

Supplementary files

Supplementary Table 1. Comparison of patients with other comorbidities between the SITT and SIDT pre-treatment

| Comorbidities | SITT, n = 52 | SIDT, n = 49 | P-value |
|---------------------------------|--------------|--------------|---------|
| Allergic rhinitis | 22 (42.3) | 16 (32.7) | 0.4 |
| Atopic dermatitis | 2 (3.9) | 1 (2.0) | 1.0 |
| Sinusitis with nasal polyps | 0 (0) | 1 (2.0) | 0.5 |
| ANCA-related disease | 0 (0) | 0 (0) | 1.0 |
| Hypertension | 8 (15.4) | 13 (26.5) | 0.2 |
| Hyperlipidemia | 6 (11.5) | 9 (18.4) | 0.4 |
| Mood disorder | 4 (7.7) | 3 (6.1) | 1.0 |
| Gastroesophageal reflux disease | 5 (9.6) | 1 (2.0) | 0.2 |
| Diabetes | 3 (5.8) | 1 (2.0) | 0.6 |
| Central nervous system disorder | 3 (5.8) | 1 (2.0) | 0.6 |
| Cardiac disease | 2 (3.9) | 1 (2.0) | 1.0 |
| Collagen vascular diseases | 1 (1.9) | 2 (4.1) | 0.6 |
| Arrhythmia | 0 (0) | 1 (2.0) | 0.5 |
| Chronic hepatitis | 1 (1.9) | 1 (2.0) | 1.0 |
| Hypothyroidism | 2 (3.9) | 0 (0) | 0.5 |
| Benign prostatic hyperplasia | 0 (0) | 1 (2.0) | 0.5 |

Notes: No patient had malignancies or chronic kidney diseases.

Abbreviations: ANCA; anti-neutrophil cytoplasmic antibody, SIDT; single-inhaler dual therapy, SITT; single-inhaler triple therapy

Supplementary Table 2. Regimens of controllers in SITT and SIDT in this study

| SITT, n = 52 | | | | |
|---|----------------------|--------------------|-----------------------|----------------------------|
| Controller via device | ICS/LABA/LAMA | Daily doses | Administration | No. (%) of patients |
| Enerzair® via Breezhaler® | | | | 16 (30.8) |
| high-dose | MF/IND/GLY | 160/150/50 µg | once daily | 5 (9.6) |
| medium-dose | MF/IND/GLY | 80/150/50 µg | once daily | 11 (21.2) |
| Trelegy® via Ellipta® | | | | 36 (69.2) |
| high-dose | FF/VI/UMEC | 200/25/62.5 µg | once daily | 10 (19.2) |
| medium-dose | FF/VI/UMEC | 100/25/62.5 µg | once daily | 26 (50.0) |
| SIDT, n = 49 | | | | |
| Controller via device | ICS/LABA | Daily doses | Administration | No. (%) of patients |
| Atecura® via Breezhaler® | | | | 4 (8.2) |
| high-dose | MF/IND | 160/150 µg | once daily | 0 (0) |
| medium-dose | MF/IND | 80/150 µg | once daily | 4 (8.2) |
| Relovair® via Ellipta® | | | | 20 (40.8) |
| high-dose | FF/VI | 200/25 µg | once daily | 19 (38.8) |
| medium-dose | FF/VI | 100/25 µg | once daily | 1 (2.0) |
| Flutiform125® via a pressurized metered dose inhaler | | | | 12 (24.5) |
| high-dose | FP/FF | >625/25 µg | twice daily | 0 (0) |
| medium-dose | FP/FF | 500/20 µg | twice daily | 12 (24.5) |
| Symbicort® via Turbuhaler® | | | | 13 (26.5) |
| high-dose | BUD/FF | 1280/36 µg | twice daily | 1 (2.0) |
| medium-dose | BUD/FF | 640/18 µg | twice daily | 12 (24.5) |

Notes: The percentage (%) of patients was estimated using the subtotal study population (SITT for n = 52 and SIDT for n = 49, respectively).

Abbreviations: BUD; budesonide, FF; fluticasone furoate, FF; formoterol fumarate, FP; fluticasone propionate, GLY; glycopyrronium bromide, IND; indacaterol acetate, MF; mometasone furoate, SIDT; single-inhaler dual therapy, SITT; single-inhaler triple therapy, UMEC; umeclidinium bromide, VI; vilanterol trifenate.

Supplementary Table 3. Relative glucocorticoid receptor-binding affinity among ICS in our study

| Corticosteroids | Relative glucocorticoid receptor-binding affinity compared to dexamethasone |
|------------------------|---|
| Fluticasone furoate | 29.9 |
| Mometasone furoate | 21.0 |
| Fluticasone propionate | 17.8 |
| Budesonide | 9.4 |
| Dexamethasone | 1 |

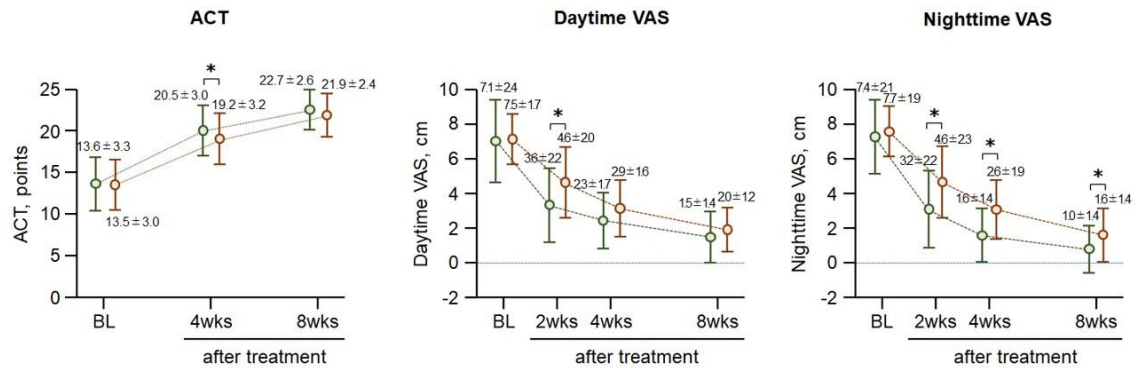
Notes: Glucocorticoid receptor-binding affinity is relative to dexamethasone, where dexamethasone affinity = 1 in accordance with a previous review reference below.

Abbreviations: ICS; inhaled corticosteroid

Reference

Daley-Yates PT. Inhaled corticosteroids: potency, dose equivalence, and therapeutic index. *Br J Clin Pharmacol* 2015; **80**: 372-80.

Supplementary Figure 1.



Notes: All data are expressed as mean \pm standard deviation. The mean ACT scores with SITT were significantly higher than those with SIDT four weeks post-treatment ($P = 0.0443$). The mean daytime VAS scores with SITT were significantly lower than those with SIDT two weeks post-treatment ($P = 0.0280$). In contrast, the nighttime VAS scores with SITT were significantly lower than those with SIDT two ($P = 0.0026$), four ($P = 0.0100$), and eight weeks ($P = 0.0288$) post-treatment.

* $P < 0.05$ when comparing between SITT and SIDT at each visit

Abbreviations: ACT; asthma control test, SIDT; single-inhaler dual therapy, SITT; single-inhaler triple therapy, VAS; visual analog scale, wks; weeks