

# Does co-payment for inhaler devices affect therapy adherence and disease outcomes? A historical, matched cohort study

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**Background:** Adherence to asthma and chronic obstructive pulmonary disease (COPD) treatment has been shown to depend on patient-level factors, such as disease severity, and medication-level factors, such as complexity. However, little is known about the impact of prescription charges – a factor at the health care system level. This study used real-life data to investigate whether co-payment affects adherence (implementation and persistence) and disease outcomes in patients with asthma or COPD.

**Methods:** A matched, historical cohort study was carried out using two UK primary care databases. The exposure was co-payment for prescriptions, which is required for most patients in England but not in Scotland. Two comparison cohorts were formed: one comprising patients registered at general practices in England and the other comprising patients registered in Scotland. Patients aged 20–59 years with asthma, or 40–59 years with COPD, who were initiated on fluticasone propionate/salmeterol xinafoate, were included, matched to patients in the opposite cohort, and followed up for 1 year following fluticasone propionate/salmeterol xinafoate initiation. The primary outcome was good adherence, defined as medication possession ratio  $\geq 80\%$ , and was analyzed using conditional logistic regression. Secondary outcomes included exacerbation rate.

**Results:** There were 1,640 patients in the payment cohort, ie, England (1,378 patients with asthma and 262 patients with COPD) and 619 patients in the no-payment cohort, ie, Scotland (512 patients with asthma and 107 patients with COPD). The proportion of patients with good adherence was 34.3% and 34.9% in the payment and no-payment cohorts, respectively, across both disease groups. In a multivariable model, no difference in odds of good adherence was found between the cohorts (odds ratio, 1.04; 95% confidence interval, 0.85–1.27). There was also no difference in exacerbation rate.

**Conclusion:** There was no difference in adherence between matched patients registered in England and Scotland, suggesting that prescription charges do not have an impact on adherence to treatment.

**Keywords:** implementation, adherence, asthma, chronic obstructive pulmonary disease, prescriptions, co-payment

## Background

Asthma and chronic obstructive pulmonary disease (COPD) are two leading causes of morbidity and economic burden worldwide, with 300 and 210 million people affected, respectively.<sup>1</sup> Both these diseases are chronic obstructive lung conditions principally managed with inhaled therapies, such as inhaled corticosteroids (ICS). Effectiveness of therapy, however, can be adversely affected by poor adherence.<sup>2</sup> Adherence is defined as the process by which patients take their medications as prescribed, comprising

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1) initiation of treatment, 2) implementation of the dosing regimen, and 3) persistence with treatment.<sup>3</sup> Previous literature has demonstrated that poor adherence can exist among patients with asthma or COPD<sup>4–6</sup> and has an impact on disease outcomes.<sup>7–9</sup> Contributing factors to poor adherence at the patient level include perceived need of treatment,<sup>10</sup> fear of side effects,<sup>11</sup> and ability to use inhaler devices correctly.<sup>12,13</sup> Less research has been done into contributing factors at the health care system level.

One potential factor is co-payment for prescriptions. In an attempt to scale back on pharmaceutical spending, many countries have introduced co-payment policies to offset growing drug bills.<sup>14–16</sup> A co-payment is a fixed fee for a prescription, which is intended to reduce drug expenditure and the collection of medications that have no role in improving health.<sup>17</sup> However, co-payments may be disadvantageous if they result in a decrease in the use of beneficial medication. A study in Australia found that a 24% increase in patient co-payment for medications had significant effects on the dispensing of some essential drugs, including decreased volumes of dispensed asthma medications.<sup>18</sup> The more vulnerable patient populations are those that have an increased sensitivity to adverse health outcomes, such as the elderly and those with low socioeconomic status.<sup>19</sup> A review of 22 studies in the USA between 1990 and 2003 found that co-payments were associated with a decrease in the health status of older patient groups.<sup>20</sup>

Currently, in England, for residents nonexempt from paying for prescriptions, there is a flat rate patient contribution of £8.40 per prescription.<sup>21</sup> A resident of England can cap the prescription cost by purchasing a monthly (£29) or annual (£104) prescription prepayment certificate (PPC). In contrast, Scotland canceled co-payments on prescriptions in 2011.<sup>22</sup> This difference in prescription payments in England and Scotland, and the availability of real-life primary care data over time, allows an opportunity to examine the effects of prescription co-payment.

The aim of this study was to use real-life data to examine the impact of co-payment on maintenance therapy implementation and persistence and disease outcomes in patients with asthma or COPD, accounting for differences between the populations.

## Methods

### Data source and permissions

This study used data from the Optimum Patient Care Research Database (OPCRD)<sup>23</sup> and the Clinical Practice Research Datalink (CPRD).<sup>24</sup> Both are primary care databases with routinely recorded clinical data from general practices in

the UK. The OPCRd has anonymized data from >3 million patients in >570 UK practices, and the CPRD has anonymized data from >5 million patients in >600 UK practices. The databases contain a range of information, including prescribing activity and diagnoses of disease, identifiable by Read codes recorded by clinicians. The CPRD (formerly GPRD) is well validated and used frequently for medical and health research.<sup>24,25</sup> The OPCRd has been used in numerous studies of real-life effectiveness in respiratory diseases;<sup>26–29</sup> the validity of the real-life endpoints commonly identified in the OPCRd (such as risk-domain asthma control) has been supported previously in a study showing the predictive value of real-life endpoints for future patient outcomes.<sup>30</sup>

The OPCRd has ethical approval from the National Health Service (NHS) Research Authority to hold and process anonymized research data (Research Ethics Committee reference: 15/EM/0150). This study was approved by the Anonymised Data Ethics Protocols and Transparency (ADEPT) committee – the independent scientific advisory committee for the OPCRd, commissioned by the Respiratory Effectiveness Group.<sup>31</sup> Informed consent was not required from the participants, as this was a retrospective study, and all data was anonymized. The study is registered with the European Network of Centres for Pharmacoepidemiology and Pharmacovigilance (ENCEPP/SDPP/13586).<sup>32</sup>

### Study design

A historical, matched cohort study design was used to compare patients with asthma or COPD with likely co-payment with patients with asthma or COPD with no co-payment for prescriptions, in terms of their adherence to fluticasone propionate/salmeterol xinafoate (FP/SAL) therapy. We chose FP/SAL as we aimed to examine adherence to maintenance therapy only and so did not select medications that are used as both maintenance and reliever therapy. The cohorts will be referred to as the payment cohort (patients from general practices in England) and the no-payment cohort (patients from general practices in Scotland). The study period consisted of 2 years – a baseline year for patient characterization and identification of suitable matching variables and an outcome year for observing the outcomes of interest. The baseline and outcome years were separated by an index date, which was the date of first prescription of FP/SAL.

### Inclusion and exclusion criteria

Patients were selected based on the following inclusion criteria: diagnosis for asthma and/or COPD; received prescription for FP/SAL, with no prescription in the preceding year; aged ≥40 and <60 years at the index date (date of initial

prescription of FP/SAL) for patients with COPD, and aged  $\geq 20$  and  $< 60$  years at the index date for patients with asthma; at least 2 years of valid, continuous data (at least 1 year before and at least 1 year after the index date); and at least one further prescription for FP/SAL during the outcome period. The upper age boundary of 60 years was used because patients aged  $\geq 60$  years are exempt for co-payment in England.

Patients were excluded if they had any chronic respiratory disease other than asthma or COPD, at any time, or they met other conditions for co-payment exemption, which included being pregnant during or in the 12 months prior to the study period, record of “prescription exemption status” before or during the study period, and hospitalization during the study period.

Patients with only an asthma diagnosis were considered the “asthma group”, and patients with a COPD diagnosis, who may or may not have had an asthma diagnosis in addition, were considered the “COPD group”. Thus, there was no overlap between the two groups.

## Outcome measures

The primary outcome was adherence to FP/SAL therapy over the 1-year outcome period. This was evaluated in terms of the percentage of prescription refill rate, with good adherence being defined as a medication possession ratio (MPR)<sup>33</sup> of  $\geq 80\%$ . The normal duration of a prescription in the UK is 30 days. Since the MPR was assessed in a fixed 1-year period, this adherence measure incorporates both implementation and persistence. Details of the calculations used to compute MPR can be found in the [Supplementary materials](#).

Secondary outcome measures included number of severe exacerbations (or moderate/severe exacerbations for patients in the COPD group), risk-domain control, average daily dose of short-acting beta<sub>2</sub> agonist (SABA), and number of acute respiratory events (patients in asthma group).

The definition of severe asthma exacerbations was the sensitive definition based on the American Thoracic Society/European Respiratory Society (ATS/ERS) guidelines.<sup>34</sup> A severe exacerbation included one or more occurrences of any of the following: asthma-related hospital admission or accident and emergency (A&E) attendance and an acute course of oral corticosteroids (OCS) with lower respiratory consultation.

For patients in the COPD group, a moderate or severe exacerbation included one or more occurrences of any of the following: COPD-related, unscheduled hospital admission or A&E attendance; an acute course of OCS with lower respiratory consultation; and antibiotics prescribed with

lower respiratory consultation. The risk-domain control of COPD was defined as the absence of any moderate/severe exacerbations.

For patients in the asthma group, an acute respiratory event was defined as one or more occurrences of any of the following: asthma-related hospital admission or A&E attendance; an acute course of OCS; and antibiotics prescribed with lower respiratory consultation. The risk-domain control of asthma was defined as the absence of any acute respiratory events.

Finally, the average daily dose of SABA during the outcome year was calculated as the average number of puffs per day over the year multiplied by the strength of dose (in  $\mu\text{g}$  of salbutamol equivalents).

More detailed definitions of the above outcomes can be found in the [Supplementary materials](#).

## Statistical analysis

### Baseline characterization

Baseline variables were summarized and compared between cohorts. This was done separately in the asthma and COPD groups to identify potential matching variables, ie, those that differed between cohorts. Binary and categorical variables were summarized using counts and percentages by category. Continuous variables were summarized using mean and standard deviation or median and interquartile range. Variables were compared between cohorts to assess imbalance and bias potential. Imbalance was measured using the standardized mean difference (SDD)<sup>35</sup> between cohorts, while bias potential was measured using the relative change in coefficient (RCC).<sup>36</sup>

### Matching

The baseline variables with high bias potential ( $> 2\%$ ) and high level of imbalance ( $\text{SDD} > 10\%$ )<sup>37</sup> were considered for matching. This was done separately within the asthma group and the COPD group, ie, patients with asthma only were matched with patients with asthma only (which formed the matched asthma group) and patients with COPD were matched with patients with COPD (which formed the matched COPD group), regardless of coexisting asthma diagnosis. Exact matching combined with a propensity score was carried out with a mixture of ratios of patients in the no-payment cohort to patients in the payment cohort (1:1, 1:2, and 1:3). After matching, bias potential was again assessed to identify areas where residual confounding was present for the different outcomes. More details on matching methods can be found in the [Supplementary materials](#).

## Outcome analysis

Conditional logistic regression was used to compare the odds of good adherence of patients in the payment cohort with that of the matched patients in the no-payment cohort. This was carried out in the asthma and COPD groups combined.

Conditional logistic regression was used to estimate odds ratios (ORs) for risk-domain control, comparing the payment cohort with the no-payment cohort. Conditional Poisson regression was used to estimate rate ratios (RRs), for the number of severe exacerbations (or moderate/severe in the COPD group) and the number of acute respiratory events (in the asthma group only). Finally, ordinal logistic regression was used to estimate the proportional odds of being in a higher category of SABA dose. For this analysis, the average daily dose of SABA was categorized as 0, >0–200, >200–400, >400–600, and >600 µg/day (salbutamol equivalent). These categories have been used in previous studies.<sup>38,39</sup>

In all cases, multivariable models were fitted, adjusting for identified residual confounders, and model estimates were computed with 95% confidence intervals (CIs). The level of statistical significance was set at 5%. The study had 90% power to detect a difference in the proportion of adherent patients of 7.7%, with a two-sided significance of 5%. All statistical analyses were carried out using Stata MP6 Version 12 and Stata SE Version 14 (StataCorp, College Station, TX, USA).

## Sensitivity analysis

The analysis of the primary outcome was repeated in a subgroup of patients who had evidence of 9 months (270 days) of persistent use of FP/SAL medication. This meant that the patient received prescription for FP/SAL continuously over 9 months, with no more than 30 days between the dates of issue of prescriptions. Discontinuation was defined as having no repeat prescription of FP/SAL in 6 months (180 days).

The main analysis was carried out in cohorts deriving from all three matching ratio strategies, ie, 1:1, 1:2, and 1:3. A sensitivity analysis was carried out using cohorts that resulted from the 1:3 matching strategy only.

## Results

### Patients

After applying study criteria, 6,716 patients with asthma or COPD were available from the primary care databases. Details of specific inclusions and exclusions are shown in [Figure S1](#). Of the 6,716 unmatched patients, 6,000 were likely to require co-payments (ie, registered in England) and 716 were not required to pay for prescriptions (ie, registered

in Scotland). After matching, these figures were reduced to 1,640 patients in the payment cohort and 619 patients in the no-payment cohort. Of the payment cohort, 1,378 patients had asthma only and 262 patients had COPD, and of the no-payment cohort, 512 patients had asthma only and 107 patients had COPD (Table 1).

An overview of the variables with relevant differences between the cohorts is listed in [Table S1](#). The proportions of patients matched 1:1, 1:2, or 1:3 are listed in [Table S2](#) and show that the majority of patients were matched to three others.

Baseline characteristics of the matched cohorts, with disease groups combined, are listed in Tables 2 and 3. Patients had a mean age of 44 years in both cohorts; 47–48% of them were male, 28–30% of them were current smokers, and 35–36% of them had a body mass index (BMI) of  $\geq 30$ . All of these characteristics were well matched between cohorts, such that *P*-values for comparison were nonsignificant, and SDDs were <10%. Prevalence of comorbidities was similar between cohorts, and the median Charlson comorbidity index (CCI) was 4.0 (Table 2).

The average daily dose of SABA was >600 µg/day in 11.2% of the patients in the payment cohort and 13.1% of the patients in the no-payment cohort, and ~15% of the patients in each cohort had two or more prescriptions of OCS, during the baseline year (Table 3). Approximately 42–45% of the patients with asthma had at least one severe exacerbation and 30–32% of the patients with COPD had at least one moderate or severe exacerbation. All baseline variables had an SDD of <10%, indicating good balance between the matched cohorts. Definitions of the baseline variables in Table 3 are found in the [Supplementary materials](#). Further baseline characteristics are listed in [Table S3](#). Baseline characteristics for the asthma and COPD groups separately are listed in [Tables S4–S6](#).

## Primary outcome

During the outcome year, 34.3% of the patients in the payment cohort and 34.9% of the patients in the no-payment cohort had good adherence to FP/SAL therapy (ie, MPR

**Table 1** Number of patients by cohort and disease group

Disease group	Unmatched (n=6,716)		Matched (n=2,259)	
	Payment cohort	No-payment cohort	Payment cohort	No-payment cohort
Asthma	4,882	582	1,378	512
COPD	1,118	134	262	107
Total	6,000	716	1,640	619

**Abbreviation:** COPD, chronic obstructive pulmonary disease.

**Table 2** Baseline demographic characteristics and comorbidities in combined disease groups, by matched cohort

Variables	Payment cohort (n=1,640)	No-payment cohort (n=619)	P-value <sup>a</sup>	SDD
Index year, n (%)			0.176	5.9
2012	879 (53.6)	322 (52.0)		
2013	509 (31.0)	191 (30.9)		
2014	231 (14.1)	90 (14.5)		
2015	21 (1.3)	16 (2.6)		
Age, mean (SD)	44.0 (11.1)	44.4 (11.2)	0.341	3.7
Male, n (%)	775 (47.3)	298 (48.1)	0.707	1.8
Smoking status, n (%)			0.687	3.6
Never smoked	762 (46.5)	275 (44.4)		
Current smoker	469 (28.6)	184 (29.7)		
Exsmoker	409 (24.9)	160 (25.8)		
BMI (kg/m <sup>2</sup> ), n (%)			0.356	2.5
<18.5	15 (0.9)	11 (1.8)		
18.5–<25	557 (34.0)	206 (33.3)		
25–<30	478 (29.1)	186 (30.0)		
≥30	590 (36.0)	216 (34.9)		
Comorbidities, n (%)				
Cardiovascular disease	114 (7.0)	41 (6.6)	0.784	1.3
Ischemic heart disease	37 (2.3)	13 (2.1)	0.822	1.1
Hypertension	218 (13.3)	80 (12.9)	0.817	1.1
Cancer	142 (8.7)	70 (11.3)	0.054	8.8
Diabetes	76 (4.6)	36 (5.8)	0.249	5.3
Rhinitis	469 (28.6)	191 (30.9)	0.292	4.9
Active rhinitis	191 (11.6)	81 (13.1)	0.349	4.4
Gastroesophageal reflux disease	166 (10.1)	73 (11.8)	0.249	5.3
Active gastroesophageal reflux disease	121 (7.4)	49 (7.9)	0.666	2.0
Eczema	486 (29.6)	170 (27.5)	0.311	4.8
Active eczema	31 (1.9)	14 (2.3)	0.573	2.6
Pneumonia	72 (4.4)	24 (3.9)	0.590	2.6
Oral candidiasis	15 (0.9)	7 (1.1)	0.641	2.1
Charlson comorbidity index, median (IQR)	4.0 (4.0)	4.0 (4.0)	0.554	3.1

**Note:** <sup>a</sup>P-value estimated from Kruskal–Wallis equality-of-populations rank test, or the Pearson's chi-square test of independent categories, where appropriate.

**Abbreviations:** SDD, standardized mean difference; SD, standard deviation; BMI, body mass index; IQR, interquartile range.

≥80%). This proportion was lower in patients with asthma (31.4% in both cohorts) and higher in patients with COPD (49.6 and 51.4% in the payment and no-payment cohorts, respectively; Table 4). In unadjusted comparisons, adherence was not significantly different between cohorts ( $P=0.801$ ). The distribution of the adherence levels is shown graphically in Figure 1. Summary outcome data for the asthma and COPD groups separately are listed in Table S7.

When adjusting for potential confounders, the adjusted OR for adherence across both disease groups was 1.04 (95% CI, 0.85–1.27; Table 5), comparing the payment cohort with the no-payment cohort, showing that adherence was similar between cohorts. Variables used for adjusting were selected based on the residual confounding observed (Table S8).

## Secondary outcomes

During the outcome year, 62–64% of the patients with asthma achieved risk-domain control and 45–47% of the patients with

COPD achieved risk-domain disease control (Table 4). These proportions were not significantly different between cohorts. There were fewer exacerbations in the outcome year, but exacerbations remained similar between cohorts. Furthermore, there was no difference between acute respiratory events (in patients with asthma) and the average daily dose of SABA, when cohorts were compared in unadjusted analyses (Table 4).

When adjusting for residual confounders, the adjusted ORs for risk-domain control were 0.89 (95% CI 0.71–1.11) in patients with asthma and 0.89 (95% CI 0.53–1.48) in patients with COPD (Table 5). Adjusted RRs for severe exacerbations (or moderate/severe exacerbations in COPD) were 1.01 (95% CI 0.85–1.21) in patients with asthma and 1.27 (95% CI 0.98–1.65) in patients with COPD, comparing the payment cohort with the no-payment cohort. None of these associations were statistically significant. For patients with asthma, there was no difference between cohorts in terms of acute respiratory events (RR 1.10; 95% CI 0.96–1.27), after adjustment



**Table 3** Baseline clinical characteristics in combined disease groups, by matched cohort

Variables	Payment cohort (n=1,640)	No-payment cohort (n=619)	P-value <sup>a</sup>	SDD
Average daily dose of SABA (µg, salbutamol equivalent), n (%)			0.597	6.9
0	745 (45.4)	270 (43.6)		
>0–≤200	397 (24.2)	144 (23.3)		
>200–≤400	221 (13.5)	82 (13.2)		
>400–≤600	94 (5.7)	42 (6.8)		
>600	183 (11.2)	81 (13.1)		
Any OCS prescriptions, n (%)			0.864	2.2
0	1,132 (69.0)	420 (67.9)		
1	259 (15.8)	102 (16.5)		
≥2	249 (15.2)	97 (15.7)		
Antibiotic prescriptions, with LR consultation, <sup>b</sup> n (%)			0.432	5.5
0	1,108 (67.6)	406 (65.6)		
1	317 (19.3)	119 (19.2)		
≥2	215 (13.1)	94 (15.2)		
A&E attendances (at least 1), n (%)	4 (0.2)	2 (0.3)	0.744	1.5
Outpatient visits <sup>c</sup> (at least 1), n (%)	33 (2.0)	13 (2.1)	0.895	0.6
Moderate/severe exacerbations (COPD), n (%)			0.688	5.6
0	941 (57.4)	339 (54.8)		
1	376 (22.9)	147 (23.7)		
2	199 (12.1)	80 (12.9)		
≥3	124 (7.6)	53 (8.6)		
Severe exacerbations (asthma), n (%)			0.845	4.0
0	1,141 (69.6)	422 (68.2)		
1	306 (18.7)	117 (18.9)		
2	121 (7.4)	48 (7.8)		
≥3	72 (4.4)	32 (5.2)		

**Notes:** <sup>a</sup>P-value estimated from Kruskal–Wallis equality-of-populations rank test, or the Pearson's chi-square test of independent categories, where appropriate. <sup>b</sup>LR consultations were identified by: Read codes (including asthma, COPD and LRTI); asthma/COPD review codes excluding any monitoring letter codes; and lung function and/or asthma monitoring. <sup>c</sup>Outpatient visits for asthma or LR code or generic outpatient code on same day as respiratory consultation.

**Abbreviations:** SDD, standardized mean difference; SABA, short-acting beta<sub>2</sub> agonist; OCS, oral corticosteroids; LR, lower respiratory; LRTI, lower respiratory tract infection; A&E, accident and emergency; COPD, chronic obstructive pulmonary disease.

for confounders. Proportional ORs for the average daily dose of SABA showed no difference in odds of being in a higher category of SABA dose in the payment cohort than in the no-payment cohort. Definitions of variables used to adjust the outcome models can be found in the [Supplementary materials](#).

## Sensitivity analyses

Since the adherence measure (MPR in a fixed 1-year period) included both implementation and persistence, a sensitivity analysis was performed, restricting the population to patients with at least 9 months of persistent use of FP/SAL after initiation. Of the matched cohorts, 80.2% of the patients in the payment cohort and 77.7% of the patients in the no-payment cohort met this criterion. In this subgroup, 42.1% of the patients in the payment cohort and 44.3% of the patients in the no-payment cohort had good adherence to treatment, in both disease groups combined (Table 4). In an adjusted model, there was no significant difference between cohorts in terms of adherence (Table S9).

Results were similar in sensitivity analysis restricted to patients matched 1:3 (Table 5). One exception, however,

was in the comparison of severe exacerbations in the asthma group, which found that patients in the payment cohort had double the rate of exacerbations as patients in the no-payment cohort (RR, 2.03; 95% CI, 1.05–3.93).

The main analysis investigated adherence in both disease groups combined, but an ad hoc sensitivity analysis was carried out in patients with asthma only. This was due to poorer matching of patients with COPD, evidenced by the presence of residual confounding. Patients with COPD were, therefore, excluded to test the robustness of the main results. In patients with asthma only, there was no difference in adherence between the two cohorts and no difference in the subgroup with 9 months of persistence (Table S9).

## Discussion

This study showed that patients with asthma or COPD registered at general practices in England – who were likely to pay for their prescriptions – and patients with asthma or COPD registered in Scotland – who were not paying for prescriptions – had similar adherence to FP/SAL therapy. This was observed after matching by demographic characteristics and

**Table 4** Summary of outcomes, by cohort

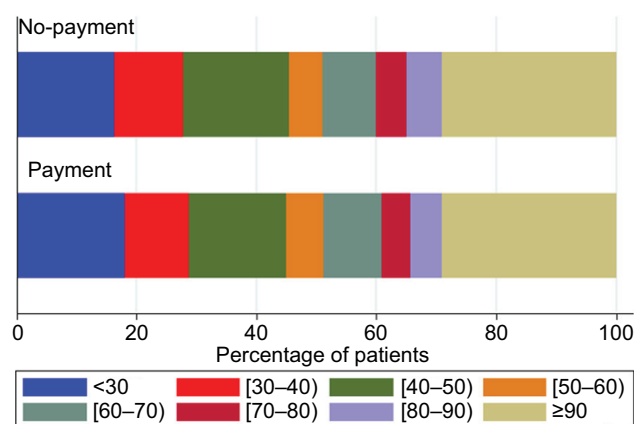
Variables, n (%)	Payment cohort (n=1,640)	No-payment cohort (n=619)	P-value <sup>a</sup>
Adherence achieved (MPR >80%)			
Both disease groups	563 (34.3)	216 (34.9)	0.801
Asthma group	433 (31.4)	161 (31.4)	0.992
COPD group	130 (49.6)	55 (51.4)	0.756
With 9-month persistence <sup>b</sup>	554 (42.1)	213 (44.3)	0.414
Risk domain control achieved			
Asthma group	854 (62.0)	325 (63.5)	0.549
COPD group	117 (44.7)	50 (46.7)	0.717
Severe asthma exacerbations			0.950
0	1,053 (76.4)	389 (76.0)	
1	216 (15.7)	79 (15.4)	
2	70 (5.1)	27 (5.3)	
≥3	39 (2.8)	17 (3.3)	
Moderate/severe COPD exacerbations			0.393
0	117 (44.7)	50 (46.7)	
1	62 (23.7)	26 (24.3)	
2	31 (11.8)	17 (15.9)	
≥3	52 (19.8)	14 (13.1)	
Acute respiratory events, asthma			0.865
0	854 (62.0)	325 (63.5)	
1	341 (24.7)	121 (23.6)	
2	103 (7.5)	40 (7.8)	
≥3	80 (5.8)	26 (5.1)	
SABA, average daily dose (µg/day, salbutamol equivalent), asthma			0.706
0	463 (33.6)	174 (34.0)	
>0–≤200	389 (28.2)	133 (26.0)	
>200–≤400	257 (18.7)	98 (19.1)	
>400–≤600	121 (8.8)	42 (8.2)	
>600	148 (10.7)	65 (12.7)	
SABA, average daily dose (µg/day, salbutamol equivalent), COPD			0.155
0	74 (28.2)	33 (30.8)	
>0–≤200	52 (19.8)	14 (13.1)	
>200–≤400	48 (18.3)	28 (26.2)	
>400–≤600	46 (17.6)	12 (11.2)	
>600	42 (16.0)	20 (18.7)	

**Notes:** <sup>a</sup>P-value estimated from Kruskal–Wallis equality-of-populations rank test, or the Pearson's chi-square test of independent categories, where appropriate. <sup>b</sup>A total of 80.2% of the patients in the payment cohort and 77.7% of the patients in the no-payment cohort had 9 months of persistent use of FP/SAL.

**Abbreviations:** MPR, medication possession ratio; COPD, chronic obstructive pulmonary disease; SABA, short-acting beta<sub>2</sub> agonist; FP/SAL, fluticasone propionate/salmeterol xinafoate.

indicators of disease severity and after adjusting for residual confounding.

The level of adherence to maintenance therapy observed in this study – ~35% in patients with asthma and 50% in patients with COPD – was within the range reported in

**Figure 1** Distribution of medication possession ratio (%) in matched cohorts.

other studies.<sup>9,40–43</sup> In terms of the association between co-payments for prescription and adherence, the current study was not consistent with recent literature. A meta-analysis published in 2013,<sup>44</sup> which included only US studies of varying medication types, estimated an overall increase in odds of nonadherence of 11% in co-payment versus no co-payment. The authors noted several limitations of the results, however – that the follow-up times varied between studies, between 3 months and 2 years, and that the quality of studies was poor in some cases. Another meta-analysis on studies done in the USA or Canada<sup>45</sup> found that 56 of the 66 studies reported an inverse relationship between co-payment and adherence, with 10 studies reporting a nonsignificant relationship, or limited data to detect a relationship. These studies also covered a range of medication types. The authors acknowledged that the analysis of adherence should account for factors affecting adherence, such as disease severity, and that this was not done consistently across the studies included. The review covered two studies on ICS. One study, using electronic medical records, did not find a difference in adherence between co-payment levels.<sup>46</sup> The second study, using pharmacy records, showed a significant negative association with cumulative co-payment level and implementation adherence.<sup>40</sup> Another study in the same patient population in Pennsylvania showed that patients with higher co-payment costs were less likely to initiate asthma medication.<sup>47</sup> A discrete choice experiment of patients with COPD from the USA showed medication costs for the patients to be an important factor in treatment decisions.<sup>48</sup>

An explanation for the inconsistency between the current study and what is reported in the literature could be publication bias: negative studies tend to be less likely to be published,<sup>49</sup> but this only counts for studies with the association between co-payment and adherence as their main

**Table 5** Results of adjusted outcome models

Outcome variable	All matching ratios used (N=2,259)		Only 1:3 matching ratio used (N=1,928)	
	Adjusted odds ratio/rate ratio <sup>a</sup> (with 95% CI)	P-value	Adjusted odds ratio/rate ratio <sup>a</sup> (with 95% CI)	P-value
Combined disease groups				
Adherence <sup>b</sup>	1.04 (0.85–1.27)	0.704	1.58 (0.80–3.14)	0.191
Asthma				
Risk-domain control <sup>b</sup>	0.89 (0.71–1.11)	0.294	0.56 (0.27–1.17)	0.125
Number of severe exacerbations <sup>c</sup>	1.01 (0.85–1.21)	0.907	2.03 (1.05–3.93)	0.035
Number of acute respiratory events <sup>d</sup>	1.10 (0.96–1.27)	0.163	1.56 (0.99–2.47)	0.055
SABA average daily dose <sup>e</sup>	1.05 (0.86–1.27)	0.655	1.22 (0.61–2.47)	0.576
COPD				
Risk-domain control <sup>f</sup>	0.89 (0.53–1.48)	0.641	NC	
Number of moderate/severe exacerbations <sup>g</sup>	1.27 (0.98–1.65)	0.067	NC	
SABA average daily dose <sup>h</sup>	1.03 (0.65–1.63)	0.907	NC	

**Notes:** NC means model did not reach convergence. <sup>a</sup>Odds/rate ratios compare payment cohort with no-payment cohort. <sup>b</sup>Adjusted by antibiotic prescriptions with lower respiratory indication. <sup>c</sup>Adjusted by acute OCS courses (sensitive definition) LTRA prescription, and maintenance OCS prescription. <sup>d</sup>Adjusted by antibiotic prescriptions with lower respiratory indication and LTRA prescription. <sup>e</sup>Adjusted by the average daily dose of SABA. <sup>f</sup>Adjusted by eczema, cancer, active gastroesophageal reflux disease, rhinitis, number of SABA inhalers, any OCS prescription and acute OCS course (sensitive definition). <sup>g</sup>Adjusted by ICS prescriptions, acute OCS prescriptions (probable definition), SABA inhaler prescriptions, Medical Research Council dyspnea score, and GOLD severity. <sup>h</sup>Adjusted by average daily dose of SABA and GOLD severity.

**Abbreviations:** CI, confidence interval; SABA, short-acting beta<sub>2</sub> agonist; COPD, chronic obstructive pulmonary disease; NC, no convergence; OCS, oral corticosteroids; LTRA, leukotriene receptor antagonist; ICS, inhaled corticosteroids; GOLD, Global Initiative for Chronic Obstructive Lung Disease.

objective. The cited studies came from the USA and Canada. Cultural and socioeconomic differences existing between these countries and the UK could be another explanation for the discrepancy in findings.

This study found no difference in disease outcomes between patients likely paying for prescriptions and patients not paying for prescriptions in the majority of cases. The association between adherence and effectiveness of treatment has been demonstrated previously,<sup>9</sup> and hence it is unsurprising that matched cohorts with similar adherence would observe similar disease outcomes. However, there was some inconsistency in the results: patients in the payment cohort were found to have significantly more severe asthma exacerbations than those in the no-payment cohort when the analysis was restricted to cohorts derived from one matching strategy, rather than a mixture of matching strategies. This inconsistency cannot easily be explained but is potentially an anomaly as the results were consistent on all other endpoints.

The strength of this study is its size and the real-life data used, which is likely to be representative of patients with asthma and patients with COPD. Furthermore, the length of follow-up allowed for a good measure of adherence to be estimated. As this was an observational study to compare two cohorts, methods were applied to ensure that the cohorts were as similar as possible, so that associations with the exposure (co-payment of prescriptions, or no co-payment of prescriptions) could be studied. The

study design included matching patients based on important demographic and clinical factors and adjusting models by further potential confounders. When the level of residual confounding was high, as was observed in the COPD group, ad hoc analyses were carried out in the asthma group only, but the exclusion of the COPD group did not appear to impact the results.

## Limitations

Limitations of the current study include the possibility of residual confounding. Despite matching methods and adjustment for measured confounders, the nature of the study – being observational, rather than a randomized controlled trial – is such that confounding by unmeasured variables may be present in the results. Medication-taking behavior can be affected by many factors. Such factors may be subjective and unique to the patients – for example, attitude toward the health care system and personal perception of their disease. This is an added difficulty in the analysis of adherence.<sup>50</sup> Furthermore, the two cohorts being compared in this study were from different geographical locations in the UK, which are known to differ in terms of chronic disease and life expectancy.<sup>51,52</sup> Matching by the history of disease outcomes will have gone some way to alleviate differences between patients in England and Scotland, and the study cohorts were well-matched in terms of comorbidities. Other potential differences, such as income of the patients,



could not be accounted for in these data. The results did not include practice-level clustering and, therefore, may have been affected by potential differences between practices, such as the level of health education offered. In addition, despite the quality of the databases used, it cannot be guaranteed that all information is complete.<sup>53</sup>

The study did not distinguish between the three components of adherence (initiation, implementation, and persistence). By requiring at least two prescriptions for a patient to be included in the study, we excluded patients who did not initiate their prescribed treatment. Therefore, our study was not able to detect differences in that aspect of adherence. In a sensitivity analysis restricting the population to patients with at least 9 months of persistent FP/SAL use, we showed that our conclusion of no difference in adherence between the payment and no-payment cohorts was not driven by the inclusion of nonpersistent patients.

We did not exclude patients who switched to other ICS medication, and such patients will have appeared to have poor adherence to FP/SAL therapy in our dataset. However, since the observed adherence levels were within the range of results from other studies, and both cohorts are likely to have been affected equally, we do not think this affected the results.

Our algorithm for MPR did not adjust observed prescription dates and durations for stockpiling, dose changes, or medication switches, which could have resulted in a reduced precision of adherence quantification.

The study endeavored to exclude patients registered in general practices in England who were exempt from prescription charges. This was done by searching for common reasons for exemption, such as pregnancy, and by searching for a record of exemption status. In doing this, the study relied on the accuracy of the primary care source databases. It is likely that the vast majority of patients with exemptions have been excluded, but it remains a possibility that a small proportion of exempt patients were in the study sample.

## Conclusion

Patient adherence is vital if treatment is to be effective. At a time when prescription costs are high and put a strain on the health care system,<sup>54</sup> it is important to ascertain whether measures to relieve the strain – such as co-payments – have an adverse effect on adherence to treatment, and consequently, on patient outcomes. In this study, there was no difference in adherence between matched patients registered in England and Scotland, suggesting that prescription charges do not have an impact on adherence to maintenance therapy for asthma or COPD.

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## Author contributions

BV, JFMvB, DR, MM, and DBP developed the research question. JV and DBP developed the study design. JV conducted the data preparation and analyses. LML drafted the manuscript. All authors were involved in the interpretation of the data and the critical review and revision of the manuscript. All authors read and approved the final manuscript.

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