# **Supplemental Material**

#### ATTRIBUTE LEVELS AND DEFINITIONS

Discrete choice experiment (DCE) attributes were defined based on key efficacy and risk outcomes relevant for asthma and chronic obstructive pulmonary disease (COPD) maintenance inhaler treatments as well as features characterizing the convenience of using these maintenance inhalers and out-of-pocket costs for maintenance medicine. Because the attributes that are most relevant to patients differ for asthma versus COPD (e.g., increased risk of pneumonia is only relevant for COPD patients), two unique DCEs were administered to patients with asthma and patients with COPD. The DCE attributes were selected based on a targeted literature review, opinions generated from focus groups consisting of patients with asthma (n=15) or COPD (n=22),[1] and consultations with clinical experts on the project team.

The attributes for this study span three domains: (1) efficacy, characterized by onset of action of the inhaled therapy and exacerbations per year; (2) safety, characterized by 5-year risk of osteoporosis, and for COPD patients, 5-year risk of pneumonia; and (3) non-clinical features of the inhaler related to convenience, characterized by device type, dosing frequency, dose counters, and priming. The attribute levels were constructed based on: asthma and COPD treatment product characteristics; clinical expert feedback; and a review of clinical data sources, published studies on patients' perceptions of asthma and COPD endpoints, and preference studies in patients with asthma or COPD. Definitions of attributes and attribute levels are provided in **Supplemental Table 1**.

# **Supplemental Table 1. Attribute and Level Definitions**

Attribute	Attribute Description	Level	Level Description
Number of	For Asthma: An asthma exacerbation, or	1	You experience one exacerbation in a
exacerbations	asthma attack, is when your airways	exacerbation	year.
	become swollen and the muscles around	per year	
	your airways contract. This may cause you	2	You experience two exacerbations in a
	to experience difficulty breathing,	exacerbations	year.
	coughing, or wheezing. You will need to	per year	
	visit your health care provider or	3	You experience three exacerbations in
	emergency room to get extra medication,	exacerbations	a year.
	such as oral steroids, and may require an	per year	
	overnight stay in the hospital.		
	For COPD: A COPD exacerbation is		
	when you experience a "flare-up", or		
	sudden worsening of respiratory		
	symptoms, and become sick. You will		
	need to visit your health care provider or		
	emergency room to get extra medication,		
	such as oral steroids or antibiotics, and		
	may require an overnight stay in the		
	hospital.		
Onset of	After you use your maintenance inhaler, it	5 minutes	Your medication will begin to work
action	may take a few minutes before it starts		within 5 minutes after using your
	working. With some inhalers, the		maintenance inhaler.
	medication will start working within just 5	15 minutes	Your medication will begin to work
	minutes, but with others it may take up to		within 15 minutes after using your
	30 minutes. The amount of time it takes		maintenance inhaler.
	for your medication to begin working is	20 minutes	Your medication will begin to work
	called onset of action.		within 20 minutes after using your
			maintenance inhaler.
		30 minutes	Your medication will begin to work
			within 30 minutes after using your
			maintenance inhaler.
Osteoporosis	Some maintenance medications increase	4% (4 out of	You have a 4% chance of
,	your chance of getting osteoporosis.	100 patients)	experiencing a hip fracture within 5
	Osteoporosis is a disease that affects your	, , , , , , , , , , , , , , , , , , ,	years of using your inhaler.
	bones. Your body may then lose bone	5% (5 out of	You have a 5% chance of
	mass and your bones may become weak	100 patients)	experiencing a hip fracture within 5
	mass and your cones may become weak	100 patients)	years of using your inhaler.
			Jours of using your finiater.

Attribute	Attribute Description	Level	Level Description
	and break easily. As a result, you may	6% (6 out of	You have a 6% chance of
	experience a hip fracture.	100 patients)	experiencing a hip fracture within 5
			years of using your inhaler.
Dosing	Dosing frequency refers to the number of	Once daily	You need to take one or two puffs of
Frequency	times per day your doctor recommends		your maintenance inhaler once per
	you use your maintenance inhaler. Some		day.
	maintenance medications should be used	Twice daily	You need to take one or two puffs of
	once per day while others should be used		your maintenance inhaler two times
	twice per day (once in the morning and		per day - once in the morning and
	once in the evening, approximately 12		again in the evening
	hours apart).		
Priming	Before you use your maintenance	One or two	Each time you use your inhaler, you
	medication for the first time, it may	simple steps	will need to do one or two simple
	require you to insert a cartridge containing	to get dose	steps to get it ready. These steps only
	all the doses. Some inhalers require you to	ready	include opening the device and pulling
	follow a few simple steps to get it ready		either one or two levers. The steps are
	each time you use them. Others require		the same for your first dose and all
	you to remove a capsule from separate foil		other doses.
	packaging and inserting it into the device.	Need to	The device requires you to discharge
		discharge test	several doses into the air before you
		doses on first	use it for the first time. Between each
		use. One	discharge of a test dose, you must
		simple step	shake the device for five seconds.
		to get dose	Each time you use your inhaler for all
		ready.	other doses, you will just need to open
			the device.
		Need to	The device requires you to go through
		insert	several steps to get it ready before you
		canister on	use it for the first time. First, you will
		first use.	need to remove the base of the inhaler
		Two steps to	and insert a separate canister of the
		get dose	medicine into the inhaler and push
		ready.	down hard until it clicks into place.
			Then, you must reattach the base into
			place. Each time you use the inhaler,
			you need to keep it upright, turn the
			base, and open the cap.

Attribute	Attribute Description	Level	Level Description
		Need to	Each time you use this inhaler, you
		insert a new	will need to remove a capsule
		capsule into	containing the medicine from a
		inhaler each	separate package, insert it into the
		time to get	inhaler, and then close the inhaler to
		dose ready.	get it ready for use. The steps are the
			same for your first dose and all other
			doses.
Device type	Maintenance inhalers deliver your	Pressurized	These inhalers are pressurized and
	medicine in two different ways: 1) One	inhaler	force the medication out when you
	type of inhaler is pressurized, like an		depress the canister or push the
	aerosol can. The medication is forced out		button. When you use these, you must
	and you inhale moderately deeply and		inhale at slow speed, moderately
	somewhat slowly. 2) The other type of		deeply.
	inhaler contains the medicine in dry	Soft mist	These inhalers gently release the
	powder. For dry powder inhalers, you take	inhaler	medication out in a continuous liquid
	a fast, very deep breath.		mist when you press the dose-release
			button. When you use these, you must
			inhale slowly and deeply as you press
			the button, continuing until your lungs
			are full.
		Dry powder	These inhalers are not pressurized and
		inhaler	hold the medicine in a dry powder
			form. When you use these inhalers,
			you must inhale very fast and deeply.
Dose counter	Inhalers often come with a dose counter	Every dose	The dose counter displays exactly the
	built in that tells you how much medicine		number of doses left.
	you have left. Some dose counters are very	Every 10	The dose counter displays the number
	precise and display the exact number of	doses	of doses left but counts only every 10
	doses that are left. Some dose counters are		doses.
	very precise but tell you how many doses	Metered dose	The dose counter is displayed as a
	you have left in counts of 10. Other dose	counter	needle on a meter, much like you
	counters are in the form of a meter,		would see a gas gauge in a car. It
	displaying an approximate amount of		moves after every dose.
	medicine left and look similar to a gas		
	gauge in a car.		

Attribute	<b>Attribute Description</b>	Level	Level Description
Pneumonia	Some maintenance medications increase	10% (10 out	You have a 10% chance of getting
(COPD	your chance of getting pneumonia.	of 100	pneumonia in the coming 5 years.
patients only)	Pneumonia is a type of lung infection that	patients)	
	can cause coughing, fever, shortness of	15% (15 out	You have a 15% chance of getting
	breath, fatigue, weakness, nausea,	of 100	pneumonia in the coming 5 years.
	vomiting, and diarrhea.	patients)	
		20% (20 out	You have a 20% chance of getting
		of 100	pneumonia in the coming 5 years.
		patients)	

### **DOMINATED-CHOICE QUESTION**

For the dominated-choice question, one choice option was at least as good or better than the other on all attributes (**Supplemental Figure 1**). This dominated-choice question was used to assess whether patients understand the discrete choice task and were responding appropriately to the choices. The dominated-choice question was presented as the last question of the DCE exercise.

Although the patient population overall preferred once-daily over twice-daily dosing (**Figure 3**), some individual patients might have preferred twice-daily over once-daily dosing. For such patients, the dominated-choice question in **Supplemental Figure 1** is inappropriate.

#### **Supplemental Figure 1. Dominated-Choice DCE Question**

Your doctor asks you to decide between two maintenance inhaler medications, Inhaler A and Inhaler B. The outcomes you will experience after using either Inhaler A or B are described in the table below.

Take a moment to review the outcomes produced by each treatment, and then indicate which treatment you would choose at the bottom of the table.

Outcomes	Inhaler A	Inhaler B
Number of exacerbations	3 exacerbations per year	1 exacerbation per year
Onset of action	30 minutes	15 minutes
Osteoporosis	4% (4 out of 100 patients)	4% (4 out of 100 patients)
	risk of hip fracture over 5	risk of hip fracture over 5
	years	years
Pneumonia	20% (2 out of 100 patients)	10% (1 out of 100 patients)
	risk of pneumonia over 5	risk of pneumonia over 5
	years	years
Dosing frequency	Once daily	Twice daily
Device type	Pressurized inhaler	Pressurized inhaler
Dose counter	Metered dose counter	Metered dose counter

#### **DCE DESIGN**

Using NGene version 1.1 (ChoiceMetrics, Sydney, Australia), separate D-efficient experimental designs were generated for the asthma and COPD surveys, applying directional priors when applicable. The designs consisted of 96 questions split into eight survey versions (blocks) with 12 choice tasks per block and four blocks each for asthma and COPD. For each choice task, patients were asked to choose between two unlabeled treatments (A and B) described by the treatment attributes. The design did not include constraints for level combinations. Although some attribute combinations do not occur with currently available inhalers, such as capsule

priming with pressurized inhaler type, no issues with level combinations were detected in the pilot interviews.

#### STATISTICAL ANALYSIS OF DCE DATA

The analysis of the DCE data followed random utility maximization theory, by assuming that within every choice task (t), each respondent (n) always chose the treatment (j) that resulted in the highest utility.[2-4]

Baseline utility of patients with asthma was defined as

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\begin{split} u_{jnt}^{\text{asthma}} &= \alpha_{jnt}^{\text{asthma}} + \beta_{1}^{\text{asthma}} \text{one\_exacerbation}_{jnt} + \beta_{2}^{\text{asthma}} \text{two\_exacerbations}_{jnt} + \\ \beta_{3}^{\text{asthma}} &20 \text{min\_onset}_{jnt} + \beta_{4}^{\text{asthma}} 15 \text{min\_onset}_{jnt} + \beta_{5}^{\text{asthma}} 5 \text{min\_onset}_{jnt} + \\ \beta_{6}^{\text{asthma}} &5\% \text{\_osteoporosis\_risk}_{jnt} + \beta_{7}^{\text{asthma}} 4\% \text{\_osteoporosis\_risk}_{jnt} + \\ \beta_{8}^{\text{asthma}} &\text{once\_daily}_{jnt} + \beta_{9}^{\text{asthma}} \text{canister}_{jnt} + \beta_{10}^{\text{asthma}} \text{discharge}_{jnt} + \\ \beta_{11}^{\text{asthma}} &1 \text{to2\_easy\_steps}_{jnt} + \beta_{12}^{\text{asthma}} \text{soft\_mist}_{jnt} + \beta_{13}^{\text{asthma}} \text{pressurized\_device}_{jnt} + \\ \beta_{14}^{\text{asthma}} &1 \text{to2\_easy\_steps}_{jnt} + \beta_{15}^{\text{asthma}} \text{count\_evey\_10th\_dose}_{jnt} + \\ \beta_{16}^{\text{asthma}} &\text{count\_evey\_dose}_{jnt} + \varepsilon_{jnt} \end{split}
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Baseline utility of patients with COPD was defined as

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\begin{split} u_{jnt}^{\text{COPD}} &= \alpha_{jnt}^{\text{COPD}} + \beta_{1}^{\text{COPD}} \text{one\_exacerbation}_{jnt} + \beta_{2}^{\text{COPD}} \text{two\_exacerbations}_{jnt} + \\ \beta_{3}^{\text{COPD}} &20 \text{min\_onset}_{jnt} + \beta_{4}^{\text{COPD}} 15 \text{min\_onset}_{jnt} + \beta_{5}^{\text{COPD}} 5 \text{min\_onset}_{jnt} + \\ \beta_{6}^{\text{COPD}} &5\%\_\text{osteoporosis\_risk}_{jnt} + \beta_{7}^{\text{COPD}} 4\%\_\text{osteoporosis\_risk}_{jnt} + \beta_{8}^{\text{COPD}} \text{once\_daily}_{jnt} + \\ \beta_{9}^{\text{COPD}} & \text{canister}_{jnt} + \beta_{10}^{\text{COPD}} \text{discharge}_{jnt} + \beta_{11}^{\text{COPD}} 1 \text{to2\_easy\_steps}_{jnt} + \beta_{12}^{\text{COPD}} \text{soft\_mist}_{jnt} + \\ \beta_{13}^{\text{COPD}} & \text{pressurized\_device}_{jnt} + \beta_{14}^{\text{COPD}} 1 \text{to2\_easy\_steps}_{jnt} + \\ \beta_{15}^{\text{COPD}} & \text{count\_evey\_10th\_dose}_{jnt} + \beta_{16}^{\text{COPD}} \text{count\_evey\_dose}_{jnt} + \\ \beta_{17}^{\text{COPD}} & \text{pneumonia\_risk}_{jnt} + \beta_{18}^{\text{COPD}} 15\%\_\text{pneumonia\_risk}_{jnt}) + \varepsilon_{jnt} \end{split}
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where  $\alpha_{\text{left}}$  was the constant of the left alternative to control for left-right bias,  $\beta_1$  to  $\beta_{18}$  are marginal utilities (i.e. estimated preference parameters) and  $\varepsilon_{jnt}$  was an extreme value distributed error that allowed the function to be estimated in a multinomial logit model.

All attributes were dummy-coded. Reference levels are indicated in Supplemental Table 1. Each of the estimated marginal utilities measured respondents' sensitivity to deviations from the reference level of the corresponding attribute. The sign (+ or -) of a marginal utility denoted whether patients valued this deviation positively or negatively. Three behavioral output measures were calculated to explore the trade-offs that patients are willing to make:

- Relative attribute importance (RI) was estimated as the range of the marginal utilities
  within an attribute (i.e., the difference between the highest valued attribute and the lowest
  valued attribute), divided by the sum of the attribute ranges. RI scores sum up to one
  across each attribute and measure variation in overall utility that can be explained by
  changes in each attribute.
- 2. The maximum acceptable exacerbations (MAE) measured how many additional exacerbations a respondent was willing to accept for each of the attribute levels, relative to their respective reference level. This measure was obtained by estimating the baseline utility function with the number of exacerbations being coded as linear (i.e. one marginal utility is estimated instead of  $\beta_1$  and  $\beta_2$ ). Each marginal utility was then divided by the negative of the obtained marginal utility of the number of exacerbations.
- 3. The maximum acceptable onset time (MAO) measured how many extra minutes of onset of action patients were willing to accept for each of the attribute levels, relative to their

respective reference level. This measure was obtained by estimating the baseline utility function with the onset of action being coded as linear (i.e. one marginal utility is estimated instead of  $\beta_3$  to  $\beta_5$ ). Each marginal utility was then divided by the negative of the obtained marginal utility of the onset of action attribute.

The linearity assumptions made by MAE and MAO were tested by estimating the baseline utility function and fitting a linear function through the marginal utilities of the dummy coded attributes. Linearity was rejected for an  $R^2$  <0.9. Standard errors of MEA and MAO were obtained using the delta method.

#### ASSESSMENT OF INTERNAL VALIDITY

Internal validity indicators included:

- Not choosing the better option in the dominated-choice question
- Choosing a different answer in the repeated question
- Always choosing the alternative better on one of the attributes with unambiguous attribute ordering (all attributes apart from method of treatment)
- Always choosing A or B; in addition, the discrete choice models included a constant to
  assess biases that are not binary (i.e., choosing sometimes A, sometimes B, but choosing
  A more often than they should, based on the level differences)
- Assessing time to complete the DCE survey

Internal validity was assessed for the full sample and stratified by educational level and health literacy and numeracy. Level of education was dichotomized into a high and low score, whereby bachelor's and/or postgraduate degrees were defined as a high educational level and all other educational levels are defined as a low educational level. Health literacy and numeracy were dichotomized as recommended by their developers.[5, 6] Results are presented in **Supplemental Table 2**.

# **Supplemental Table 2. Internal Validity Assessment Results**

	n (%)								_
Validity assessment	Overall	Asthma	COPD	Low Literacy	High Literacy	Low Numeracy	High Numeracy	Low Education	High Education
	(N=1957)	(N=810)	(N=1147)	(N=295)	(N=1662)	(N=228)	(N=1729)	(N=1107)	(N=850)
Dominance test									_
Failed	226 (12)	133 (16)	93 (8)	98 (33)	128 (8)	94 (41)	132 (8)	157 (14)	69 (8)
Passed	1,731 (88)	677 (84)	1,054 (92)	197 (67)	1,534 (92)	134 (59)	1,597 (92)	950 (86)	781 (92)
Repeated question									
Failed	410 (21)	155 (19)	255 (22)	74 (25)	336 (20)	55 (24)	355 (21)	251 (23)	159 (19)
Passed	1,547 (79)	655 (81)	892 (78)	221 (75)	1,326 (80)	173 (76)	1,374 (79)	856 (77)	691 (81)
Always choosing A or									
В									
A	40 (2)	35 (4)	5 (0)	29 (10)	11 (1)	34 (15)	6 (0)	36 (3)	4(0)
В	1 (0)	1 (0)	0 (0)	0 (0)	1 (0)	0 (0)	1 (0)	0 (0)	1 (0)
Neither (some A,	1,916 (98)	774 (96)	1,142 (100)	266 (90)	1,650 (99)	194 (85)	1,722 (100)	1,071 (97)	845 (99)
some B)	1,910 (90)	774 (90)	1,142 (100)	200 (90)	1,030 (99)	194 (63)	1,722 (100)	1,071 (97)	043 (99)
Always choosing the									
alternative better on									
one attribute									
None	1,593 (81)	673 (83)	920 (80)	239 (81)	1,354 (81)	200 (88)	1,393 (81)	929 (84)	664 (78)
Exacerbations	160 (8)	58 (7)	102 (9)	27 (9)	133 (8)	11 (5)	149 (9)	75 (7)	85 (10)
Onset of action	150 (8)	52 (6)	98 (9)	21 (7)	129 (8)	9 (4)	141 (8)	71 (6)	79 (9)
Osteoporosis	31 (2)	26 (3)	5 (0)	5 (2)	26 (2)	7 (3)	24 (1)	15 (1)	16 (2)
Pneumonia	23 (1)	1 (0)	22 (2)	3 (1)	20 (1)	1 (0)	22 (1)	17 (2)	6 (1)
Time to complete									
survey (min:s)									
3:00-4:59	1 (0)	1 (0)	0 (0)	1 (0)	0 (0)	1 (0)	0 (0)	1 (0)	0 (0)
5:00-6:59	33 (2)	25 (3)	8 (1)	19 (6)	14 (1)	14 (6)	19 (1)	22 (2)	11 (1)
7:00-9:59	83 (4)	61 (8)	22 (2)	26 (9)	57 (3)	22 (10)	61 (4)	44 (4)	39 (5)
9:00-14:59	515 (26)	265 (33)	250 (22)	89 (30)	426 (26)	77 (34)	438 (25)	265 (24)	250 (29)
15:00-19:59	427 (22)	166 (20)	261 (23)	47 (16)	380 (23)	40 (18)	387 (22)	252 (23)	175 (21)
≥20:00	898 (46)	292 (36)	606 (53)	113 (38)	785 (47)	74 (32)	824 (48)	523 (47)	375 (44)

Abbreviation: COPD, chronic obstructive pulmonary disease

### **RESULTS OF MULTINOMIAL LOGIT MODELING**

Results of multinomial logit modeling by disease are shown in **Supplemental Tables 3–5**, by asthma severity groups in **Supplemental Table 6**, and by COPD severity groups in **Supplemental Table 7**.

Supplemental Table 3. Multinomial Logit Results (Dummy Coding) by Disease

	Astl	nma (N=810)		CO	PD (N=1147)	
	Marginal	•		Marginal	,	
	utility			utility		
Attribute/level	(coefficient)	95% CI	RI	(coefficient)	95% CI	RI
Left alternative	0.16***	0.12, 0.20	-	0.02	-0.02, 0.06	-
Exacerbations						
3 per year	Referen	ce level			nce level	
2 per year	0.43***	0.37, 0.49	0.21	0.52***	0.47, 0.57	0.27
1 per year	0.68***	0.61, 0.74		1.07***	1.02, 1.13	
Onset time						
30 min	Referen	ce level		Referen	nce level	
20 min	0.38***	0.30, 0.45	0.33	0.46***	0.40, 0.53	0.28
15 min	0.61***	0.53, 0.68	0.55	0.74***	0.67, 0.80	0.28
5 min	1.05***	0.97, 1.13		1.11***	1.04, 1.18	
5-year risk of						
osteoporosis						
6%	Referen	ce level		Referer	nce level	
5%	0.21***	0.15, 0.27	0.14	0.21***	0.15, 0.26	0.08
4%	0.46***	0.39, 0.52		0.32***	0.26, 0.38	
Dosing frequency		,			,	
Twice daily	Referen	ce level	0.05	Referer	nce level	0.05
Once daily	0.15***	0.10, 0.20	0.05	0.21***	0.16, 0.25	0.05
Priming		,			,	
Capsule	Referen	ce level		Referer	nce level	
Canister: two steps	0.39***	0.32, 0.46		0.32***	0.26, 0.38	
Discharge: one step	0.36***	0.28, 0.44	0.13	0.20***	0.12, 0.27	0.08
One to two easy					,	
steps	0.40***	0.32, 0.48		0.32***	0.25, 0.39	
Device type		,			,	
Dry powder inhaler	Referen	ce level		Referei	nce level	
Soft mist inhaler	0.24***	0.18, 0.30	0.11	0.06*	0.01, 0.11	0.04
Pressurized inhaler	0.34***	0.28, 0.41		0.16***	0.11, 0.21	
Dose counter		,			*****	
Metered	Referen	ce level		Referen	nce level	
Every 10 <sup>th</sup> dose	0.03	-0.03, 0.10	0.03	-0.13***	-0.19, -0.08	0.05
Every dose	0.08*	0.02, 0.15	0.00	0.07*	0.02, 0.13	0.00
5-year risk of	0.00	0.02, 0.12		0.07	0.02, 0.12	
pneumonia						
20%				Refere	nce level	
15%	N	ot included		0.268***	0.21, 0.32	0.15
10%	11	or meradea		0.603***	0.55, 0.66	0.15
Log-likelihood		-6027		0.003	-8220	
McFadden's pseudo-		-0027			-0220	
R <sup>2</sup>		0.11			0.14	
IX		0.11			U.14	

Abbreviations: COPD, chronic obstructive pulmonary disease; RI, relative importance score; SE, standard error. \*P<0.05. \*\*P<0.01. \*\*\*P<0.001.

### Supplemental Table 4. Multinomial Logit Results (Continuous Exacerbation Coding) by Disease

		Asthma (N	N=810)			CO	PD (N=1147)	
Attribute/level	Coefficient	95% CI	MAE	95% CI	Coefficient	95% CI	MAE	95% CI
Left alternative	0.16***	0.11, 0.20	-	-	0.02	-0.02, 0.06	-	-
Exacerbations (1 per year)	-0.34***	-0.37, -0.31	-	-	-0.54***	-0.57, -0.51	-	-
Onset time								
30 min		Reference	e level			Re	ference level	
20 min	0.37***	0.29, 0.45	1.1	0.8, 1.3	0.46***	0.40, 0.53	0.9	0.7, 1.0
15 min	0.60***	0.53, 0.68	1.8	1.5, 2.0	0.74***	0.67, 0.80	1.4	1.2, 1.5
5 min	1.05***	0.97, 1.13	3.1	2.7, 3.4	1.11***	1.04, 1.18	2.1	1.9, 2.2
5-year risk of osteoporosis								
6%		Reference	e level			Re	ference level	
5%	0.21***	0.15, 0.27	0.6	0.4, 0.8	0.21***	0.15, 0.26	0.4	0.3, 0.5
4%	0.45***	0.38, 0.52	1.3	1.1, 1.5	0.32***	0.26, 0.38	0.6	0.5, 0.7
Dosing frequency		•				•		•
Twice daily		Reference	e level			Re	ference level	
Once daily	0.15***	0.10, 0.20	0.4	0.3, 0.6	0.21***	0.16, 0.25	0.4	0.3, 0.5
Priming								
Capsule		Reference	e level			Re	ference level	
Canister: Two steps	0.38***	0.31, 0.46	1.1	0.9, 1.4	0.32***	0.26, 0.37	0.6	0.5, 0.7
Discharge: One step	0.36***	0.28, 0.44	1.1	0.8, 1.3	0.20***	0.12, 0.27	0.4	0.2, 0.5
One to two easy steps	0.40***	0.32, 0.47	1.2	0.9, 1.4	0.32***	0.25, 0.39	0.6	0.5, 0.7
Device type								
Dry powder inhaler		Reference	e level			Re	ference level	
Soft mist inhaler	0.24***	0.18, 0.31	0.7	0.5, 0.9	0.06*	0.01, 0.11	0.1	0.0, 0.2
Pressurized inhaler	0.34***	0.28, 0.41	1.0	0.8, 1.2	0.16***	0.11, 0.21	0.3	0.2, 0.4
Dose counter		•				•		•
Metered		Reference	e level			Re	ference level	
Every 10th dose	0.04	-0.02, 0.10	0.1	-0.1, 0.3	-0.13***	-0.19, -0.08	-0.2	-0.3, -0.1
Every dose	0.08*	0.02, 0.15	0.2	0.1, 0.4	0.07**	0.02, 0.13	0.1	0.0, 0.2
5-year risk of pneumonia								
20%						Re	ference level	
15%		Not incl	uded		0.27***	0.21, 0.32	0.5	0.4, 0.6
10%					0.60***	0.58, 0.66	1.1	1.0, 1.2
Log-likelihood		-603	3				-8220	
McFadden's pseudo-R <sup>2</sup>		0.11					0.14	

Abbreviations: COPD, chronic obstructive pulmonary disease; MAE, maximum acceptable exacerbations; SE, standard error.

\*P<0.05. \*\*P<0.01. \*\*\*P<0.001.

Supplemental Table 5. Multinomial Logit Results (Continuous Onset Coding) by Disease

		Asthma (	N=810)			COPD	(N=1147)	
Attribute/level	Marginal utility (coefficient)	95% CI	MAO	95% CI	Marginal utility (coefficient)	95% CI	MAO	95% CI
Left alternative	0.16***	0.12, 0.20	-	-	0.02	-0.02, 0.06	-	-
Exacerbations								
3 per year		Reference	ce level			Refere	nce level	
2 per year	0.43***	0.37, 0.49	10.1	8.6, 11.7	0.52***	0.47, 0.57	11.6	10.3, 12.9
1 per year	0.68***	0.61, 0.74	16.0	14.2, 17.7	1.07***	1.02, 1.13	24.0	22.3, 25.7
Onset time (min)	-0.04***	-0.05, -0.04	-	-	-0.04***	-0.05, -0.04	-	-
5-year risk of osteoporosis								
6%		Reference	e level			Refere	nce level	
5%	0.21***	0.15, 0.27	4.9	3.5, 6.4	0.20***	0.15, 0.26	4.5	3.4, 5.7
4%	0.46***	0.39, 0.52	10.8	9.2, 12.4	0.32***	0.26, 0.38	7.1	5.9, 8.4
Dosing frequency								
Twice daily		Reference	e level			Refere	nce level	
Once daily	0.15***	0.10, 0.20	3.6	2.4, 4.8	0.20***	0.16, 0.25	4.6	3.6,5.6
Priming								
Capsule		Reference	ce level			Refere	nce level	
Canister: two steps	0.39***	0.32, 0.46	9.2	7.4, 11.0	0.32***	0.26, 0.38	7.1	5.7, 8.4
Discharge: one step	0.36***	0.28, 0.44	8.5	6.6, 10.4	0.20***	0.12, 0.27	4.4	2.8, 6.1
One to two easy steps	0.40***	0.32, 0.48	9.4	7.5, 11.3	0.32***	0.25, 0.40	7.2	5.6, 8.9
Device type								
Dry powder inhaler		Reference	e level			Refere	nce level	
Soft mist inhaler	0.24***	0.18, 0.30	5.6	4.1, 7.1	0.06*	0.01, 0.11	1.4	0.21, 2.6
Pressurized inhaler	0.34***	0.28, 0.40	8.1	6.5, 9.6	0.16***	0.11, 0.22	3.6	2.4, 4.8
Dose counter								
Metered		Reference	e level			Refere	nce level	
Every 10th dose	0.03	-0.03, 0.09	0.7	-0.7, 2.2	-0.13***	-0.19, -0.08	-3.0	-4.2, -1.8
Every dose	0.08*	0.02, 0.14	1.9	0.4, 3.4	0.07*	0.01, 0.13	1.6	0.3, 2.8
5-year risk of pneumonia								
20%						Refere	nce level	
15%		Not inc	luded		0.27***	0.21, 0.32	6.0	4.8, 7.2
10%					0.61***	0.55, 0.66	13.5	12.2, 14.9
Log-likelihood		-67:	37			-9	540	
McFadden's pseudo-R <sup>2</sup>		0.1					.14	

Abbreviations: COPD, chronic obstructive pulmonary disease; SE, standard error.

\*P<0.05. \*\*P<0.01. \*\*\*P<0.001.

Supplemental Table 6. Multinomial Logit Results (Dummy Coding): Asthma by ACQ Groups

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	ACQ:	≤0.75 (N=219)		0.75 <a< th=""><th>CQ &lt;1.5 (N=19</th><th>5)</th><th>A</th><th>CQ ≥1.5 (N=396)</th><th></th></a<>	CQ <1.5 (N=19	5)	A	CQ ≥1.5 (N=396)	
	Marginal	, ,		Marginal	•		Marginal	,	
Attribute/level	utility	95% CI	RI	utility	95% CI	RI	utility	95% CI	RI
	(coefficient)			(coefficient)			(coefficient)		
Left alternative	0.02	-0.07, 0.10	-	0.10*	0.01, 0.19	-	0.28***	0.21, 0.34	-
Exacerbations									
3 per year	Referen	ce level		Referen	ice level		Refere	nce level	
2 per year	0.53***	0.41, 0.66	0.22	0.44***	0.32, 0.57	0.25	0.38***	0.28, 0.47	0.19
1 per year	0.94***	0.80, 1.07		0.73***	0.60, 0.86		0.54***	0.45, 0.63	
Onset time									
30 min	Referen	ce level		Referen	ice level		Refere	nce level	
20 min	0.31***	0.15, 0.46	0.24	0.30***	0.15, 0.46	0.34	0.46***	0.35, 0.57	0.39
15 min	0.60***	0.45, 0.76	0.24	0.49***	0.33, 0.64	0.34	0.68***	0.57, 0.79	0.39
5 min	1.02***	0.86, 1.18		1.00***	0.84, 1.17		1.13***	1.02, 1.25	
5-year risk of osteoporosis									
6%	Referen	ce level		Referen	ice level		Refere	nce level	
5%	0.39***	0.26, 0.51	0.18	0.19**	0.06, 0.31	0.10	0.13**	0.05, 0.22	0.13
4%	0.77***	0.64, 0.91		0.31***	0.17, 0.44		0.38***	0.28, 0.48	
Dosing frequency									
Twice daily	Referen	ce level	0.08	Referen	ice level	0.04	Refere	nce level	0.03
Once daily	0.33***	0.22, 0.43	0.08	0.11*	0.01, 0.22	0.04	0.08*	0.01, 0.15	0.03
Priming									
Capsule	Referen	ce level		Referen	ice level		Refere	nce level	
Canister: two steps	0.53***	0.39, 0.67	0.15	0.42***	0.27, 0.56	0.15	0.31***	0.21, 0.41	0.11
Discharge: one step	0.55***	0.40, 0.71	0.15	0.44***	0.28, 0.60	0.15	0.24***	0.12, 0.35	0.11
One to two easy steps	0.63***	0.48, 0.79		0.46***	0.31, 0.62		0.26***	0.15, 0.37	
Device type									
Dry powder inhaler	Referen	ce level		Referen	ice level		Refere	nce level	
Soft mist inhaler	0.22***	0.11, 0.34	0.09	0.25***	0.12, 0.37	0.10	0.25***	0.17, 0.34	0.12
Pressurized inhaler	0.38***	0.26, 0.50		0.29***	0.17, 0.42		0.36***	0.27, 0.45	
Dose counter									
Metered	Referen	ce level		Referen	ice level		Refere	nce level	
Every 10th dose	0.08	-0.05, 0.20	0.05	-0.03	-0.15, 0.10	0.02	0.05	-0.04, 0.13	0.03
Every dose	0.21**	0.08, 0.34		-0.07	-0.20, 0.06		0.09*	0.00, 0.19	
Log-likelihood		-1580			-1445			-2946	
McFadden's pseudo-R <sup>2</sup>		0.13			0.11			0.11	

Abbreviations: ACQ, Asthma Control Questionnaire®; RI, relative importance score; SE, standard error. \*P<0.05. \*\*P<0.01. \*\*\*P<0.001.

# Supplemental Table 7. Multinomial Logit Results (Dummy Coding): COPD by CAT Groups

Left alternative  Exacerbations 3 per year	Marginal utility oefficient) -0.04 Referen	<b>95% CI</b> -0.11, 0.03	RI	Marginal utility (coefficient)			Marginal utility	, ,	
Left alternative  Exacerbations 3 per year	-0.04		RI	(coefficient)			aciii,		
Exacerbations 3 per year		-0.11, 0.03		(Journal of the Control of the Contr	95% CI	RI	(coefficient)	95% CI	RI
3 per year	Referer		-	0.01	-0.05, 0.07	-	0.08*	0.02, 0.15	-
1 2	Referer								
	recreici	nce level		Referen	ice level		Referen	ce level	
2 per year	0.60***	0.50, 0.71	0.27	0.48***	0.39, 0.56	0.24	0.51***	0.41, 0.61	0.27
1 per year	1.32***	1.21, 1.43		1.00***	0.91, 1.09		0.96***	0.86, 1.07	
Onset time									
30 min	Referer	nce level			ice level		Referen	ce level	
20 min	0.45***	0.32, 0.57	0.20	0.46***	0.36, 0.56	0.28	0.48***	0.36, 0.60	0.32
15 min	0.65***	0.52, 0.77	0.20	0.78***	0.68, 0.88	0.28	0.77***	0.65, 0.90	0.32
5 min	0.98***	0.84, 1.11		1.19***	1.08, 1.30		1.16***	1.03, 1.29	
5-year risk of osteoporosis									
6%	Referer	nce level			ice level		Referen	ce level	
5%	0.18***	0.07, 0.28	0.07	0.23***	0.15, 0.32	0.08	0.20***	0.10, 0.29	0.07
4%	0.36***	0.25, 0.47		0.33***	0.24, 0.42		0.26***	0.16, 0.37	
Dosing frequency									
Twice daily		nce level	0.07	Referen	ice level	0.05	Referen	ce level	0.02
Once daily	0.34***	0.25, 0.43	0.07	0.20***	0.13, 0.27	0.03	0.09*	0.00, 0.17	0.02
Priming									
Capsule	Referer	nce level		Referen	ice level		Referen	ce level	
Canister: two steps	0.46***	0.35, 0.58	0.09	0.32***	0.28, 0.41	0.08	0.18***	0.08, 0.29	0.08
Discharge: one step	0.34***	0.20, 0.49	0.09	0.15*	0.03, 0.26	0.08	0.13	-0.01, 0.27	0.08
One to two easy steps	0.41***	0.28, 0.55		0.29***	0.17, 0.40		0.28***	0.14, 0.42	
Device type									
Dry powder inhaler	Referer	nce level		Referen	ice level		Referen	ce level	
Soft mist inhaler	0.07	-0.03, 0.17	0.03	0.10*	0.02, 0.18	0.05	0.00	-0.09, 0.10	0.04
Pressurized inhaler	0.14**	0.04, 0.24		0.21***	0.13, 0.29		0.13**	0.03, 0.23	
Dose counter									
Metered	Referer	nce level		Referen	ice level		Referen	ce level	
Every 10th dose -	0.18***	-0.28, -0.08	0.04	-0.09*	-0.17, -0.01	0.02	-0.15**	-0.25, -0.05	0.04
Every dose	0.10	-0.00, 0.21		0.09*	0.00, 0.18		0.02	-0.08, 0.12	
5-year risk of pneumonia		,			,			<del>,</del> <del>-</del>	
20%	Referer	nce level	0.24	Referen	ice level	0.21	Referen	ce level	0.16
	0.36***	0.26, 0.46	0.24	0.28***	0.20, 0.37		0.16**	0.06, 0.26	0.16

10%	0.83*** 0.73, 0.94	0.58*** 0.50, 0.67	0.42*** 0.32, 0.53
Log-likelihood	-2299	-3423	-2445
McFadden's pseudo-R	0.17	0.14	0.13

Abbreviations: CAT, COPD Assessment Test; COPD, chronic obstructive pulmonary disease; RI, relative importance score; SE, standard error. \*P<0.05. \*\*P<0.01. \*\*\*P<0.001.

## Supplemental Table 8. Multinomial Logit Results for Asthma by Age, Sex, Level of Education, and ACQ Score

	Coefficient (95% CI) University degree								
			or higher vs. less						
				than university	0.75 < ACQ < 1.5  vs.	ACQ ≥1.5 vs. ACQ			
Attribute/level	Marginal utility	≥65 y vs. <65 y	Female vs. male	degree	ACQ ≤0.75 or ≥1.5	<1.5			
Left alternative	0.17 (0.13, 0.22)***								
Exacerbations									
3 per year	Reference level								
2 per year	0.36 (0.17, 0.54)***	0.13 (-0.03, 0.30)	0.14 (0.01, 0.27)*	0.13 (0.00, 0.26)	-0.08 (-0.26, 0.10)	-0.14 (-0.30, 0.20)			
1 per year	0.66 (0.47, 0.85)***	0.14 (-0.03, 0.32)	0.21 (0.07, 0.35)**	0.21 (0.08, 0.35)**	-0.18 (-0.37, 0.01)	-0.35 (0.52, -0.18)***			
Onset time									
30 min	Reference level								
20 min	0.19 (-0.04, 0.41)	-0.02 (-0.23, 0.18)	0.05 (-0.11, 0.21)	0.19 (0.03, 0.35)*	0.00 (-0.22, 0.23)	0.16 (-0.04, 0.36)			
15 min	0.42 (0.19, 0.64)***	-0.08 (-0.28, 0.12)	0.13 (-0.03, 0.29)	0.25 (0.09, 0.41)**	-0.11 (-0.33, 0.11)	0.09 (-0.10, 0.29)			
5 min	0.47 (0.23, 0.70)***	0.10 (-0.12, 0.31)	0.34 (0.17, 0.51)***	0.52 (0.35, 0.69)***	0.06 (-0.17, 0.30)	0.23 (0.02, 0.43)*			
5-year risk of osteoporosis									
6%	Reference level								
5%	0.28 (0.09, 0.46)**	0.03 (-0.13, 0.20)	0.15 (0.02, 0.29)*	0.02 (-0.11, 0.15)	-0.19 (-0.37,-0.02)*	-0.25 (-0.42, -0.09)***			
4%	0.66 (0.47, 0.86)***	0.01 (-0.17, 0.19)	0.09 (-0.05, 0.24)	0.11 (-0.03, 0.25)	-0.45 (-0.65, -0.26)***	-0.39 (-0.57, -0.22)*			
Dosing frequency									
Twice daily	Reference level								
Once daily	0.19 (0.04, 0.34)*	0.10 (-0.04, 0.24)	0.05 (-0.06, 0.16)	0.14 (0.04, 0.25)**	-0.20 (-0.35, -0.05)	-0.20 (-0.34, -0.07)			
Priming									
Capsule	Reference level								
Canister: Two-Steps	0.29 (0.08, 0.49)**	0.29 (0.10, 0.48)**	0.14 (-0.01, 0.30)	0.12 (-0.03, 0.27)	-0.07 (-0.28, 0.14)	-0.15 (-0.34, 0.03)			
Discharge: One-Step	0.34 (0.10, 0.58)**	0.27 (0.06, 0.49)*	0.11 (-0.05, 0.28)	0.11 (-0.06, 0.28)	-0.07 (-0.30, 0.16)	-0.25 (-0.46, -0.05)*			
One to two easy steps	0.47 (0.24, 0.71)***	0.08 (-0.13, 0.28)	0.06 (-0.10, 0.23)	0.19 (0.02, 0.35)*	-0.15 (-0.37, 0.08)	-0.34 (-0.54, -0.14)***			
Device type									
Dry powder inhaler	Reference level								
Soft mist inhaler	0.11 (-0.07, 0.28)	0.04 (-0.12, 0.20)	0.13 (0.00, 0.26)*	0.06 (-0.07, 0.19)	0.03 (-0.14, 0.21)	0.03 (-0.12, 0.19)			
Pressurized inhaler	0.25 (0.06, 0.43)**	0.08 (-0.08, 0.25)	0.17 (0.04, 0.30)*	0.06 (-0.07, 0.19)	-0.10 (-0.28, 0.08)	-0.03 (-0.19, 0.13)			
Dose counter									
Metered	Reference level								
Every 10th dose	0.10 (-0.08, 0.28)	-0.02 (-0.19, 0.14)	-0.01 (-0.14, 0.12)	-0.02 (-0.15, 0.11)	-0.10 (-0.27, 0.08)	-0.04 (-0.20, 0.12)			
Every dose	0.09 (-0.09, 0.28)	0.26 (0.10, 0.43)**	0.02 (-0.11, 0.16)	0.04 (-0.09, 0.17)	-0.27 (-0.46, -0.09)**	-0.07 (-0.23, 0.09)			

Abbreviations: ACQ, Asthma Control Questionnaire®; CI, confidence interval

# Supplemental Table 9. Multinomial Logit Results for COPD by Age, Sex, Level of Education, and CAT score

	Coefficient (95% CI)								
Attribute/level	Marginal Utility	≥65 y vs. <65 y	Female vs. male	University degree or higher vs. less than university degree	20 < CAT < 30 vs. CAT ≤20 or ≥30	CAT ≥30 vs. CAT <30			
Left alternative	0.02 (-0.02, 0.06)	<u>_000 y 150 100 y</u>	1 chare vs. mare	uegree	_50	100			
Exacerbations	0.02 ( 0.02, 0.00)								
3 per year				Reference level					
2 per year	0.43 (0.28, 0.59)***	0.02 (-0.11, 0.14)	0.18 (0.07, 0.30)**	0.16 (0.04, 0.27)**	-0.12 (-0.25, 0.02)	-0.09 (-0.25, 0.06)			
1 per year	1.04 (0.88, 1.21)***	0.15 (0.02, 0.28)	0.23 (0.09, 0.33)***	0.23 (0.10, 0.35)***	-0.28 (-0.43, -0.13)***	-0.32 (-0,48, -0.16)***			
Onset time									
30 min				Reference level					
20 min	0.40 (0.22, 0.58)***	-0.12 (-0.27, 0.03)	0.09 (0.04, 0.23)	0.16 (0.02, 0.30)*	0.00 (-0.16, 0.17)	-0.02 (-0.21, 0.16)			
15 min	0.57 (0.38, 0.76)***	=0.17 (-0.32, -0.01)*	0.14 (0.00, 0.28)*	0.20 (0.05, 0.34)**	0.12 (-0.05, 0.29)	0.06 (-0.13, 0.25)			
5 min	0.98 (0.78, 1.18)***	-0.27 (-0.43, -0.10)**	0.04 (-0.11, 0.19)	0.30 (0.15, 0.46)***	0.18 (0.0, 0.36)	0.06 (-0.14, 0.26)			
5-year risk of osteoporosis									
6%				Reference level					
5%	0.18 (0.30, 0.33)*	-0.02 (-0.14, 0.11)	-0.03 (-0.14, 0.09)	0.03 (-0.08, 0.15)	0.07 (-0.07, 0.20)	0.02 (-0.13, 0.17)			
4%	0.37 (0.20, 0.53)***	-0.08 (0.02, 0.24)	-0.10 (-0.22, 0.02)	0.03 (-0.09, 0.16)	0.00 (-0.14, 0.15)	-0.06 (-0.23, 0.10)			
Dosing frequency									
Twice daily				Reference level					
Once daily	0.26 (0.13, 0.39)***	0.13 (0.02, 0.24)*	0.04 (-0.05, 0.14)	-0.02 (-0.11, 0.08)	-0.12 (-0.23, 0.00)	-0.20 (-0.33, -0.07)**			
Priming									
Capsule				Reference level					
Canister: Two-Steps	0.42 (0.25, 0.59)***	0.11 (-0.03, 0.25)	-0.04 (-0.17, 0.09)	0.00 (-0.13, 0.13)	-0.12 (-0.27, 0.04)	-0.21 (-0.38, -0.05)*			
Discharge: One-Step	0.41 (0.19. 0.63)***	-0.01 (-0.18, 0.17)	-0.03 (-0.19. 0.13)	-0.12 (-0.29. 0.04)	-0.20 (-0.40, -0.01)*	-0.20 (-0.42, 0.02)			
One to two easy steps	0.52 (0.30, 0.73)***	-0.14 (-0.31, 0.03)	0.02 (-0.14, 0.17)	-0.08 (-0.24, 0.08)	-0.017 (-0.36, 0.02)	-0.19 (-0.40, 0.02)			
Device type									
Dry powder inhaler				Reference level					
Soft mist inhaler	0.09 (-0.06, 0.24)	-0.01 (-0.13, 0.11)	0.04 (-0.07, 0.15)	-0.12 (-0.23, -0.01)*	0.02 (-0.11, 0.15)	-0.05 (-0.20, 0.10)			
Pressurized inhaler	0.21 (0.06, 0.36)**	-0.01 (-0.13, 0.11)	-0.16 (-0.17, 0.05)	-0.11 (-0.22, 0.00)	0.07 (-0.07, 0.20)	0.00 (-0.15, 0.15)			
Dose counter									
Metered				Reference level					
Count every 10th dose	-0.22 (-0.38, -0.07)**	0.01 (-0.11, 0.14)	0.01 (-0.10, 0.12)	0.04 (-0.07, 0.16)	0.11 (-0.02, 0.25)	0.05 (-0.10, 0.20)			
Count every dose	0.02 (-0.14, 0.18)	0.09 (-0.04, 0.22)	0.03 (-0.08, 0.15)	0.05 (-0.07, 0.17)	0.01 (-0.13, 0.16)	-0.05 (-0.20, 0.11)			
5-year risk of pneumonia 20%				Reference level					
15%	0.42 (0.27, 0.57)***	-0.02 (-0.15, 0.10)	-0.08 (-0.19, 0.03)	0.00 (-0.12, 0.11)	-0.08 (-0.21, 0.06)	-0.21 (-0.36, -0.06)**			
10%	0.86 (0.70, 1.02)***	0.05 (-0.08, 0.18)	-0.11 (-0.23, 0.01)	0.04 (-0.08, 0.16)	-0.023 (-0.37, -0.09)	-0.40 (-0.56, -0.24)***			

Abbreviations: CAT, COPD Assessment Test; CI, confidence interval; COPD, chronic obstructive pulmonary disease

Supplemental material

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