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Efficacy of mepolizumab for patients with severe asthma and eosinophilic chronic rhinosinusitis

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Abstract

Background: Several major randomized control studies have demonstrated that mepolizumab, an anti-IL-5 monoclonal antibody, is effective for patients with severe eosinophilic asthma who show exacerbation or require systemic corticosteroid maintenance therapy. However, the predictive factors of the response to mepolizumab other than blood eosinophil count are unclear in clinical practice.

Objective: To elucidate the predictive factors of the response to mepolizumab for patients with severe eosinophilic asthma.

Methods: From July 2016 to December 2017, 28 patients with severe asthma received mepolizumab in our hospital. To determine the predictive factors, we retrospectively evaluated patient characteristics, comorbidities, biomarkers, pulmonary function, maintenance dose of systemic corticosteroids and number of exacerbations.

Results: The response rate to mepolizumab treatment was 70% (19/27; one pregnant woman was excluded from analysis). Compared with 11 patients without eosinophilic chronic rhinosinusitis (ECRS), 16 patients with ECRS showed significantly improved systemic corticosteroid-sparing effects [$-71.3 \pm 37.0\%$ vs $-10.7 \pm 20.1\%$, $P = 0.006$], change from baseline FeNO [-19 ± 57 (%) vs 30 ± 77 (%), $P = 0.023$] and symptoms [14 patients (88%) vs five patients (45%), $P = 0.033$]. ECRS was identified as a predictive factor of the response to mepolizumab in a multivariate logistic regression analysis [odds ratio = 22.5, 95% CI (1.5–336), $P = 0.024$]. Of the eight patients previously administered omalizumab, five responded to mepolizumab. *Staphylococcus aureus* enterotoxin B IgE results were negative in 80% of responders ($P = 0.14$).

Conclusion: Both groups showed improved symptom scores and a decreased number of exacerbations. Mepolizumab substantially improved the clinical variables of patients with eosinophilic asthma complicated with ECRS.

Keywords: Asthma, Mepolizumab, Eosinophilic chronic rhinosinusitis, Predictive factor, *Staphylococcus* enterotoxin

Introduction

It is estimated that as many as 300 million people of all ages and ethnic backgrounds suffer from asthma [1]. Asthma is a common chronic respiratory disease that affects approximately 5–10% of Japanese adults [2, 3]. The prevalence of severe and uncontrolled asthma among

adults with asthma has been reported to be 3 to 10% [4]. Severe asthma is associated with morbidity, mortality, and high healthcare costs [5]. Recently, severe asthma was classified into several clinical phenotypes that differ in severity and response to therapy [6]. Severe eosinophilic asthma is characterized by Th2 inflammation, and patients with severe eosinophilic asthma experience recurrent asthma exacerbations even when treated with high doses of inhaled corticosteroids (ICS) and controllers such as long-acting bronchodilators, leukotriene receptor antagonists and oral corticosteroids (OCS) [7].

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Twenty to 25% of patients with difficult-to-control asthma have nasal polyps [8]. Furthermore, patients with asthma and high blood eosinophil counts have poor asthma control [9].

IL-5, which induces the proliferation, differentiation and migration of eosinophils, plays an important role in the pathogenesis of eosinophilic asthma [10]. Mepolizumab, a humanized monoclonal antibody against IL-5, selectively inhibits eosinophilic airway inflammation. Several major randomized control trials (RCTs) have demonstrated that this antibody reduces the number of eosinophils in both sputum and blood, resulting in reductions in exacerbations, improvements in pulmonary function and a glucocorticoid-sparing effect in patients with uncontrolled asthma and a peripheral blood eosinophil count $\geq 150/\text{mm}^3$ at baseline or $\geq 300/\text{mm}^3$ during the previous year [11–13]. Although previous reports have demonstrated the efficacy of this treatment in patients with severe eosinophilic asthma, patients who meet the criteria of eosinophil counts do not always respond to mepolizumab. Accordingly, the identification of predictive factors of a response is very important from a medical economic perspective. However, few reports have examined the predictive factors for mepolizumab; these have been limited to RCTs and their subgroup analyses [11, 14] and a cluster analysis of the response to mepolizumab [15]. The reported main distinctive features of patients with good response are blood eosinophilia $\geq 150/\text{mm}^3$, airway reversibility $> 16.2\%$ and a body mass index (BMI) $\geq 30 \text{ kg/m}^2$ [15]. BMI has been reported to be lower in Japanese patients with asthma than in Caucasian ones [16]. To elucidate the clinical efficacy of mepolizumab in Japanese patients with severe asthma, we performed a single-center, retrospective study.

Methods

Subjects

From July 2016 to December 2017, 28 adult patients with severe uncontrolled asthma received mepolizumab injections (100 mg every 4 weeks) in our hospital. Severe asthma was defined according to the Global Initiative of Asthma (GINA) guidelines and required high doses of ICS plus at least one of the following additional controllers: long-acting beta-agonists (LABA), long-acting muscarinic antagonists (LAMA), leukotriene receptor antagonists (LTRA), xanthine derivative and daily OCS [5]. All patients with asthma were diagnosed by respiratory physicians based on past medical histories of asthma, typical symptoms (cough, wheezing, circadian variation and seasonal episodes) and pulmonary function compatible with asthma. Severity at baseline and during exacerbation was defined according to the GINA guidelines [5]. The present study was approved by the Ethical Committee of Jikei University (28–259 (8502)), and

informed consent was obtained from all subjects. The patient inclusion criteria were as follows: 1. at least two exacerbations requiring OCS treatment in the previous year, and 2. a blood eosinophil count of at least $150/\text{mm}^3$ [12].

We retrospectively examined the following patient characteristics and parameters at baseline and at the last follow-up: gender, age, underlying eosinophilic diseases, smoking status, body mass index, peripheral blood eosinophil count, serum IgE, serum *Staphylococcus aureus enterotoxin* (SE)-IgE level, FeNO, Asthma Control Test (ACT) score, pulmonary function test results [forced volume capacity (FVC), forced expiratory volume in one second (FEV_1), FEV_1/FVC , and $\%\text{FEV}_1$], the severity of asthma according to the GINA guidelines [5], history of previous omalizumab treatment, daily systemic corticosteroid maintenance therapy, the number of exacerbations of asthma symptoms requiring systemic corticosteroids, duration of asthma, number of injections and observation period.

The underlying eosinophilic diseases considered included eosinophilic chronic rhinosinusitis (ECRS), eosinophilic granulomatosis with polyangiitis and eosinophilic otitis media. The FeNO level was measured using a NIOX VERO™ analyzer (Aerocrine AB, Stockholm, Sweden) according to the American Thoracic Society/European Respiratory Society recommendations at a 50 mL/s flow rate [17]. Experienced technicians measured FeNO and remeasured if necessary. We recorded the presence of ECRS or chronic rhinosinusitis with nasal polyps (CRSwNP) as evaluated by computed tomography findings, peripheral blood eosinophil count and the presence of nasal polyps according to the Japanese Epidemiological Survey of Refractory Eosinophilic Chronic Rhinosinusitis (JESREC) study [18] or by pathological findings. We analyzed all patients and compared two groups: patients with ECRS (ECRS group) and those without (non-ECRS group). The levels of IgEs specific to SEs, *Staphylococcus aureus* enterotoxin A (SEA) and *Staphylococcus aureus* enterotoxin B (SEB), were measured with a fluorescence enzyme immunoassay (SRL, Tokyo). The lower limit of SE-IgE detection was 0.10 kU/L. Sensitization was defined as a specific IgE level greater than 0.10 kU/L.

Evaluation of the response to mepolizumab

The mepolizumab response in all cases with inconclusive results were categorized as ineffective. For the remaining cases, we defined a response to mepolizumab as improvement in one or more of the following parameters: change in ACT from baseline to the last follow-up, change in the number of asthma exacerbations per year and reduction of the daily dose of OCSs. The minimal clinically relevant difference was defined as an ACT

score of three points [19]. Systemic corticosteroid doses are shown as prednisone equivalents (mg).

Statistical analysis

All statistical analyses were performed using StatView version 5 (SAS Institute Inc., Cary, NC, USA). All values are expressed as the mean \pm standard deviation. A *P*-value of < 0.05 was considered statistically significant. The factors associated with patient characteristics were examined using Mann-Whitney *U* tests, Fisher's exact tests, or Wilcoxon signed rank tests. Moreover, we re-evaluated several primary outcomes by performing a post-hoc power analysis. Logistic regression analysis was performed to evaluate potential predictive factors (in a multivariate model), including age (≥ 65 years), gender (male), and other variables that achieved a value of *P* < 0.10 in the univariate models. The cut-off value of FeNO

(≥ 50) was applied based on RCTs [11] and guidelines [20].

Results

Assessment of all patients

We excluded one pregnant woman who had received only one injection of mepolizumab from evaluation. The characteristics of the remaining 27 patients are shown in Table 1. Eight patients (30%) previously received omalizumab treatment. Among the 27 patients with severe asthma, 16 patients (59%) needed OSC maintenance therapy, and 16 patients (59%) suffered from ECRS as a comorbidity. The median number of mepolizumab injection was nine (range 2–17), and there was no significant difference between the ECRS and non-ECRS groups. The median duration of observation was 11 months (range 4–17), and there was a significant difference in observation duration between the two groups. Initial

Table 1 patients' characteristics at baseline

	all patients <i>n</i> = 27	ECRS (+) group <i>n</i> = 16	ECRS (–) group <i>n</i> = 11	<i>p</i> value between two groups
gender (male), <i>n</i> (%)	8 (30)	5 (31)	3 (27)	0.99 < ^a
age (year-old), (range)	56.3 (11.8), (35–79)	54.2 (11.3), (35–71)	59.3 (12.5), (36–79)	0.31 ^b
duration of asthma (year), (range)	19.6 (11.9), (4–43)	23.5 (10.6), (8–43)	45.4 (22.6), (4–39)	0.020 ^b
BMI(kg/m ²)	22.8 (4.7)	22.1 (4.5)	23.9 (4.9)	0.26 ^b
smoking (never / former / current), <i>n</i>	20 / 7 / 0	12 / 4 / 0	8 / 3 / 0	0.99 < ^a
atopic / non atopic, <i>n</i>	21 / 6	15 / 1	6 / 5	0.027 ^a
previous omalizumab, <i>n</i> (%)	8 (30)	6 (38)	2 (18)	0.40 ^a
initial treatments use				
—ICS/LABA, <i>n</i> (%)	27 (100)	16 (100)	11 (100)	NA
—ICS dose(μ g) ^c , mean (SD)	963 (222)	950 (137)	982 (316)	0.60
—LAMA, <i>n</i> (%)	13 (48)	5 (31)	8 (73)	0.054
—LTRA, <i>n</i> (%)	23 (85)	13 (81)	10 (91)	0.62
—xanthine derivative, <i>n</i> (%)	17 (63)	9 (56)	8 (73)	0.45
—maintenance therapy of OCS, <i>n</i> (%)	16 (59)	10 (63)	6 (55)	0.71 ^a
—daily dose of OCS (mg) ^d , (range)	8.4 (5.6), (0.5–20)	7.0 (4.8), (0.5–15)	10.8 (6.5), (5–20)	0.29 ^b
comorbidities, <i>n</i> (%)				
—ECRS	16 (59)	16 (100)	–	NA
—eosinophilic otitis media	9 (33)	9 (56)	0 (0)	NA
—EGPA	4 (15)	4 (25)	0 (0)	NA
number of injection, median (range)	9 (2–17)	9.5 (4–17)	7 (2–12)	0.17 ^b
duration of observation (month), median (range)	11 (4–17)	12.5 (6–17)	10 (4–12)	0.014 ^b

ECRS eosinophilic chronic rhinosinusitis, BMI body mass index, NA not available

EGPA eosinophilic granulomatous with polyangitis, ICS inhaled corticosteroid

LABA long-acting beta agonist, LAMA long-acting muscarinic antagonist

LTRA leukotriene receptor antagonist

Data presented as *n*(%) or mean (standard deviation), unless otherwise stated

^aFisher's exact test, ^bMann-Whitney *U* test

^cICS doses are provided as fluticasone propionate equivalents (μ g)

^dOral corticosteroid(OCS) doses are provided as prednisone equivalents (mg)

treatment at baseline is shown in Table 1. All patients received ICS/LABA, and 25 patients (93%) received high-dose ICS ($\geq 800 \mu\text{g}$, fluticasone propionate or equivalent). There was no significant difference regarding initial treatment between the ECRS and non-ECRS groups.

The variables at baseline and the last follow-up of mepolizumab are shown in Table 2. We performed a post-hoc power analysis regarding the following primary outcomes in all patients, which are shown in Table 2: ACT score, daily dose of OCS and exacerbation. The power was over 0.80 except for the daily dose of OCS in all patients (0.62) (data not shown). The peripheral blood eosinophil count and serum IgE level were significantly decreased from baseline to the last follow-up, and the ACT score was significantly improved (Fig. 1a). The number of asthma exacerbations requiring systemic corticosteroids and the maintenance dose of OCS were significantly reduced after introducing mepolizumab (Fig.

1b). The mean BMI was $22.9 \text{ (kg/m}^2\text{)}$ in all patients. Of the 19 patients for whom SEA- and SEB-IgE tests were performed, 11 had negative SEB-IgE test results (57.9%) (data not shown). Furthermore, of the 13 patients who responded to mepolizumab, nine had negative SEB-IgE results (69.2%) (data not shown).

Assessment of asthma patients with or without ECRS

We divided all 27 patients into two groups: and ECRS group (with ECRS) and a non-ECRS group (without ECRS). The characteristics of these patients are shown in Tables 1 and 2. Peripheral blood eosinophil count and FeNO level in the ECRS group were significantly higher than those in the non-ECRS group at baseline (Table 2 and Fig. 2a). Moreover, at baseline, FEV₁/FVC was significantly lower in the ECRS group than in the non-ECRS group (Table 2 and Fig. 2b). The power was over 0.80 for the three primary outcomes in the ECRS group, however it was 0.31 for exacerbation in the non-ECRS

Table 2 change from baseline to last follow-up in asthma patients with or without ECRS

	all patients, n = 27			ECRS (+) group, n = 16			ECRS (-) group, n = 11			between two groups p value [‡]
	baseline	last follow-up	p value ^a	at baseline	at last follow-up	p value ^a	at baseline	at last follow-up	p value ^a	
blood eosinophil count/mm ³	873 (1851)	71 (64)	< 0.0001	1219 (2353)	89 (74)	0.0004	338 (364)	44 (31)	0.005	0.0049
serum IgE (IU/ml)	485 (713)	305 (583)	0.016	440 (415)	261 (263)	0.0052	505 (992)	367 (873)	0.95	0.15 ^b
FeNO (ppb)	62 (57)	44 (33)	0.056	84 (62)	53 (37)	0.021	32 (24)	32 (24)	0.53	0.007 ^b
—Δ%FeNO (%), (range)	0.7 (69.2), (-68—180)			-19 (57), (-68—157)			30 (77), (-53—180)			0.023
ACT (pts)	15.2 (5.1)	19.5 (5.2)	0.0005	16.3 (5.5)	21.7 (4.8)	0.0023	13.6 (4.2)	16.4 (4.2)	0.066	0.16 ^b
—ΔACT (pts), (range)	5.3 (4.5), (-4—16)			5.4 (5.4), (-3—16)			2.7 (4.4), (-4—9)			0.24
%FVC (%)	91.1 (14.8)	99.2 (17.5)	0.031	94.7 (14.9)	100.3 (12.9)	0.14	85.1 (13.2)	97.7 (23.8)	0.09	0.16 ^b
%FEV ₁ (%)	83.3 (27.6)	83.9 (24.2)	0.62	77.4 (24.5)	79.0 (19.9)	0.36	92.1 (30.7)	91.5 (29.3)	0.44	0.35 ^b
FEV ₁ / FVC (%)	71.1 (13.3)	68.9 (12.9)	0.32	66.6 (12.8)	65.2 (12.5)	0.78	77.8 (11.6)	74.6 (11.8)	0.11	0.026 ^b
%PEF (%)	86.2 (28.2)	89.4 (23.6)	0.011	82.3 (30.8)	85.3 (24.0)	0.011	92.1 (24.1)	95.9 (22.7)	0.53	0.29 ^b
V ₅₀ /V ₂₅	3.5 (1.1)	3.5 (1.1)	0.61	3.5 (1.2)	3.1 (0.9)	0.81	3.5 (1.1)	4.1 (1.1)	0.26	0.78 ^b
daily dose of OCS (mg) [§] , (range)	8.4 (5.6), (0.5—20) (n = 16)	5.0 (5.8), (0—20) (n = 16)	0.0032	7.0 (4.8), (0.5—15) (n = 10)	2.0 (2.4), (0—6) (n = 10)	0.008	10.8 (6.5), (5—20) (n = 6)	10.0 (6.5), (2.5—20) (n = 6)	0.16	0.29 ^b
—ΔOCS (%), (range)	-48.6 (43.3), (-100—0)			-71.3 (37.0), (-100—0)			-10.7 (20.1), (-50—0)			0.006
exacerbation (/year), n	5.6 (4.9)	2.9 (3.7)	0.002	5.6 (5.6)	2.4 (3.3)	0.01	5.5 (3.9)	3.6 (4.2)	0.0498	0.77 ^b
—Δexacerbation (%), (range)	-44.7 (61.4), (-100—125)			- 48.4 (62.6),(- 100—125)			-39.9 (62.6), (- 100—100)			0.82

ECRS eosinophilic chronic rhinosinusitis, NA not available, ACT Asthma Control Test Δ: change from baseline to the last follow-up Data presented as mean (standard deviation), unless otherwise stated. Blood eosinophil counts at last follow-up were examined at 3 months (median, range 1–6). Serum IgE tests at last follow-up were examined at 6 months (median, range 2–6)

FeNO tests at last follow-up were examined at 9 months (median, range 1–12). Asthma Control Tests (ACT) score at last follow-up was examined at 6 months (median, range 2–12)

Pulmonary function tests at last follow-up were conducted at 4 months (median, range 2–10)

^aWilcoxon signed rank test [‡] Mann-Whitney U test [§] Oral corticosteroid (OCS) doses are provided as prednisone equivalents (mg). ^b at baseline

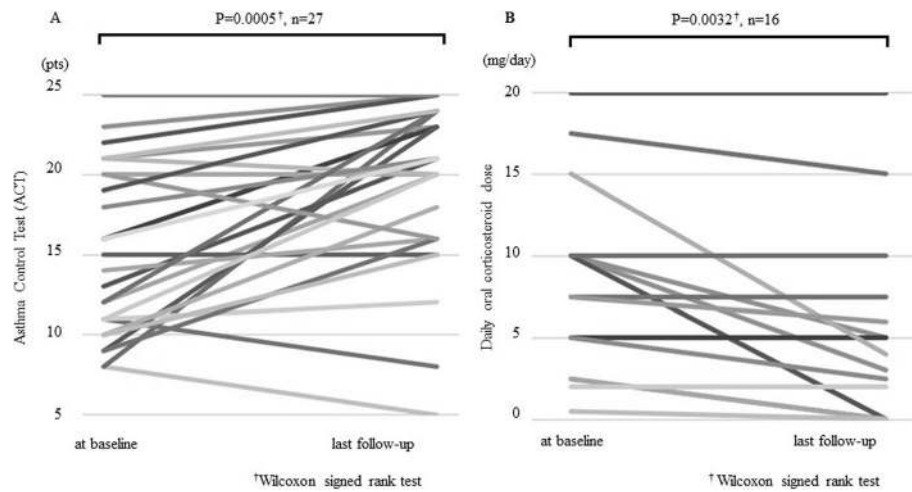


Fig. 1 Efficacy of introduction of mepolizumab. **a** There was a significant change from baseline to the last follow-up in the Asthma Control Test (ACT) score ($P = 0.0005$, Wilcoxon signed rank test, $n = 27$). **b** There was a significant change from baseline to the last follow-up in the daily dose of oral corticosteroids ($P = 0.0032$, Wilcoxon signed rank test, $n = 16$)

group (data not shown). Between the ECRS and non-ECRS group, there was a significant difference in the percentage reduction of the daily dose of OCS [$-71.3 \pm 37.0\%$ and $-10.7 \pm 20.1\%$, respectively, $P = 0.006$, Mann-Whitney U test] and in the change in FeNO level from baseline [$-19 \pm 57\%$ and $30 \pm 77\%$, respectively, $P =$

0.023 , Mann-Whitney U test] (Table 2, Fig. 3a and b). Of the 16 patients in the ECRS group, 10 received daily OCS at baseline, and five patients stopped OCS treatment after the introduction of mepolizumab. There were two patients that exhibited an increased number of exacerbations despite improvement in ACT score. After

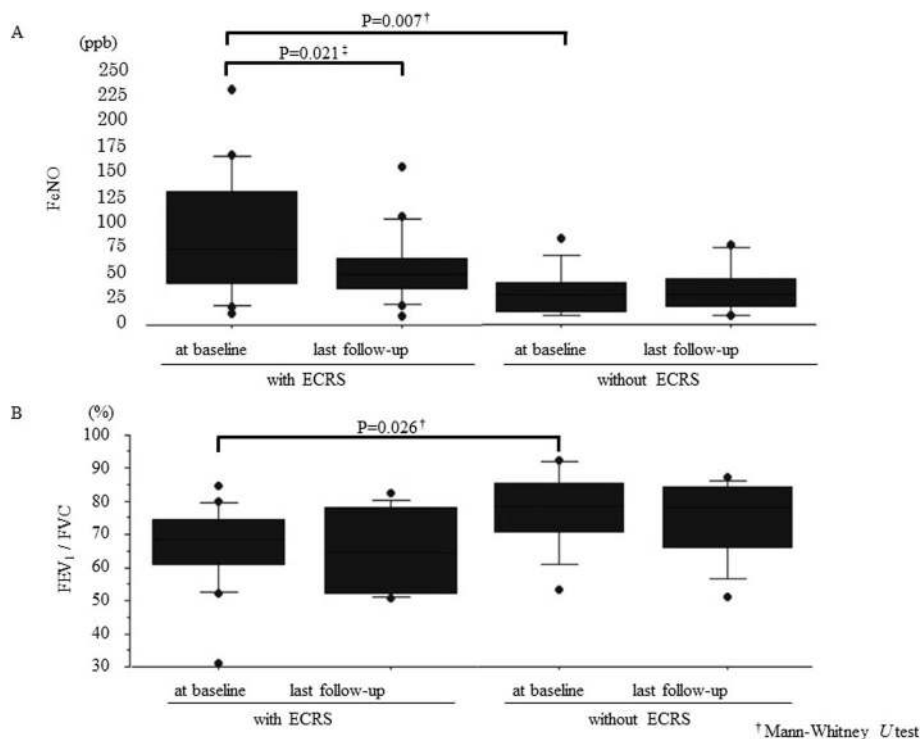
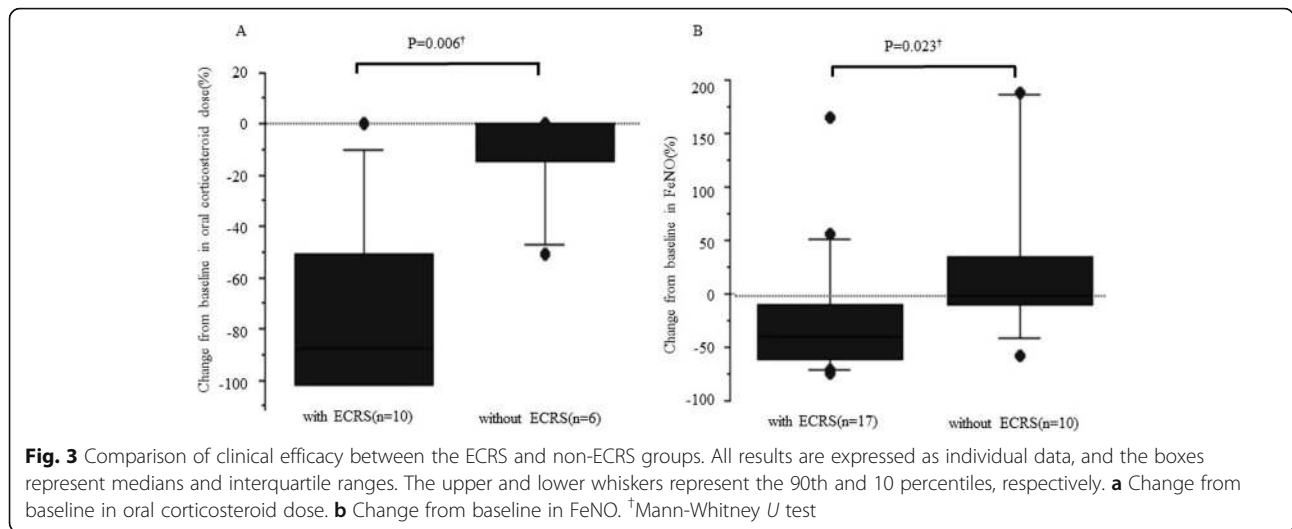


Fig. 2 FeNO level and FEV₁/FVC in the ECRS and non-ECRS groups. All results are expressed as individual data, and the boxes represent medians and interquartile ranges. The upper and lower whiskers represent the 90th and 10th percentiles, respectively. **a** FeNO level. **b** FEV₁/FVC. †Mann-Whitney U test, ‡Wilcoxon signed rank test



considering these factors, we concluded that 14 patients (88%) with ECRS and five patients (45%) without ECRS showed clinical improvement ($P = 0.033$, Fisher’s exact test).

Predictive factors

Among the 27 assessed patients, 16 patients (59%) showed a clinically relevant amelioration of ACT score (Table 1 and Fig. 1a). Furthermore, the maintenance dose of daily OCS (Table 2 and Fig. 1b) and the number of exacerbations were significantly reduced from 8.4 ± 5.6 to 5.0 ± 5.8 (mg/day) ($P = 0.0032$, Wilcoxon signed rank test) and from 5.6 ± 4.9 to 2.9 ± 3.7 (per year) ($P = 0.002$, Wilcoxon signed rank test), respectively, following the introduction of mepolizumab (Table 2). As mentioned above, we concluded that 19 patients (70%) showed clinical improvement. To identify predictive factors of mepolizumab response, we first analyzed each variable, such as patient background, comorbidities and

biomarkers, by univariate logistic regression analysis (Table 3). We identified ECRS as a predictive factor of a mepolizumab response [odds ratio (OR) = 8.4, 95% CI (1.3–56), $P = 0.028$]. The variables that achieved a value of $P < 0.10$ in the univariate models included ECRS and maintenance OCS therapy. We then evaluated predictive factors including age (≥ 65 years), gender (male), ECRS and maintenance OCS therapy by multivariate logistic regression analysis. Only ECRS was identified as a predictive factor by multivariate logistic regression analysis [OR = 22.5, 95% CI (1.5–336), $P = 0.024$].

Previous treatment with omalizumab

Recently, a cluster analysis of patients with chronic rhinosinusitis with nasal polyp (CRSwNP) reported that both IL-5-positive clustering and SE-IgE-positive clustering were important factors for eosinophilic inflammation when asthma was included in the analysis as a comorbidity [21]. Among 27 patients, eight patients with

Table 3 predictive factors by logistic regression analysis

variables	effective <i>n</i> = 19	non effective <i>n</i> = 8	odds ratio(95% CI) (monovariate)	<i>P</i> value	odds ratio(95% CI) (multivariate)	<i>P</i> value
gender (male), <i>n</i> (%)	7 (37)	1 (13)	4.1 (0.4–40)	0.23	5.6 (0.28–115)	0.26
age (≥ 65 year-old), <i>n</i> (%)	6 (32)	2 (25)	1.4 (0.2–9.0)	0.73	1.2 (0.05–26.6)	0.92
duration of asthma (≥ 20 years), <i>n</i> (%)	11 (58)	4 (50)	1.4 (0.3–7.2)	0.71	–	–
BMI(kg/m ²) ≥ 25 , <i>n</i> (%)	6 (32)	2 (25)	1.4 (0.2–9.0)	0.73	–	–
atopic type, <i>n</i> (%)	16 (84)	5 (63)	3.2 (0.5–21)	0.23	–	–
blood eosinophil count ≥ 150 (/mm ³) ^a , <i>n</i> (%)	18 (95)	7 (88)	2.6 (0.1–47)	0.54	–	–
FeNO ≥ 50 (ppb) ^a , <i>n</i> (%)	9 (47)	3 (38)	1.5 (0.3–8.1)	0.64	–	–
with maintenance therapy of OCS, <i>n</i> (%)	10 (59)	7 (75)	0.1 (0.01–1.3)	0.078	0.05 (0.001–1.5)	0.08
with ECRS as comorbidity, <i>n</i> (%)	14 (74)	2 (25)	8.4 (1.3–56)	0.028	22.5 (1.5–336)	0.024

CI confidence interval, BMI body mass index, OCS oral corticosteroid, ECRS eosinophilic chronic rhinosinusitis, ^a at baseline

inadequately controlled asthma despite omalizumab treatment were transitioned to mepolizumab treatment. Their ACT scores increased from 17.0 ± 4.6 to 21.1 ± 3.6 ($P = 0.063$, Wilcoxon signed rank test, data not shown), and all five patients (63%) who we assessed as effective cases had ECRS. All four patients whose serum SEB-IgE levels were negative (< 0.10 kU/L) tended to respond better to mepolizumab than to omalizumab ($P = 0.14$, Fisher's exact test, Table 4). A significant difference was observed between the SEB-IgE-positive and SEB-IgE-negative groups in the change from baseline of the daily dose of OCSs [$-4.8 \pm 8.2\%$ vs $-64 \pm 41\%$, respectively, $P = 0.046$, Mann-Whitney U test] (Table 4); with the power was greater than 0.80 according to the post-hoc power analysis (data not shown).

Discussion

We demonstrated that mepolizumab substantially improved the clinical variables of patients with asthma complicated with ECRS. In the present study, we identified ECRS as a predictive factor based on multivariate logistic regression analysis. BMI was among the predictive factors of mepolizumab reported in the previous cluster analysis reported [15]; in the present study, we adjusted BMI to the Japanese obese standard (≥ 25 kg/m²). A recent US database study reported that the risk factors for asthma attack include blood eosinophilia, rhinitis and nasal polyps [22]. Patients with ECRS often have eosinophilia, and reduction of blood eosinophil count is important for the stability of asthma symptoms. Although endoscopic sinus surgery improves asthma symptom scores in patients with ECRS [23], the recurrence rate of severe ECRS is approximately 70% or higher, and repeat surgery is often required [24]. The deterioration of ECRS is related to asthmatic symptoms, and patients are often dependent on corticosteroids. We hypothesize that the anti-IL-5 antibody inhibits eosinophilic inflammation more effectively than corticosteroids not only in the lower airway but also in the upper airway. Because ACT scores may deteriorate if mepolizumab treatment

reduces the daily dose of OCS, we evaluated multidisciplinary measures, such as ACT, the annual rate of asthma exacerbation and the OCS-sparing effect.

FeNO is useful for the diagnosis of asthma and for evaluating lower-airway inflammation. In the present study, FeNO levels in the ECRS group were higher than those in the non-ECRS group. This result is similar to that of a previous report [25]. Under eosinophilic inflammation, sinus tissue releases not only mediators and cytokines that directly induce inflammation of the lower airways but also chemotactic factors that recruit eosinophils from the bone marrow and circulation into the upper and lower airways [26, 27]. We observed that FeNO level significantly decreased from baseline in the ECRS group but not in the non-ECRS group. A potential reason for this result is that in the ECRS group, mepolizumab reduced not only the eosinophil counts in sinus tissue, but also eosinophilic airway inflammation. It is commonly reported that mepolizumab does not necessarily decrease FeNO level. Eosinophil-derived major basic protein (MBP) injures epithelial cells, which activates the innate immune system. Furthermore, group 2 innate lymphoid cells (ILC2s) and eosinophils release large amounts of IL-4, 5 and 13 [28, 29]. Hence, it is possible that anti-IL-5 therapy decreases peripheral blood and tissue eosinophils and indirectly inhibits the IL-4/IL-13 pathway, reducing serum IgE and FeNO levels.

Of the eight patients with previous omalizumab treatment in the present study, four patients (50%) had serum SEB-IgE levels lower than 0.10 kU/L; these patients responded to mepolizumab. On the other hand, three of the remaining four patients with positive serum SEB-IgE test results did not respond to mepolizumab. The prevalence of specific SE-IgE (≥ 0.10 kU/L) is approximately 40 to 75% in patients with asthma, and SE-IgE levels are associated with high IgE in eosinophilic asthma [30, 31]. Previous studies have suggested the potential involvement of SEB in the pathophysiology of not only eosinophilic asthma but also CRSwNP [32–35]. We

Table 4 association with SEB-IgE in the patients with previously omalizumab treatment ($n = 8$)

	SEB-IgE(+) $n = 4$	SEB-IgE(-) $n = 4$	p value
positive number with favours mepolizumab, n(%)	1 (25)	4 (100)	0.14 ^a
ECRS as comorbidity, n(%)	2 (50)	4 (100)	0.43 ^a
change from baseline in ACT score(pts)	4.5 (7.0)	3.8 (3.0)	0.77 ^b
change from baseline in OCS dose ^c (%)	-4.8 (8.2)	-64 (41)	0.046 ^b
change from baseline in exacerbation(%)	-8.8 (98)	-77 (42)	0.24 ^b

SEB-IgE *Staphylococcus aureus* enterotoxin B specific immunoglobulin E antibody, ECRS eosinophilic chronic rhinosinusitis, OMA omalizumab, ACT Asthma Control Test, OCS oral corticosteroid

Data presented as mean (standard deviation), unless otherwise stated

^aFisher's exact test

^bMann-Whitney U test

^cOCS doses are provided as prednisone equivalents

There are three patients received maintenance OCS treatment in each groups

occasionally observe patients with eosinophilic asthma who show slight or no response to mepolizumab in clinical practice. We suspect that such asthma patients are not IL-5-positive patients with CRSwNP/ECRS but are SE-IgE-positive patients. However, the number of patients with ECRS was small in the present study. Hence, additional data are needed to elucidate the usefulness of serum SE-IgE in predicting a response to mepolizumab in patients with severe asthma and ECRS.

There are several limitations to the present study. One limitation is that it is a small, single-center, retrospective study. To increase the reliability of the statistical analysis, a power analysis was warranted. Thus, we performed a post-hoc power analysis regarding the primary outcomes in every group. A higher number of patients and a prolonged follow-up period are needed for more accurate evaluations. A second is that the limited induction of biologic therapy due to high cost may have resulted in selection bias. Accordingly, data collection and analysis in clinical studies that are distinct from major RCTs are important. In the future, detailed assessments of comorbidities in patients with asthma will be important for clinical practice.

In conclusion, the results of the present study showed that the improvement in the asthmatic symptoms of patients with ECRS was stronger than that of patients without ECRS. This information may be useful for managing the introduction of biologic treatments. The SE-IgE test might be helpful for selecting mepolizumab or omalizumab therapy for patients with severe asthma.

Abbreviations

ACT: Asthma Control Test; CRSwNP: Chronic rhinosinusitis with nasal polyp; ECRS: Eosinophilic chronic rhinosinusitis; GINA: Global Initiative of Asthma; OCS: Oral corticosteroids; RCT: Randomized control trial; SE-IgE: *Staphylococcus aureus* specific IgE

Authors' contributions

TN, KN and JA designed the study and wrote the manuscript. TN, KN, HU, KKo, HY, MH, SM, TI, HH, JA and Kku contributed to data collection. TN and KN performed the statistical analyses and interpretation of the results. All authors were involved in preparation and review of the manuscript and approved the final version to be submitted.

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Availability of data and materials

The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

Ethics approval and consent to participate

For this retrospective study, written informed consent from the participants involved was unavailable and unnecessary. The Ethical Committee of Jikei University School of Medicine stated that patient consent was not necessary for this retrospective study. The informed consent was obtained in the form of the bulletin board, or opt-out on the website of our hospital. And this study approved by the Ethical Committee (28–259 (8502)). The director/administer of Jikei University hospital granted us the permissions to access the medical records.

Consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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References

- WHO. Global surveillance, prevention and control of chronic respiratory diseases: a comprehensive approach; 2007. p. 1–146. Available at http://whqlibdoc.who.int/publications/2007/9789241563468_eng.pdf?ua=1. Accessed 20 Feb 2018
- Fukutomi Y, Taniguchi M, Watanabe J, Nakamura H, Komase Y, Ohta K, et al. Time trend in the prevalence of adult asthma in Japan: findings from population-based surveys in Fujiyama City in 1985, 1999, and 2006. *Allergol Int.* 2011;60:443–8.
- Ichinose M, Sugiura H, Nagase H, Yamaguchi M, Inoue H, Sagara H, et al. Japanese guidelines for adult asthma 2017. *Allergol Int.* 2017;66:163–89.
- Hekking PPW, Wener RR, Amelink M, Zwinderman AH, Bouvy ML, Bel EH. The prevalence of severe refractory asthma. *J Allergy Clin Immunol.* 2015; 135:896–902.
- Global strategy for Asthma Management and Prevention. 2016; Available at www.ginasthma.org. Accessed 21 June 2016.
- Wenzel S. Severe asthma: from characteristics to phenotypes to endotypes. *Clin Exp Allergy.* 2012;42:650–8.
- Chung KF, Wenzel SE, Brozek JL, Bush A, Castro M, Sterk PJ, et al. International ERS/ATS guidelines on definition, evaluation and treatment of severe asthma. *Eur Respir J.* 2014;43:343–73.
- Hekking PPW, Amelink M, Wener RR, Bouvy ML, Bel EH. Comorbidities in difficult-to-control asthma. *J Allergy Clin Immunol Pract.* 2017;6:108–13.
- Price DB, Rigazio A, Campbell JD, Bleecker ER, Corrigan CJ, Thomas M, et al. Blood eosinophil count and prospective annual asthma disease burden: a UK cohort study. *Lancet Respir Med.* 2015;3:849–58.
- Rosenberg HF, Phipps S, Foster PS. Eosinophil trafficking in allergy and asthma. *J Allergy Clin Immunol.* 2007;119:1303–10.
- Pavord ID, Korn S, Howarth P, Bleecker ER, Buhl R, Keene ON, et al. Mepolizumab for severe eosinophilic asthma (DREAM): a multicentre, double-blind, placebo-controlled trial. *Lancet.* 2012;380:651–9.
- Ortega HG, Liu MC, Pavord ID, Brusselle GG, FitzGerald JM, Chetta A, et al. Mepolizumab treatment in patients with severe Eosinophilic asthma. *N Engl J Med.* 2014;371:1198–207.
- Bel EH, Wenzel SE, Thompson PJ, Prazma CM, Keene ON, Yancey SW, et al. Oral glucocorticoid-sparing effect of Mepolizumab in Eosinophilic asthma. *N Engl J Med.* 2014;371:1189–97.
- Yancey SW, Keene ON, Albers FC, Ortega H, Bates S, Bleecker ER, et al. Biomarkers for severe eosinophilic asthma. *J Allergy Clin Immunol.* 2017;140: 1509–18.
- Ortega H, Li H, Suruki R, Albers F, Gordon D, Yancey S. Cluster analysis and characterization of response to mepolizumab: a step closer to personalized medicine for patients with severe asthma. *Ann Am Thorac Soc.* 2014;1:1011–7.
- Shimoda T, Odajima H, Okamasa A, Kawase M, Komatsubara M, Mayer B, Yancey S, et al. Efficacy and safety of mepolizumab in Japanese patients with severe eosinophilic asthma. *Allergol Int.* 2017;66:445–51.
- American Thoracic Society Documents / European Respiratory Society. Recommendations for Standardized Procedures for the Online and Offline Measurement of Exhaled Lower Respiratory Nitric Oxide and Nasal Nitric Oxide, 2005. *Am J Respir Crit Care Med.* 2005;171:912–30.
- Takeno S, Hirakawa K, Ishino T. Pathological mechanisms and clinical features of Eosinophilic chronic Rhinosinusitis in the Japanese population. *Allergol Int.* 2010;59:247–56.
- Schatz M, Kosinski M, Yarlas AS, Hanlon J, Watson ME, Jhingran P, et al. The minimally important difference of the asthma control test. *J Allergy Clin Immunol.* 2009;124:719–23.
- Dweik R, Boggs P, Erzurum S, Irvin C. An Official ATS Clinical practice guideline: interpretation of exhaled nitric oxide levels (FENO) for. *Am J Respir Crit Care Med.* 2011;184:1–33.

21. Tomassen P, Vandeplass G, Van Zele T, Cardell LO, Arebro J, Olze H, et al. Inflammatory endotypes of chronic rhinosinusitis based on cluster analysis of biomarkers. *J Allergy Clin Immunol*. 2016;137:1449–56.
22. Blakey JD, Price DB, Pizzichini E, Popov TA, Dimitrov BD, Postma DS, et al. Identifying risk of future asthma attacks using UK medical record data: a respiratory effectiveness group initiative. *J Allergy Clin Immunol Pract*. 2017; 5:1015–24.
23. Schlosser RJ, Smith TL, Mace J, Soler ZM. Asthma quality of life and control after sinus surgery in patients with chronic rhinosinusitis. *Allergy*. 2017;72: 483–91.
24. Tokunaga T, Sakashita M, Haruna T, Asaka D, Takeno S, Ikeda H, et al. Novel scoring system and algorithm for classifying chronic rhinosinusitis: the JESREC study. *Allergy*. 2015;70:995–1003.
25. Guida G, Rolla G, Badiu I, Marsico P, Pizzimenti S, Bommarito L, et al. Determinants of exhaled nitric oxide in chronic rhinosinusitis. *Chest*. 2010; 137:658–64.
26. Denburg JA, Sehmi R, Saito H, Pil-Seob J, Inman MD, O'Byrne PM. Systemic aspects of allergic disease: bone marrow responses. *J Allergy Clin Immunol*. 2000;106:S242–6.
27. Ten Brinke A, Grootendorst DC, Schmidt JT, De Bruïne FT, Van Buchem MA, Sterk PJ, et al. Chronic sinusitis in severe asthma is related to sputum eosinophilia. *J Allergy Clin Immunol*. 2002;109:621–6.
28. Smith SG, Chen R, Kjarsgaard M, Huang C, Oliveria JP, O'Byrne PM, et al. Increased numbers of activated group 2 innate lymphoid cells in the airways of patients with severe asthma and persistent airway eosinophilia. *J Allergy Clin Immunol*. 2016;137:75–86.e8.
29. Doherty TA, Broide DH. Airway innate lymphoid cells in the induction and regulation of allergy. *Allergol Int*. 2019;68:9–16.
30. Bachert C, Van Steen K, Zhang N, Holtappels G, Cattaert T, Maus B, et al. Specific IgE against *Staphylococcus aureus* enterotoxins: an independent risk factor for asthma. *J Allergy Clin Immunol*. 2012;130:376–81.
31. Kowalski ML, Cieślak M, Pérez-Novo CA, Makowska JS, Bachert C. Clinical and immunological determinants of severe/refractory asthma (SRA): association with staphylococcal superantigen-specific IgE antibodies. *Allergy*. 2011;66: 32–8.
32. Patou J, Gevaert P, Van Zele T, Holtappels G, van Cauwenberge P, Bachert C. *Staphylococcus aureus* enterotoxin B, protein a, and lipoteichoic acid stimulations in nasal polyps. *J Allergy Clin Immunol*. 2008;121:110–5.
33. Hellings PW, Hens G, Meyts I, Bullens D, Vanoirbeek J, Gevaert P, et al. Aggravation of bronchial eosinophilia in mice by nasal and bronchial exposure to *Staphylococcus aureus* enterotoxin B. *Clin Exp Allergy*. 2006;36: 1063–71.
34. Chung KF. Staphylococcal enterotoxin-specific IgE: a biomarker for a distinct phenotype of severe asthma? *Clin Exp Allergy*. 2016;46:387–9.
35. Cui XY, Miao JL, Lu HQ, Qi QH, Chen XI, Xu J, et al. Serum levels of specific IgE to *Staphylococcus aureus* enterotoxins in patients with chronic rhinosinusitis. *Exp Ther Med*. 2015;9:1523–7.

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