td>
<td>2 (11.8)</td>
<td>0</td>
<td>2 (4.4)</td>
<td></td>
</tr>
</tbody>
</table>

*Physician-reported pulmonary exacerbation; coded as cystic fibrosis lung according to MedRA term.

### Table 3

VX-809 Pharmacokinetic parameters following once-daily dosing of VX-809 for 28 days

<table>
<thead>
<tr>
<th>Dose Parameter</th>
<th>VX-809 25 mg (n = 18)</th>
<th>VX-809 50 mg (n = 18)</th>
<th>VX-809 100 mg (n = 17)</th>
<th>VX-809 200 mg (n = 19)</th>
<th>Total (n = 45)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Day 1</td>
<td>Cmax (µg/ml)</td>
<td>tmax (h)</td>
<td>Cmax (µg/ml)</td>
<td>tmax (h)</td>
<td></td>
</tr>
<tr>
<td>Mean</td>
<td>0.760</td>
<td>3.5*</td>
<td>1.85</td>
<td>3.1*</td>
<td></td>
</tr>
<tr>
<td>SD</td>
<td>0.280</td>
<td>1.5–9.0†</td>
<td>0.92</td>
<td>1.6</td>
<td></td>
</tr>
<tr>
<td>Mean</td>
<td>1.85</td>
<td>3.1*</td>
<td>2.93</td>
<td>3.0*</td>
<td></td>
</tr>
<tr>
<td>SD</td>
<td>0.70</td>
<td>1.5–12.1†</td>
<td>0.92</td>
<td>1.6</td>
<td></td>
</tr>
<tr>
<td>Mean</td>
<td>2.93</td>
<td>3.0*</td>
<td>2.93</td>
<td>3.0*</td>
<td></td>
</tr>
<tr>
<td>SD</td>
<td>0.92</td>
<td>1.5–6.2†</td>
<td>0.92</td>
<td>1.6</td>
<td></td>
</tr>
<tr>
<td>Mean</td>
<td>6.41</td>
<td>4.0*</td>
<td>6.41</td>
<td>4.0*</td>
<td></td>
</tr>
<tr>
<td>SD</td>
<td>2.55</td>
<td>1.5–6.0†</td>
<td>2.55</td>
<td>1.5–6.1†</td>
<td></td>
</tr>
</tbody>
</table>

*Median.
†Min–Max range.
AR, accumulation ratio; AUC0–24 h, area under the plasma concentration versus time curve from time 0 to 24 h; Cmax, maximum observed drug concentration; CL/F, oral clearance at steady state; t1/2, apparent terminal half-life; tmax, time to reach Cmax after dosing; V/F, apparent volume of distribution.
All discontinuations were due to the occurrence of respiratory AEs. There were no clinically significant changes in laboratory findings during the study.

PK results
Plasma concentrations of VX-809 were measured by a fully validated bioanalytical method. PK parameters estimated at the four dose levels of VX-809 on days 1 and 28 are summarised in table 3. VX-809 appeared to be slowly absorbed from the gut in subjects with CF with median time to reach maximum concentration (t_{max}) values ranging from 3 to 4 h across all doses on days 1 and 28. Based on predose PK samples collected from day 1 to day 28, plasma steady-state concentrations appeared to be reached by day 7 at all dose levels with a mean accumulation ratio ranging from 1.7 to 2.0 on day 28 (based on area under the curve (AUC) over the 24 h dosing period, AUC_{0–24 h}). The time to reach steady state and the extent of accumulation observed in this study were consistent with a terminal half-life approaching 24 h. The estimated mean values of this PK parameter in this study were slightly lower (range from 15 to 18 h) which may be explained by the low number of PK samples collected beyond 24 h on the last dosing day. The estimated oral VX-809 clearance (ie, volume of plasma purified per unit of time) at steady state was relatively low (<2% hepatic blood flow) and comparable for all doses. The estimated volume of distribution of VX-809 at steady state (V_{z/F}) suggested a potential diffusion of the drug into tissues.

- Maximum (C_{max}) and total (AUC_{0–24 h}) exposure to VX-809 increased proportionally with VX-809 dose increases from 25 to 200 mg over the 28 days of treatment. The intersubject variability observed for VX-809 in subjects with CF appeared to be moderate and comparable across all doses for C_{max} (50–40%) but with a trend to increase slightly with dose for AUC_{0–24 h} (from 40% to 60%) on day 28.

CFTR bioactivity
The effects of VX-809 on F508del-CFTR processing and function were evaluated throughout the trial. The assessments included measurements of sweat chloride concentration (in all subjects) and NPD (in 71 subjects). The relative amounts of the immature (B-Band) and mature (C-Band) forms of F508del-CFTR, which are markers of protein maturation beyond the endoplasmic reticulum and Golgi, were measured in biopsies of the rectal epithelium in 34 subjects.

Sweat chloride
Figure 1 shows changes in sweat chloride values, which were reduced in a dose-dependent manner (p=0.0013) in VX-809-treated subjects. The reduction in sweat chloride values was rapid and sustained, with measurable changes seen within 7 days of VX-809 dosing (figure 1A). The mean change from baseline in sweat chloride concentration (mmol/l) after 7 days was 2.2 in the placebo group, −0.5 in the 25 mg group, −3.7 in the 50 mg group (95% CI −7.1 to −0.28, p=0.05), −2.3 in the 100 mg group and −6.6 in the 200 mg group (95% CI −10.27 to −2.83, p=0.0008). At day 28, the mean treatment differences from baseline (−placebo) for the 25, 50, 100 and 200 mg groups were +0.10, −4.61, −6.13 (95% CI −12.25 to −0.01) and −8.21 mmol/l (95% CI −14.33 to −2.10), respectively (figure 1B). These differences were statistically significant versus placebo for the 100 mg (p<0.05) and 200 mg (p<0.01) groups. Following 7 days of drug washout, mean sweat chloride values returned to approximately pretreatment levels (figure 1A).

The number of subjects whose sweat chloride values were reduced by the threshold (‘responder’) values was evaluated using predefined and posthoc study criteria. Using a posthoc responder criterion of ≥10 mmol/l reduction, six subjects (38%) in the 200 mg group and six subjects (40%) in the 100 mg group responded compared with none of the placebo-treated patients (p=0.02 for both VX-809 treatment groups). In the 25 and 50 mg dose groups, there were no subjects classified as responders by either criterion. Using the predefined responder criterion, only one subject (6.3%, in the 200 mg dose group) had a ≥20 mmol/l response to VX-809 (p=NS).

Nasal potential difference
Seventy-one subjects underwent NPD measurements throughout the study. There was no significant change in CFTR-dependent NPD parameters (chloride or sodium transport) in any of the dose groups. Supplementary figure 1 summarises the CFTR chloride ion transport parameters (change in chloride-free isoproterenol response) for the VX-809 treatment groups from baseline to day 28 after removing the change in the placebo group (treatment difference).

CFTR B- to C-Band maturation
Thirty-four subjects provided rectal biopsy tissue for evaluation of F508del-CFTR maturation. Among the 33 subjects who were homozygous for the F508del-CFTR allele, no mature C-Band was detected (supplementary figure 2). Only one subject in the highest dose group (VX-809 200 mg) provided rectal biopsy tissue.

Clinical outcomes
Spirometry
There were no significant changes in lung function (FEV_{1}, FVC, FEF_{25–75%}) in any of the dose groups, including changes in percentage predicted values versus baseline and placebo, or raw measures of litre flow (data not shown). Figure 2 shows the percentage change relative to baseline FEV_{1} (percentage predicted) for each study group across all study visits. After 28 days of treatment, the mean percentage change from baseline in FEV_{1} percentage predicted was 0.07, −2.46, −2.15, 0.32 and 0.47 in the placebo, 25, 50, 100 and 200 mg dose groups, respectively (p=NS).

Patient-reported outcomes
Supplementary table 1 summarises the change in CFQ-R measures (day 28 vs baseline) across the placebo- and VX-809-treated subjects. There were no clear or sustained changes in the respiratory domain or in any other subdomains of the CFQ-R in any dose group. After 28 days, the respiratory domain score in the placebo group increased by 4.5. After 28 days of treatment in the 25, 50, 100 and 200 mg dose groups, the changes were −5.2, −6.3, −1.30 and +2.2, respectively.

DISCUSSION
These data provide evidence of a safety and tolerability profile sufficient to support further clinical evaluation of VX-809 in subjects with CF homozygous for the F508del-CFTR mutation, the most common CF mutation worldwide. Over a dosing range of 25–200 mg, study participants reported symptoms and AEs that were similar to those in the placebo group, and similar to those commonly found in adult patients with CF. One subject in each of the four dose groups developed a pulmonary exacerbation during the 28 day period of treatment, and the incidence of symptoms such as cough was similar across the dose groups and to that of the placebo group.
Sweat chloride concentration measurements demonstrated modest, statistically significant improvements in VX-809-treated subjects. This effect was dose dependent, sustained during the treatment period and rapidly reversed following discontinuation of VX-809. This suggests that VX-809 increased the chloride transport function of CFTR in the sweat gland, and supports the idea that VX-809 is bioactive in the sweat gland of patients homozygous for the F508del-CFTR mutation. These results also support the hypothesis that small molecule correction of F508del-CFTR is feasible. The results provide support for the use of previously obtained CFTR biomarker data sets for study planning of new CFTR modulators. Using this paradigm, the treatment effects on sweat chloride reported with the CFTR potentiator VX-770 in patients with CF with the G551D-CFTR mutation were successfully used to power the current study to detect improvements in sweat chloride (within-group comparisons and with placebo). It is not clear if the reductions in sweat chloride in this study achieved a peak effect (figure 1), and this raises the question of whether higher doses of VX-809 may produce greater effects on F508del-CFTR. The peak corrective effect of VX-809 on F508del-CFTR in primary HBE cells has been estimated at 3 μM (F. Van Goor, personal communication, 2011). Such concentrations of VX-809 were observed in the plasma of patients in the 100 and 200 mg dose cohorts at steady state. However, the drug exposure levels achieved in patient target tissues at these doses of VX-809 is not known.

The other biomarkers of CFTR activity (NPD, immunoblot from rectal biopsy tissue) failed to demonstrate changes over the course of VX-809 treatment. This may reflect the trial’s lack of power to detect a predicted change in these measurements based on in vitro results with VX-809 in F508del/F508del primary HBE cells (maximum of ~15% of wild-type CFTR function). The intrinsically greater variability of NPD measurement (potentially compounded by the large number of NPD sites used in this study) may have contributed to the lack of detectable changes in this study. The results provide support for the use of previously obtained CFTR biomarker data sets for study planning of new CFTR modulators. Using this paradigm, the treatment effects on sweat chloride reported with the CFTR potentiator VX-770 in patients with CF with the G551D-CFTR mutation were successfully used to power the current study to detect improvements in sweat chloride (within-group comparisons and with placebo). It is not clear if the reductions in sweat chloride in this study achieved a peak effect (figure 1), and this raises the question of whether higher doses of VX-809 may produce greater effects on F508del-CFTR. The peak corrective effect of VX-809 on F508del-CFTR in primary HBE cells has been estimated at 3 μM (F. Van Goor, personal communication, 2011). Such concentrations of VX-809 were observed in the plasma of patients in the 100 and 200 mg dose cohorts at steady state. However, the drug exposure levels achieved in patient target tissues at these doses of VX-809 is not known.

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been shown to suppress PTCs and improve NPD parameters in
and colleagues treated a study. Using a separate CFTR modulator strategy, Rubenstein
nantly limited to a subset of patients with the Y122X muta-
two pilot studies, but effects on sweat chloride were predomi-
function and cough frequency were not observed until months
responsiveness to VX-809 treatment.

PTC124 has been shown to have detectable bioactivity by NPD
termination codons in CFTR, while sweat chloride
improvements in NPD-dependent chloride secretion were seen,
but there were no effects on sweat chloride. The reasons for
discrepancies in CFTR biomarkers across different CFTR
modulator strategies are not clear, but suggest that organ effects
may vary due to tissue drug availability, CFTR regulation in
different cell types or the responsiveness of mutant CFTR across
different tissue compartments and modulator strategies. These
uncertainties should be considered in future study planning, as
should the selection and continued development of CFTR
biomarkers to demonstrate study drug bioactivity.

In conclusion, the results provide support for the continued
evaluation of VX-809 in patients with CF with the F508del-CFTR
mutation. This study demonstrated that VX-809 has
an acceptable safety profile. It also provided evidence of the effect
of VX-809 on improving CFTR function based on dose-depen-
dent reductions in F508del-CFTR activity in the sweat gland
using a CFTR corrector administered orally to subjects with CF,
and further investigation is warranted to evaluate the potential
of this therapeutic strategy to increase F508del-CFTR activity.

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No improvements in clinical outcomes, including lung function
and quality of life (CFQ-R), were observed over the course of this
study. The trial was not powered to detect such improvements in
these measures, and longer or larger trials of VX-809 (alone or
in combination with CFTR potentiators) may be necessary to
determine the clinical effect of these CFTR modulators.

Previous trials of systemically dosed CFTR modulators have
frequently described discordant effects on CFTR-dependent
biomarkers and pulmonary outcome measures. For example,
PTC124 has been shown to have detectable bioactivity by NPD
over 2 weeks of treatment in patients with CF possessing
premature termination codons in CFTR, while sweat chloride
measurements remained unchanged. Improvements in lung
function and cough frequency were not observed until months
of treatment were completed. Systemic gentamicin has also
been shown to suppress PTCs and improve NPD parameters in
two pilot studies, but effects on sweat chloride were predomi-
nantly limited to a subset of patients with the Y122X muta-
tion. Using a separate CFTR modulator strategy, Rubenstein
and colleagues treated F508del-CFTR homozygous patients with
CF with the F508del-CFTR modulator 4-phenyl butyrate.
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This article has been corrected since it was published Online First. The author name Kris DeBoeck has been corrected to Kris DeBoeck.

Competing interests
None.

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This study was conducted with the approval of the Institutional Review Board at each site.

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REFERENCES
Results of a phase IIa study of VX-809, an investigational CFTR corrector compound, in subjects with cystic fibrosis homozygous for the F508del-CFTR mutation


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