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95% CI

Tobacco Heating System 2.2 Use in Mild to Moderate Chronic **Obstructive Pulmonary Disease Subjects Switching from Cigarettes: An Exploratory Analysis**

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Purpose

The Tobacco Heating System (THS) is a product that electronically heats tobacco at temperatures significantly lower (<350°C) than in combusted cigarettes (CC), producing substantially lower levels of harmful and potentially harmful constituents, while providing nicotine and a ritualistic and sensorial experience that approach that of CC.

A randomized, two-arm parallel group, multicenter "Exposure Response Study" (ERS) conducted in the United States (US) (NCT02396381) demonstrated statistically significant favorable changes in five of eight biomarkers of effect (BoEff) at 6 months among adult smokers (not willing to quit smoking) switching to THS 2.2 compared with those who continued smoking, with all BoEffs moving in the direction of smoking cessation (SC). The study was extended for an additional 6 months (NCT02649556) for the selected BoEffs.

In parallel, a 12-month Study "Smoking Cessation Response" (SCR) was conducted among smokers willing to quit smoking in the US, Europe, and Japan (NCT02432729). All studies were conducted in accordance with the International Council for Harmonisation of Technical Requirements for Pharmaceuticals for Human Use (ICH) and Good Clinical Practice guidelines (Figure 1).

This analysis explored the health effects of switching to THS 2.2 compared with those of continued smoking in a subgroup of smokers with mild to moderate chronic obstructive pulmonary disease (COPD) enrolled in the ERS study relative to those enrolled in the SCR.

Figure 1. 6-Month + 6-Month Exposure Response and Smoking Cessation Response Studies



Pre-BD FEV₁ (%pred) Month 6 -0.02, 2.00 eneral Study Populatio 0.99 OPD (Previous ATS)* 1.47 -0.52, 3.46 OPD (GOLD 2018) subjects 0.71 2.97, 4.39 Month 12 neral Study Population 1.16 -0.06. 2.39 OPD (Previous ATS)* 1.92 -0.49, 4.32 COPD (GOLD 2018) subjects 3.48 -1.12, 8.07 Pre-BD Best FEV₁ (mL)

LS Mean

Table 2: Results of Pre-Bronchodilator Spirometry

THS Use - CC Use Difference

	Month 6						
General Study Population	30.1	-6.99, 67.1					
COPD (Previous ATS)*	46.4	-27.5, 120					
COPD (GOLD 2018) subjects	11.1	-125, 147					
Month 12							
General Study Population	40.7	-2.15, 83.5					
COPD (Previous ATS)*	60.2	-27.9, 148					
COPD (GOLD 2018) subjects	71.8	-101, 244					
Note: Adjusted least squares (LS) means and confidence intervals (LS) from a mode model analysis conducted on original values with visit, baseline values and its interaction with visit, and baseline smoking intervals values difficult sources and the interaction with visit, and baseline smoking intervals value difficult curve difficult sources and the interaction with visit, and baseline smoking intervals values are difficult and with an and an effect factor. BD & benchroliditor, short-scaling beta agoinst.							

Methods

The analysis included mild to moderate COPD subjects (GOLD 1 & 21) from the ERS, which was broadened to include airway obstruction defined by FEV₁/FVC <0.75 (previous ATS guidelines²), described as COPD subjects, as per the Lung Health Study³. The results from a subset of subjects with mild COPD from the SCR, using the same definition for airway obstruction as the ERS, were used as the reference.



The subset of COPD subjects randomized to the THS 2.2 or CC arm from the ERS and subjects from the SCR were analyzed at 6 and 12 months. The selected respiratory and cardiovascular BoEffs in this subgroup analysis were forced expiratory volume in one second (FEV₁), white blood cell (WBC) count, soluble intercellular adhesion molecule 1 (sICAM-1) level, and 8-epiprostaglandin F2α (8-epi-PGF2α) level.

Results

Table 1: Baseline Characteristics

Variable	THS Use		CC Use		sc		
	General Population	Obstruction Subset	General Population	Obstruction Subset	General Population	Obstruction Subset	
	(n = 230)	(n = 52)	(n = 424)	(n = 97)	(n = 358)	(n = 53)	
Male: n (%)	143 (62.2)	35 (67.3)	244 (57.5)	63 (64.9)	181 (50.6)	28 (52.8)	
Female: n (%)	87 (37.8)	17 (32.7)	180 (42.5)	34 (35.1)	177 (49.4)	25 (47.2)	
Age (years): Mean (SD)	43.8 (9.68)	48.5 (10.3)	45.2 (9.54)	52.0 (10.1)	43.8 (9.21)	49.8 (8.91)	
Caucasian: n (%)	168 (73.0)	38 (73.1)	312 (73.6)	78 (80.4)	203 (56.7)	39 (73.6)	
Not Caucasian: n (%)	62 (27.0)	14 (26.9)	112 (26.4)	19 (19.6)	155 (43.3)	14 (26.4)	
BMI (kg/m ²): Mean (SD)	27.0 (4.06)	26.2 (4.08)	27.1 (4.13)	25.9 (3.65)	24.8 (3.75)	25.2 (3.14)	
Smoking duration (years): Mean (SD)	25.6 (9.48)	30.8 (9.62)	26.7 (10.1)	33.0 (11.3)	22.8 (8.76)	28.4 (8.57)	
Smoking intensity over the past year (cig/day): Mean (SD)	18.4 (6.88)	18.7 (5.69)	19.5 (7.87)	18.6 (6.87)	16.6 (5.27)	16.8 (5.92)	
COPD (Previous ATS)*: n (%)	35 (15.2)	35 (67.3)	69 (16.3)	69 (71.1)	35 (66.0)	35 (66.0)	
GOLD 1: Mild n (%)	14 (6.1)	14 (26.9)	24 (5.7	24 (24.7)	18 (34.0)	18 (34.0)	
GOLD 2: Moderate n (%)	3 (1.3)	3 (5.8)	4 (0.9)	4 (4.1)	0	0	
Note: THS use refers to THS 2.2 use of 70% or more; CC use refers to THS 2.2 use of less than 1%; SC refers to continuous smoking abstinence from the							

0.70 < FEV₂/FVC < 0.75, according to the previous ATS guidelines for obstruction

Pre-BDR FEV1 WBC from Baseline (95% CI)]

Figure 2: Respiratory and Cardiovascular Biomarkers of Effect in COPD subset





The present results indicated, for the group switching to THS, a favorable THS-CC difference of 72 mL in FEV1 among COPD subjects after 12 months. They also indicated a reduction of 1.95 GI/L in WBC count, 6.8% in sICAM-1 level, and 28% in 8-epi-PGF2α level for THS relative to CC.

FRS

SCR

After 12 months of follow-up, most of the BoEffs in subjects with COPD who used the THS showed favorable shifts in the direction of the changes observed with SC.

The magnitude of these shifts was more pronounced than those observed in the ERS, which included a majority of healthy subjects.

The results of this preliminary analysis are consistent with the findings of the main ERS⁴ and suggest a reduced harm potential for THS relative to smoking in subjects predominantly switching to THS with mild to moderate COPD.

References

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