**AN EARLY HEALTH ECONOMIC ANALYSIS OF THE POTENTIAL COST-EFFECTIVENESS OF AN ADHERENCE INTERVENTION TO IMPROVE OUTCOMES FOR PATIENTS WITH CYSTIC FIBROSIS**

**Short title: Cost-effectiveness of CF adherence intervention**

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**ABSTRACT**

**Background:** Cystic fibrosis (CF) negatively impacts upon health-related quality of life (HRQoL) and survival. Adherence to nebulised treatments is low; improving adherence is hypothesised to reduce rates of exacerbation requiring intravenous antibiotics and lung function decline.

**Methods:** A state transition model was developed to assess the cost-effectiveness of an intervention aimed at increasing patient adherence to nebulised and inhaled antibiotics compared with current CF care, in advance of the forthcoming CFHealthHub randomised controlled trial (RCT). The model estimates the costs and health outcomes for each option from the perspective of the National Health Service (NHS) and Personal Social Services (PSS) over a lifetime horizon. Health gains are valued in terms of quality-adjusted life years (QALYs) gained. Forced Expiratory Volume in 1 Second (FEV1) trajectories are predicted over three lung function strata: (i) FEV1 ≥70%; (ii) FEV1 40-69%, and; (iii) FEV1 <40%. Additional states are included to represent post-lung transplantation and dead. The model was populated using CF Registry data, literature and expert opinion. Costs were valued at 2016 prices. Uncertainty was assessed using deterministic and probabilistic sensitivity analyses.

**Results:** The adherence intervention is expected to produce an additional 0.19 QALYs and cost savings of £64,078 per patient. Across all analyses, the intervention dominated current care. Over a 5-year period, the intervention is expected to generate cost savings of £49.5million for the estimated 2,979 CF patients with *Pseudomonas aeruginosa* currently aged 16 or above in the UK. If applied to a broader population of adult CF patients receiving any nebulised therapy, the expected savings could be considerably greater.

**Conclusions:** The adherence intervention is expected to produce additional health gains at a lower cost than current CF care. The economic analysis should however be revisited upon completion of the full RCT. More generally, the analysis suggests that considerable gains could be accrued through the implementation of adherence interventions which shift care from expensive hospital-based rescue to community-based prevention.

**KEY POINTS FOR DECISION-MAKERS**

* Adherence to nebulised treatments for CF is low. CF patients with poor adherence have significantly higher healthcare costs than those with good adherence; most of the excess costs in poor adherers are related to hospital admission for i.v. rescue therapy to treat pulmonary exacerbations.
* Based on a pre-trial analysis of the CFHealthHub study, the use of an adherence intervention for CF is expected to produce an additional 0.19 QALYs and cost savings of £64,078 per patient compared with current CF care. Over a 5-year period, this corresponds to cost savings of approximately £49.5million for the estimated 2,979 CF patients with *Pseudomonas aeruginosa* currently aged 16 or above in the UK.
* If the adherence intervention benefits a broader population of CF patients who are receiving nebulised antibiotics and/or mucolytics and are aged 16 or over (likely to represent approximately 5,800 patients), the 5-year cost savings to the NHS are expected to be in excess of £96million. Given existing uncertainty, it will be important to revisit this economic analysis upon completion of the full CFHealthHub RCT.

**1. INTRODUCTION**

Cystic fibrosis (CF) is an inherited condition characterised by the abnormal transport of chloride ions (Cl-) across transporting epithelia. This leads to the production of thick sticky mucus in the lungs, pancreas, liver, intestine, and reproductive tract, and an increase in the salt content in sweat. More than 10,000 children and adults in the UK have CF [1]. Whilst CF limits life expectancy, survival is increasing, and in 2015, approximately 6,475 CF sufferers in the UK were older than 16 years of age. Amongst other problems, for example poor digestion, patients with CF are susceptible to lung infections, in particular, *Pseudomonas aeruginosa*. This is thought to be because the thick mucus makes it difficult for the body to clear inhaled bacteria, and because people with CF have an increased airway inflammatory response to pathogens. People with CF often develop intermittent infections during childhood, which can be treated and even eradicated with nebulised or inhaled antibiotics. As infection develops, however, patients may reach a chronic stage whereby eradication is no longer possible due to the formation of biofilms. In such cases, ongoing inhaled antibiotic treatment must be continued permanently. Patients commonly also require other treatments including inhaled mucolytics, bronchodilators, steroids, and physiotherapy. Treatment is time consuming and burdensome, with administration of nebulised antibiotics taking up to an hour per day whilst patients are well and longer during periods of ill health. In addition, patients also experience pulmonary exacerbations which require treatment with intravenous (i.v.) antibiotics which are administered either at home or in hospital. In either case, this treatment compromises the patient’s ability to attend school or work and leads to increased health care costs.

Adherence to preventative nebulised CF treatments is estimated at 48%, based on medication possession ratio (MPR) data in 3,287 people with CF aged 6 years and above in the US [2]. This is a measure of persistent adherence as these are chronic medications which are taken long-term [3]. These data show that adherence is inversely related to age. Importantly, MPR measures only medication that is collected from the pharmacy and cannot measure whether that medication is ever taken. Objective UK data using chipped nebulisers in adults goes beyond the coarse MPR adherence metric to objectively identify how much treatment is actually taken. These data suggest a lower median adherence rate of 36% in nebulisers that are brought to clinic to be downloaded [4]. It is also important to ensure that estimated adherence rates take into account the patient’s clinical status (“normative adherence”) and includes both nebulisers brought to clinic and the more difficult to obtain nebulisers left at home; UK data reported by Hoo *et al* suggest that when all nebulised devices are included, normative adherence in adult clinics may be as low as 33% [5].

Treatments only work if they are taken, and estimates of drug effectiveness are usually derived from randomised controlled trials (RCTs) whereby strict inclusion criteria and trial procedures typically produce high adherence rates. A recent review of adherence to CF treatments within clinical trials by Pugatsch *et al* [6] reported an adherence rate of 80% as an aggregate estimate across the population over the entire duration of the studies (mean duration 7.3 months; range 2-24 months). The methodology for adherence measurement varied between trials, but was typically undertaken at frequent intervals across the trials and can be considered as a measure of persistent adherence [3]. Although there was a tendency for some reduction over time, adherence remained high across all phases of follow-up. Based on MPR data, it has been shown that CF patients with poor adherence (MPR <50%) have significantly higher healthcare costs than patients with good adherence (MPR>80%); most of the excess costs in poor adherers are related to hospital admission for i.v. rescue therapy to treat pulmonary exacerbations [2]. A meta-analysis reported by Demonceau *et al* assessed feedback of objective adherence data in various conditions across 5,237 patients; this study demonstrated that such feedback could increase adherence by around 20%, with a further 8% improvement if simple problem solving was added [7]. The ongoing CFHealthHub ACtiF trial (Adherence to treatment in adults with Cystic Fibrosis) is currently assessing an intervention to improve outcomes for CF patients by empowering self-management and improving adherence to nebulised therapy via the use of chipped nebulisers which have the capacity to directly monitor adherence levels, combined with an intervention to support problem solving, habit formation and self-efficacy undertaken by a member of the multidisciplinary team at regular appointments [8]. The hypothesis underpinning this RCT is that increasing adherence will reduce the number of exacerbations experienced by CF patients. Reducing the number of exacerbations experienced by CF patients may also impact on their rate of lung function deterioration. This paper presents an early health economic evaluation of the expected cost-effectiveness of the adherence intervention compared with current clinical care, in advance of the completion of the full CFHealthHub ACtiF RCT. More generally, this analysis provides a formal quantification of the potential gains which could be accrued by improving disease control in CF through improved adherence.

**2. METHODS**

**2.1 Scope of the analysis**

A model-based cost-utility analysis was undertaken to assess the incremental cost-effectiveness of the adherence intervention versus standard care in adult CF patients receiving traditional nebulised or dry powder inhaled (DPI) antibiotics from the perspective of the National Health Service (NHS) and Personal Social Services (PSS) over a lifetime horizon. The intervention relates to a newly-developed nebuliser with the capacity to monitor and report adherence levels (developed as part of the CFHealthHub ACtiF programme [8]), combined with a behavioural intervention undertaken by a physiotherapist at regular appointments. Health benefits are assessed in terms of quality-adjusted life years (QALYs) gained. All costs and health outcomes were discounted at a rate of 3.5% per annum [9]. Costs were valued at 2016 prices.

**2.2 Model structure**

The analysis uses a state transition approach, based on the structure of a previously published health economic model developed to assess the cost-effectiveness of DPIs for CF patients with chronic *Pseudomonas aeruginosa* [10;11]. The model was populated using analyses of individual patient data (IPD) from the UK CF Registry [12], literature and expert opinion. The model estimates Forced Expiratory Volume in 1 Second (FEV1) percent predicted trajectories over three strata of lung function: (1) FEV1 ≥70%; (2) FEV1 40-69%, and; (3) FEV1 <40% (Figure 1). Additional health states are included to represent post-lung transplantation and dead. During each annual cycle, patients may remain in their current FEV1 state, transit to an improved or worsened FEV1 state or die. A small proportion of patients with FEV1<40% may undergo lung transplantation and do not subsequently receive further nebulised/DPI treatment. HRQoL is modelled according to FEV1 stratum and transplant history, with disutilities applied according to the proportion of time spent receiving i.v. antibiotics to manage CF exacerbations. A half-cycle correction is applied to account for the timing of events. Total QALYs are calculated as the total sojourn time in each health state weighted by state-specific utility scores, less any QALY losses resulting from exacerbations. The model conservatively assumes that there is no survival difference between the intervention and comparator groups. Costs include UK CF tariff treatment costs, high-cost drug acquisition, costs of i.v. days spent in hospital and associated i.v. antibiotic acquisition, transplantation costs and costs associated with the adherence intervention. All other costs are assumed to be captured in the CF banding tariff [13].

The model employs the following assumptions:

* Patients in any FEV1 stratum can progress/regress to any other FEV1 stratum
* Reductions in exacerbations impact upon progression rates between FEV1 strata
* The probability of experiencing exacerbations differs by FEV1 stratum
* A small proportion of patients with FEV1 <40% undergo lung transplant, whilst those ineligible for transplant continue to receive nebulised/DPI therapy
* Reductions in lung function and the incidence of exacerbations impact upon HRQoL
* The adherence intervention will impact upon the incidence of exacerbations and FEV1 transitions
* Exacerbation rates and transition rates between FEV1 strata are time-invariant
* The costs of “high-cost therapies” is independent of adherence to those therapies.

Figure 1: Model structure

[INSERT FIGURE 1 HERE]

**2.3 Evidence used to inform the model parameters**

Model parameters and their associated distributional properties are summarised in Table 1.

**Table 1: Summary of model parameters**

[INSERT TABLE 1 HERE]

*Patient characteristics*

The population is assumed to be 16 years of age at model entry. The initial distribution of patients across the health states was based on recorded FEV1 in 2013 from the CF Registry (the most recent year available for the analysis).

*Transition probabilities*

Transition probabilities were derived from IPD from the CF Registry for a total of 10,344 patients between 2007 and 2013 [12]. The overall dataset was restricted to those patients who have been recorded (at least once) as having ‘intermittent’ or ‘chronic’ *Pseudomonas aeruginosa* status (n=7,518). Fifty-three percent of these patients were male, and their average age on first appearing in the 2007-2013 dataset was 19 years (range 0-82 years). At baseline, the proportion of patients in each FEV1 strata was as follows: FEV1<40%=0.13; FEV1 40-69%=0.32; FEV1>70%=0.55). Of these, 6,788 had at least one recorded FEV1 assessment and 1,700 had measures for all seven years. Longitudinal regression was undertaken using the methods described by Jung *et al* [14]; this involved the estimation of a series of ordered logit models which give the log odds of being in a given FEV1 group post-transition, given the time (in days) between observations, the annual rate of hospital i.v. days and the patient’s age. One model was estimated for each of three possible FEV1 starting states and the model outputs were converted into annual probabilities. This approach allowed for the inclusion of patients who had between 2 and 7 entries in the registry, even if there were gaps before, between, or after review entries. Patients who left the registry were excluded from the analysis. The time variable (days since last visit) allows for the calculation of annual transitions despite the fact that sample intervals varied in the raw registry data. The lagged rate of i.v. days variable allowed for the investigation of the effect of exacerbations on FEV1 progression. A number of models were tested with both age and sex as covariates. Age was statistically significant in most models, but sex was not significant, therefore age was retained in the final model.

*Mean days on i.v. antibiotics per year*

Mean days in hospital or at home receiving i.v. antibiotics for each FEV1 group were estimated from the same group of patients with *Pseudomonas aeruginosa* in the CF Registry used to derive transition probabilities [12].

*Effectiveness of the adherence intervention*

The intervention is assumed to reduce the number of days on i.v. antibiotics, leading to changes in the transition rates between the FEV1 strata. The CFHealthHub ACtiF trial is powered to detect a reduction of one exacerbation per annum, based on a previous trial of long-term inhaled hypertonic saline for CF (Elkins *et al [15]*); this treatment effect is assumed to reflect the minimum clinically important difference. Assuming one exacerbation is equivalent to 14 days of home or hospital i.v. antibiotics, when applied to the whole CF Registry cohort with *Pseudomonas aeruginosa*, this equates to a 55% reduction in days spent receiving i.v. antibiotics. Uncertainty surrounding the relative risk reduction in i.v. days was assumed to broadly reflect that observed in the Elkins trial, but the 95% confidence interval [CI] was widened to account for additional uncertainty surrounding the effectiveness of the adherence intervention (mean relative risk=0.45, standard error=0.09). Post-intervention exacerbation rates were also applied to the logit models to derive FEV1 transition probabilities for the adherence intervention.

*CF mortality*

The CF Registry does not include sufficient data to allow for the robust derivation of estimates of long-term survival for CF patients [12]. Instead, survival estimates were based on an analysis reported by Dodge *et al* [16]. This study reported survival data up to the end of 2003 for all subjects with CF born in the UK in the period 1968-1992 collated via active enquiry of CF clinics and other hospital consultants. The published survival curves for males and females were digitised and patient-level time-to-event data were reconstructed using methods reported by Guyot *et al* [17]. Parametric survivor functions (exponential, log normal, log logistic, Weibull, Gompertz and generalised gamma) were fitted to the replicated data to extrapolate beyond the observed follow-up period. Model discrimination was undertaken using visual inspection, an examination of the goodness-of-fit statistics for each survivor function (the Akaike Information criterion [AIC] and the Bayesian Information Criterion [BIC]), together with subjective clinical judgement regarding the plausibility of the extrapolated portion of each parametric curve. On the basis of clinical plausibility, the Gompertz survivor function was selected for use in the model. Uncertainty surrounding the parameters of the survivor function was modelled using independent normal distributions with the 95% CI width calibrated such that it was similar to that observed for patients in the current CF Registry population. The same function was applied to the intervention and comparator groups, hence the adherence intervention is not assumed to impact on patient survival.

*Probability of transplantation*

The probability that a patient with FEV1 <40% will undergo a lung transplant during each cycle was estimated based on data from the UK CF Registry [1] and the US Cystic Fibrosis Foundation, assuming a 2-3% lifetime probability of undergoing lung transplantation [10;11].

*HRQoL*

The selection of studies used to inform HRQoL parameters within the model was based on a previous systematic review [9]. Health state utilities associated with each FEV1 stratum and the disutility associated with exacerbations were based on a utility valuation study reported by Bradley *et al*;[18] within this study, the Euroqol-5 Dimensions (EQ-5D) and the Cystic Fibrosis Questionnaire-Revised (CFQ-R) were administered to patients aged ≥16 years with CF and chronic *Pseudomonas aeruginosa* and who were taking nebulised or oral antibiotics. The utility score for patients who have undergone lung transplantation was taken from Anyanwu [19].

*Resource costs*

The model includes costs associated with CF tariff banding, high cost antibiotic therapies, hospital i.v. days, i.v. antibiotics, transplantation and the adherence intervention. The proportion of patients in each band of the CF tariff according to FEV1 stratum were derived from the CF Registry [12]; banding tariff costs were taken from the latest NHS National Tariff [13]. Usage of specific antibiotic products was estimated from the CF Registry dataset [12]. The analysis assumes that patients are prescribed these treatments according to their licensed dosing schedules rather than according to patient adherence levels. Unit costs for nebulised and DPI antibiotics were derived from British National Formulary (BNF) 2016 [20]. Transplantation costs were based on personal communication (Kim Cox, NHS England). The costs of i.v. antibiotics (tobramycin and ceftazidime) were sourced from Sheffield Teaching Hospitals (personal communication: Tim Gleeson, STH). The cost of a hospital i.v. day was based on NHS Reference Costs 2014/15; as there is no inpatient cost relating to CF exacerbations, the daily cost associated with a long stay inpatient admission for bronchiectasis with complications and comorbidity score 0 was assumed (daily cost = £361.68) [21]. The model assumes that 54% i.v. days take place in hospital; the remaining 46% are assumed to take place at the patient’s home and do not lead to additional costs for the NHS. The costs of the adherence intervention were assumed to include a once-only cost for data transfer hardware of £121.20 plus an ongoing annual data transfer cost of £583.44 per patient. The analysis assumes that the training and implementation costs associated with the behavioural impact component of the adherence intervention would be absorbed into routine clinic appointments undertaken by CF healthcare practitioners and therefore these costs have been excluded.

*Model evaluation methods*

Cost-effectiveness was expressed in terms of the incremental cost per QALY gained. Uncertainty surrounding the cost-effectiveness of the adherence intervention was explored using deterministic sensitivity analysis (DSA) and probabilistic sensitivity analysis (PSA). The PSA included all uncertain model parameters and was implemented using simple Monte Carlo sampling methods. DSAs were conducted to explore the impact of alternative time horizons and shorter durations of treatment effect, assuming that the intervention impacts on exacerbation rates only, reduced impacts on exacerbation rates, and altering assumptions regarding cost and utility parameters. In addition, a further scenario was conducted whereby treatment costs were calculated according to expected adherence levels in each group, based on Daniels *et al* [4] (36% drug consumption in current CF care group) and Demonceau *et al* [7](63.7% drug consumption for the adherence intervention group).

*Model validation methods*

Several measures were taken to verify the implemented model and to the ensure the credibility of its underlying conceptual basis. These included internal peer review by clinical experts, scrutiny of the implemented model coding and formulae, checking the accuracy of all model inputs against sources, investigating potentially discrepant or unexpected results identified through black box testing and double-programming of the deterministic model. Whilst the results of the CFHealthHub ACtiF trial will not be available until at least September 2019, once available, this evidence will allow the economic analysis to be updated using prospectively collected randomised data; this will also enable the comparison of the predictions of this pre-trial and post-trial analyses.

**3. RESULTS**

**3.1 Central estimates of cost-effectiveness**

Table 2 presents the central estimates of cost-effectiveness for the adherence intervention versus current CF care based on the probabilistic version of the model.

**Table 2: Central estimates of cost-effectiveness (probabilistic)**

[INSERT TABLE 2 HERE]

The probabilistic version of the model suggests that the adherence intervention is expected to produce an additional 0.19 discounted QALYs per patient and cost-savings of approximately £64,078 per patient over their remaining lifetime; hence, the adherence intervention is expected to dominate current care. The cost savings predicted by the model are driven by a small shift in CF banding resulting from improvements in predicted FEV1 trajectory, together with a significant reduction in the expected number of hospital i.v. days (accounting for savings of approximately £70,000 per patient). As shown in the cost-effectiveness plane (Figure 2), whilst there is considerable uncertainty surrounding the health gains associated with the intervention, the probabilistic analysis consistently indicates that the adherence intervention is expected to produce substantial cost savings. Across willingness-to-pay thresholds of between £0 and £100,000 per QALY gained, the probability that the adherence intervention produces more net benefit than current care is expected to be 1.0.

**3.2 Sensitivity analysis results**

Across all of the DSAs, the adherence intervention is expected to dominate current care (Table 3). This includes the highly pessimistic situation whereby the costs of high cost drugs calculated exactly according to the level patient adherence, based on Daniels *et al* [4] and Demonceau *et al* [7]. Even in this unlikely scenario, the savings associated with costs of i.v. hospital days avoided outweigh the additional costs of drug therapy due to increased patient adherence to nebulised and inhaled therapy. Assuming that treatment costs are independent of adherence levels, the sensitivity analysis suggests that over the course of 5-years, the model estimates discounted cost savings of £16,623 per patient; this is equivalent to approximately £49.5million for the estimated 2,979 CF patients with *Pseudomonas aeruginosa* currently aged 16 or above in the UK. Should the intervention benefit a broader population of CF patients who are receiving nebulised antibiotics and/or mucolytics and are aged 16 or over (likely to represent approximately 5,800 patients), the 5-year cost savings to the NHS are expected to be in excess of £96million.

**Table 3: Deterministic sensitivity analysis results**

[INSERT TABLE 3 HERE]

**Figure 2: Cost-effectiveness plane**

[INSERT TABLE 3 HERE]

**4. DISCUSSION**

This study represents the first health economic analysis of an intervention targeted at increasing patient adherence to nebulised/DPI treatments in CF. The results of the analysis suggest that the adherence intervention has the potential to produce considerable health gains as well as cost savings for the NHS, thereby dominating current CF care. The principal driver of the anticipated cost savings is due to the expected reduction in hospital i.v. days. The sensitivity analysis suggests that even under pessimistic assumptions regarding lower levels of effectiveness of the intervention and lower unit costs per i.v. day, the adherence intervention is expected to remain dominant. This suggests that even if the CFHealthHub ACtiF trial does not meet its primary endpoint, the intervention may still produce cost savings for the health service.

Since this health economic analysis precedes the CFHealthHub ACtiF trial, there is considerable uncertainty regarding whether the findings of the analysis will concord with the data that will be collected within the trial itself. Invariably, such early modelling analyses are subject to the risk of reaching erroneous conclusions and rely on a weaker evidence base than would be available had the full trial been completed. It is therefore important to consider these evidential issues in the interpretation of the results of this analysis; these limitations are discussed briefly below.

*(i) Clinical evidence to support the effectiveness of the adherence intervention*

The most pertinent limitation of the evidence base is that the CFHealthHub ACtiF trial, which aims to assess the clinical benefit of the adherence intervention, has not yet begun. As such, there is currently no direct empirical evidence through which to quantify the benefits of the adherence intervention. Given this lack of evidence, the model uses the expected reduction in exacerbations used to inform the power calculations for the CFHealthHub ACtiF trial [8;15] as the basis for modelling expected treatment effects. Whilst this estimate reflects a legitimate prior belief, and forms the basis of the hypothesis that will be tested within the trial, there is a possibility that the anticipated reduction in exacerbations could be higher or lower than predicted. Nonetheless, the economic analysis presented here has a wider relevance in quantifying the potential gains that could be accrued through the implementation of adherence interventions which shift care from expensive hospital-based rescue to more economical community-based prevention. This analysis may therefore be useful in supporting the development and evaluation of other adherence interventions within the NHS or across other health care systems.

*(ii) Transition probabilities and exacerbation rates are assumed to apply indefinitely*

The model uses a single matrix of probabilities describing the trajectories of lung function across three FEV1 strata in each treatment group. Whilst age is included as a covariate in the logit regression analysis, these are treated as time-independent parameters within the health economic model. In reality, FEV1 transitions may be time-variant. Owing to the absence of long-term data on FEV1 trajectories with and without the intervention, the model assumes that these trajectories remain constant with respect to time. It should be noted however, that the economic conclusions drawn from the analysis remain unchanged even if the intervention has no impact upon lung function decline (Table 3).

*(iii) Treatment effect assumed to apply indefinitely*

Given the preliminary nature of the health economic analysis and the current lack of evidence regarding the effectiveness of the adherence intervention, the model assumes that the treatment effect applies indefinitely over the patient’s remaining lifetime. It may be the case that levels of adherence to antibiotic therapies may increase or wane over time following the introduction of the intervention. The sensitivity analysis suggests that both health gains and cost savings are expected to be reduced over shorter intervals, although the intervention is expected to remain dominant irrespective of the time horizon.

*(iv) Limitations in handling cost savings due to the CF banding tariff*

Within the model, benefits and costs are captured through two different processes: (i) a direct reduction in the number of days spent receiving i.v. antibiotics, and; (ii) the impact of reduced exacerbations on subsequent FEV1 trajectory. In England, CF care is currently funded via a mandatory tariff for specialist commissioning which is intended to reflect the severity of disease in individual patients; the UK CF banding tariff is intended to encapsulate both lung function and i.v. days. Consequently, an analysis which accounts only for changes in the CF tariff band would fail to fully reflect cost savings realised by the NHS due to fewer i.v. days spent in hospital; this would lead to an underestimate of the true cost-effectiveness of the adherence intervention. Alternatively, an analysis which includes CF banding costs as well as i.v. hospital costs, as has been assumed here, may overestimate the cost-savings associated with reducing exacerbations. It is anticipated that the data collection mechanisms within the CFHealthHub ACtiF trial will allow for a more sensitive and accurate analysis of the true costs associated with the adherence intervention based on the direct modelling of FEV1 status and CF banding categories.

*(v) Relationship between treatment costs and patient adherence*

There is uncertainty regarding the relationship between the costs of treatment and patient adherence to those treatments. The base case analysis assumes that antibiotic treatment costs borne by the NHS are independent of patient adherence levels. It is possible that increasing adherence levels will also lead to increases in total NHS expenditure on antibiotic treatments: as patients become more adherent to therapy, they may require more frequent prescriptions. The consequence of this situation would be an increase in the total CF drugs bill, and the expected cost savings of increased adherence would be somewhat diminished. However, the sensitivity analyses indicate that even in the presence of very pessimistic assumptions regarding the relationship between treatment adherence and treatment costs, specifically a scenario whereby treatment costs borne by the NHS exactly reflect patient consumption of those treatments, the adherence intervention is expected to remain cost-saving (Table 3).

**5. CONCLUSIONS**

Based on an early health economic analysis using high quality registry data [1;12] and the estimated reduction in exacerbations used to inform the design of the CFHealthHub ACtiF trial [8], the adherence intervention is expected to produce additional health gains at a substantially lower cost than current CF care. The findings of the analysis should be revisited upon the completion of the full RCT. More broadly, the analysis suggests that considerable gains could be accrued through the implementation of adherence interventions which shift care from hospital-based rescue to community-based prevention.

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**AUTHOR CONTRIBUTIONS**

Paul Tappenden developed the health economic model. Susannah Sadler undertook the analyses of the CF Registry dataset. Martin Wildman advised on the design of the study and the evidence used to inform the model. All authors contributed to the preparation of this manuscript. Paul Tappenden will act as the overall guarantor for this work.

**COMPLIANCE WITH ETHICAL STANDARDS**

Dr Wildman has received support from Pari to speak at conferences about the importance of adherence. Dr Wildman has also received funding from Pari to travel to meetings with Pari about setting up a trial to understand whether increasing adherence improves outcomes in CF. Dr Wildman has received funding from Philips to support research using the Ineb nebuliser to understand how the device can be used to measure adherence. Dr Wildman has received speaker fees from Forrest to give independent talks at CF meetings around the UK about the importance of adherence. Paul Tappenden and Susannah Sadler have no conflicts of interest to declare.

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