

Postmarketing surveillance of safety and effectiveness of etanercept in Japanese patients with rheumatoid arthritis

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Abstract Our aim was to evaluate real-world safety and effectiveness in a 6-month postmarketing surveillance study covering all Japanese patients with rheumatoid arthritis (RA) who received etanercept during a 2-year period. Data for 13,894 patients (1334 sites) enrolled between March 2005 and April 2007 were collected. Adverse events (AEs) and serious adverse events (SAEs) were reported in 4336 (31.2%) and 857 (6.2%) patients, respectively. The most frequent AEs were injection site reactions ($n = 610$, 4.4%) and rash ($n = 339$, 2.4%), whereas pneumonia ($n = 116$, 0.8%) and interstitial lung disease ($n = 77$, 0.6%) were the most frequent SAEs. Significant improvement in the proportion of patients with a good European League Against Rheumatism (EULAR) response was observed from week 4 (17.6%) to week 24 (31.6%) ($p < 0.001$); 84.3% of patients had good or moderate EULAR responses at week 24. The percentage of

patients achieving remission increased significantly from week 4 (9.3%) to week 24 (18.9%) ($p < 0.001$). Patients with early moderate RA were less likely to experience SAEs and were more likely to achieve remission compared with patients with more severe disease. The safety and effectiveness of etanercept was demonstrated in Japanese patients in one of the largest observational trials conducted thus far in RA patients treated with biologics.

Keywords DAS28 · Etanercept · Postmarketing surveillance study · Rheumatoid arthritis

Introduction

Rheumatoid arthritis (RA), a chronic inflammatory disease affecting joints and extra-articular tissues, is associated

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with increased pain, deterioration of physical function, and decreased life expectancy [1, 2]. Achievement of disease remission is considered a realistic and crucial goal for patients with RA, as reflected by current treatment guidelines and recommendations of the European League Against Rheumatism (EULAR), the American College of Rheumatology (ACR), and others [3–5]. The modified disease activity score (DAS28) [6] is a validated 28-joint instrument that measures joint tenderness and/or swelling and can accurately evaluate the status of disease activity and the efficacy of treatment. Because of its reliability in monitoring and defining disease activity and remission, the DAS28 is commonly used in RA trials; it has also been endorsed by the EULAR working group recommendations for management of RA [5, 7].

Recent randomized clinical trials have demonstrated the efficacy and safety profile of RA treatments that function as antagonists of tumor necrosis factor alpha (TNF- α), either as monotherapy or combined with methotrexate [7–12]. In general, these treatment regimens were effective at reducing disease activity, achieving remission, and preventing joint destruction in patients with RA. However, most of the data describing efficacy and safety for RA treatments come from late-phase clinical trials or national databases. Inclusion criteria for the studies are not always reflective of treatment in a real-world environment. Large well-designed phase 3 trials generally provide useful patient outcomes data, but these studies may not adequately define the true safety and effectiveness of a drug outside the clinical research setting.

As one of the conditions of approval of etanercept in Japan, the Pharmaceutical and Medical Device Agency (PMDA) requested that Wyeth (now integrated into Pfizer as of October 2009) conduct surveillance to confirm the safety and effectiveness of etanercept in Japanese patients in the clinical setting after the drug was marketed. To comply with this request, we performed a postmarketing surveillance (PMS) study that registered all Japanese RA patients treated with etanercept at the participating study sites for the survey period. Etanercept is a fully human soluble TNF- α receptor fusion protein with demonstrated efficacy against RA in patients not adequately responding to disease-modifying antirheumatic drugs (DMARDs) such as methotrexate [13, 14]. Etanercept has shown superior efficacy to methotrexate in patients with RA [15], and the combination of etanercept plus methotrexate was superior to either monotherapy alone [16]. The goal of this PMS study was to evaluate the real-world safety and effectiveness of etanercept for a large Japanese patient population receiving etanercept for the treatment of RA. A previous interim analysis comprising data for 7091 of the registered patients reported that etanercept was effective with no new safety signals [17]. This report covers nearly 14,000

Japanese patients with RA registered to the PMS study from 2005 to 2007.

Patients, materials, and methods

Patients

Between March 2005 and April 2007, all Japanese patients with RA from 1334 sites were enrolled in a 6-month PMS study of etanercept use (NCT00503503). Patient eligibility for treatment with etanercept was based on the Japan College of Rheumatology treatment guidelines [18]. Briefly, etanercept was indicated for patients with RA who were previously treated with DMARDs (e.g., methotrexate, salazosulfapyridine, or bucillamine) for >3 months, and had ≥ 6 tender joints, ≥ 6 swollen joints, and erythrocyte sedimentation rate (ESR) ≥ 28 mm/h or C-reactive protein (CRP) levels ≥ 2.0 mg/dL [17]. Patients had a low risk for opportunistic infections, defined as white blood count $\geq 4000/\text{mm}^3$, peripheral blood lymphocyte count $\geq 1000/\text{mm}^3$, and negative serum β -D-glucan. Mandatory chest radiographs and tuberculin tests were conducted before initiation of treatment. Information on age, sex, comorbidity, Steinbrocker radiographic stage [19], Steinbrocker functional class [19], duration of RA, smoking history, previous and concomitant use of glucocorticoids, and concomitant use of DMARDs (including methotrexate) was collected. Also, ESR and CRP levels were assessed. Etanercept 10 or 25 mg was administered subcutaneously twice weekly, with the dose determined by the prescribing physician. After an initial 1-month training period, patients were allowed to self-inject etanercept.

The PMS protocol was reviewed and approved by the Japanese Ministry of Health, Labor, and Welfare. Registration was conducted centrally at the time of treatment initiation. Data collection was performed through an electronic data capture system, the Internet, or hardcopy case report forms, and medical representative staff members from Wyeth and Takeda Pharmaceutical Company visited sites periodically to collect additional data as required.

Assessments

All patients were assessed for safety every 2 weeks. Safety evaluations included all events occurring ≤ 24 weeks after the first etanercept dose and ≤ 30 days after the last dose. All adverse events (AEs) were reported. Safety data were coded with preferred terms from the Medical Dictionary for Regulatory Activities [20]. All AEs, serious AEs (SAEs), adverse drug reactions (ADRs; noxious and unintended responses deemed to be related to the treatment), and serious ADRs, including infection, were defined

based on the International Conference on Harmonisation tripartite harmonised guideline [21]. Particular attention was paid to the occurrence of infections, especially tuberculosis, pneumonia, *Pneumocystis jiroveci* pneumonia, cytomegalovirus infection, and sepsis and to specific important ADRs that included malignant neoplasm, demyelination, congestive heart failure, injection site reaction, and lupus. Safety information was independently evaluated by the Japan College of Rheumatology PMS committee.

Treatment effectiveness was measured monthly using the EULAR response criteria [22] and the DAS28 [6]. The DAS28 is divided into 4 categories: remission (DAS28 <2.6), low disease activity (DAS28 \geq 2.6 and \leq 3.2), moderate disease activity (DAS28 >3.2 and \leq 5.1), and high disease activity (DAS28 >5.1). A good response was defined as a DAS28 improvement from baseline of >1.2 and a DAS28 of \leq 3.2 during follow-up. Patients with score improvements of \leq 0.6 or those with improvements between 0.6 and 1.2 plus a DAS28 of >5.1 during follow-up were defined as nonresponders. Moderate responders were those with DAS28 improvements from baseline between 0.6 and 1.2 plus a DAS28 of \leq 5.1 during follow-up. Treatment was considered to be effective in patients with moderate or good DAS28 responses. General health status was measured using a patients' visual analog scale (100 being the worst they can imagine, 0 being the best they can imagine), and duration of morning stiffness was also assessed.

Statistical analysis

Missing data were processed using the last-observation-carried-forward method, except for baseline values, which were not carried forward. The χ^2 or *t*-test was used to compare differences in baseline values between men and women. The *t*-test was used to compare DAS28 differences between baseline and weeks 4 and 24; χ^2 tests were used to compare differences in EULAR response rates, and Cochran-Armitage tests were used to examine evidence of trends in response rates. Cox proportional hazard models were used to estimate the influence of multiple variables on the occurrence of serious infections (AEs). These variables included the confounders assessed at baseline: age, sex, concomitant methotrexate use, concomitant glucocorticoid use, history of infectious disease, history of tuberculosis, presence of any comorbidities, Steinbrocker functional class, and duration of RA. A total of 1274 patients with missing data [duration of RA ($n = 1267$) and other ($n = 7$)] were excluded from the Cox proportional hazard models. We also used multiple logistic regression models to estimate the effect of variables on the likelihood of achieving remission. Patients who had missing DAS28 data

at either baseline or at 24 weeks, a DAS28 value of <2.6 at baseline, or missing data for other adjustment factors were excluded from these models. Furthermore, we used Cox proportional hazard models and multiple logistic regression models to examine the combined effects of multiple risk factors on the occurrence of serious infections and the likelihood of achieving remission, respectively. The hazard ratio or odds ratio and the 95% confidence interval (CI) for each of the combined factors after adjustment for major confounders and the combined effect relations with numbers of factors were further explored by trend tests. All statistical analyses were performed using SAS[®] software version 8.2 (SAS Institute, Cary, NC, USA). Two-sided *P* values of < 0.05 were considered statistically significant.

Results

Patients

A total of 13,894 patients treated with etanercept completed the 24-week study (Table 1). The majority of patients were women ($n = 11,314$; 81.4%). Mean \pm SD patient age was 58.1 ± 13.1 years; most patients (78.3%) were aged >50 years (more than one half were aged >60 years), and the mean \pm SD patient weight was 53.2 ± 10.1 kg. About 40% of patients had a disease duration of >10 years. Concomitant use of DMARDs/biologics was 74.0% ($n = 10,276$) and that of methotrexate was 55.9% ($n = 7768$). The most commonly used etanercept dose regimen (76.1%; $n = 10,578$) was 50 mg per week (i.e., 25 mg twice weekly). Previous use of glucocorticoid was 83.4% ($n = 11587$) and previous use of infliximab was 13.5% ($n = 1878$). Additionally, 57.1% of patients had comorbidities, including 877 patients (6.3%) with a medical history of tuberculosis. Significant differences were observed between men and women in most demographic characteristics, including age, weight, disease duration, Steinbrocker stage and class, history of concomitant medical conditions, comorbidities, concomitant use of methotrexate, and prior glucocorticoid use (Table 1). A total of 11,615 (83.6%) patients completed 24 weeks of therapy, with 2309 patients (16.6%) discontinuing during the 24-week period. Reasons for stopping treatment were AEs (7.6%, $n = 1049$), lack of treatment effectiveness (2.6%, $n = 368$), refusal of treatment for economic reasons (1.5%, $n = 212$), moved to another hospital (1.6%, $n = 222$), and other (3.3%, $n = 458$).

Safety

As shown in Table 2, AEs and SAEs were reported in 4336 (31.2%) and 857 (6.2%) patients, respectively. The most

Table 1 Patient demographics

Characteristic	Male patients (<i>n</i> = 2580)	Female patients (<i>n</i> = 11,314)	<i>p</i> value ^a
Mean (SD) age, years	57.5 (13.2)	60.7 (12.4)	<0.001 ^b
Age range [<i>n</i> (%), years]			
<20	11 (0.4)	65 (0.6)	
20–29	52 (2.0)	342 (3.0)	
30–39	124 (4.8)	843 (7.5)	
40–49	198 (7.7)	1382 (12.2)	
50–59	691 (26.8)	3243 (28.7)	
60–69	837 (32.4)	3358 (29.7)	
≥70	667 (25.9)	2081 (18.4)	<0.001
Mean (SD) weight, kg	61.7 (10.4)	51.3 (9.0)	<0.001 ^b
Disease duration [<i>n</i> (%), years]			
<2	436 (16.9)	1168 (10.3)	
2–<5	588 (22.8)	1844 (16.3)	
5–<10	589 (22.8)	2386 (21.1)	
10–<15	328 (12.7)	1925 (17.0)	
15–<20	194 (7.5)	1166 (10.3)	
≥20	208 (8.1)	1795 (15.9)	
Unknown (years)	237 (9.2)	1030 (9.1)	<0.001
Steinbrocker stage [<i>n</i> (%)]			
I	241 (9.4)	624 (5.5)	
II	801 (31.1)	2458 (21.7)	
III	928 (36.0)	3998 (35.4)	
IV	606 (23.5)	4225 (37.4)	<0.001
Steinbrocker class [<i>n</i> (%)]			
1	277 (10.8)	946 (8.4)	
2	1536 (59.6)	6526 (57.7)	
3	694 (26.9)	3482 (30.8)	
4	69 (2.7)	351 (3.1)	<0.001
Positive history of concomitant medical conditions [<i>n</i> (%)]			
Tuberculosis	228 (8.8)	649 (5.7)	<0.001
Interstitial pneumonitis	341 (13.2)	693 (6.1)	<0.001
Follicular bronchitis	5 (0.2)	32 (0.3)	
COPD	43 (1.7)	41 (0.4)	<0.001
Total comorbidities [<i>n</i> (%)]	1570 (60.9)	6359 (56.2)	<0.001
Hepatic	155 (6.0)	475 (4.2)	<0.001
Renal	152 (5.9)	395 (3.5)	<0.001
Hematologic	100 (3.9)	698 (6.2)	<0.001
Cardiac	217 (8.4)	569 (5.0)	<0.001
Infectious (nonserious)	56 (2.2)	168 (1.5)	0.013
Diabetes mellitus	340 (13.2)	758 (6.7)	<0.001
Weekly etanercept dose regimen [<i>n</i> (%), mg]			
50	2018 (78.2)	8560 (75.7)	
25	264 (10.2)	1298 (11.5)	
20	2 (0.1)	27 (0.2)	
20–50 ^c	296 (11.5)	1429 (12.6)	0.023 ^d

Table 1 continued

Characteristic	Male patients (<i>n</i> = 2580)	Female patients (<i>n</i> = 11,314)	<i>p</i> value ^a
Concomitant DMARD/ biologic use [<i>n</i> (%)]	1924 (74.6)	8352 (73.8)	
Prior glucocorticoid use [<i>n</i> (%)] ^e	2209 (88.4)	9378 (85.5)	<0.001
Concomitant MTX use [<i>n</i> (%)]	1371 (53.1)	6397 (56.5)	0.002
MTX dose [<i>n</i> (%), mg/week]			
0–<4	39 (2.8)	271 (4.2)	
4–<6	239 (17.4)	1583 (24.7)	
6–<8	448 (32.7)	2157 (33.7)	
8–<10	497 (36.3)	1898 (29.7)	
≥10	148 (10.8)	488 (7.6)	<0.001
Prior infliximab use [<i>n</i> (%)]	366 (14.2)	1512 (13.4)	

COPD chronic obstructive pulmonary disease, DMARD disease-modifying antirheumatic drug, MTX methotrexate

^a The *p* values indicated are for comparisons between male and female patient data, using χ^2 tests unless otherwise indicated

^b Comparisons were made using the *t*-test

^c Patients received variable dosing during the observation period

^d Comparisons were made using Fisher exact test

^e Unknown patients were omitted from the percentage calculation

frequently observed AEs were injection site reaction (*n* = 610, 4.4%), rash (*n* = 339, 2.4%), and abnormal hepatic function (*n* = 328, 2.4%). Pneumonia was the most frequently reported SAE (*n* = 116, 0.8%), followed by interstitial lung disease (*n* = 77, 0.6%), and pyrexia (*n* = 40, 0.3%). In total, 26.7 and 4.6% of patients reported ADRs and serious ADRs, respectively. Table 2 also lists selected important ADRs; various forms of pneumonia were the most common ADR (*n* = 174, 1.3%). Tuberculosis was reported as an ADR in 12 patients (0.1%), and interstitial lung disease in 81 patients (0.6%). No cases of demyelinating disease were observed. Lupus-like syndrome and congestive heart failure (including all heart failure) were recorded as ADRs in 5 and 7 patients, respectively. Thirty cases of malignancy were reported (0.2%), and there were 76 deaths (0.6%) during the 24-week study.

Risk factors for the development of serious infections are shown in Table 3. Compared with the respective referent group, female sex and concomitant methotrexate use significantly lowered the risk of infection, whereas older age (≥65 years), history of infectious disease, presence of any comorbidities, Steinbrocker functional class 4, and concomitant glucocorticoid use significantly increased serious infection risk. Prior history of tuberculosis slightly, but not significantly, raised the infection risk compared

Table 2 Incidences of the most commonly reported adverse events, serious adverse events, important adverse drug reactions, and death

Event	n (%)
Patients with at least 1 AE, ^a total	4336 (31.2)
Injection site reaction	610 (4.4)
Rash	339 (2.4)
Abnormal hepatic function	328 (2.4)
Nasopharyngitis	288 (2.1)
Pyrexia	261 (1.9)
Upper respiratory tract infection	224 (1.6)
Pruritus	202 (1.5)
Total pneumonia ^b	189 (1.0)
Herpes zoster	115 (0.8)
Erythema	114 (0.8)
Patients with at least 1 SAE, ^a total	857 (6.2)
Total pneumonia ^b	116 (0.8)
Interstitial lung disease	77 (0.6)
Pyrexia	40 (0.3)
Sepsis	27 (0.2)
Herpes zoster	23 (0.2)
<i>Pneumocystis jiroveci</i> pneumonia	24 (0.2)
Urinary tract infection	16 (0.1)
Abnormal hepatic function	15 (0.1)
Bacterial arthritis	13 (0.1)
Bronchitis	13 (0.1)
Patients with at least 1 important ADR ^a	968 (7.0) ^c
Total pneumonia ^b	174 (1.3)
Interstitial lung disease	81 (0.6)
<i>Pneumocystis jiroveci</i> pneumonia	25 (0.2)
Malignancy	30 (0.2)
Tuberculosis ^d	12 (0.1)
Pulmonary	10 (0.1)
Extrapulmonary	3 (<0.1)
Congestive heart failure	7 (0.1)
Lupus-like syndrome	5 (<0.1)
Demyelinating disease	0 (0)
Deaths	76 (0.6)

ADR adverse drug reaction, AE adverse event, SAE serious adverse event

^a Patients who had at least 1 AE, SAE, or specifically important ADR, respectively. The 10 most frequently reported AEs, SAEs, and ADRs are listed

^b Total pneumonia = pneumonia + bacterial pneumonia + bronchopneumonia + *Chlamydia* pneumonia + staphylococcal pneumonia + *Candida pneumonia* + fungal pneumonia; 1 patient developed both pneumonia and bronchopneumonia

^c 609 patients who had injection site reactions were included

^d 1 patient had both pulmonary and extrapulmonary tuberculosis

with those without a history of tuberculosis (model 1). Model 2 showed the combined effects of multiple risk factors on the occurrence of serious infections adjusted for

the remaining variables included in model 1. Compared with patients without combined risk factors, the hazard ratio of serious infection was 9.91 (95% CI 5.48–17.94; $p < 0.001$) for patients who had 3 risk factors, 4.24 (95% CI 2.83–6.34; $p < 0.001$) for patients who had 2 risk factors, and 1.96 (95% CI 1.30–2.96; $p = 0.001$) for patients who had 1 risk factor. Highly significant linear association ($p < 0.001$) was evident for increasing number of combined risk factors.

Effectiveness

The effectiveness of etanercept treatment as determined by various assessment methods is shown in Table 4. The mean DAS28 decreased significantly (both $p < 0.01$), from 5.9 at baseline to 4.3 at week 4 (27% improvement) and 3.8 at week 24 (36% improvement). The mean duration of morning stiffness also decreased significantly (both $p < 0.01$), from 111.3 min at baseline to 44.2 min at week 4 (60% improvement) and 31.1 min at week 24 (72% improvement). The mean numbers of tender joints (9.2 at baseline, 4.2 at week 4, and 3.0 at week 24) and swollen joints (8.6, 4.1, and 2.8, respectively), and the mean general health status (60.1, 36.6, and 30.7 mm, respectively, by patients' visual analog scale) all showed significant (all $p < 0.01$) improvements from baseline at weeks 4 and 24. The mean ESR improved significantly, from 58.7 mm/h (baseline) to 38.1 mm/h (week 4) and 34.8 mm/h (week 24), representing improvements of 35 and 41%, respectively (both $p < 0.01$). The mean levels of CRP decreased significantly (both $p < 0.01$), from 3.6 mg/dL at baseline to 1.4 mg/dL at week 4 (61% improvement) and 1.2 mg/dL at week 24 (67% improvement).

The EULAR response and DAS28 remission rate were also evaluated. At week 4, 779 patients (17.6%) had achieved a good EULAR response; a trend of significant improvement occurred throughout the observation period, with 2336 patients (31.6%) having a good response by week 24 ($p < 0.001$ for trend; Fig. 1a). At week 4, 77.0% of patients had achieved a good or moderate response; response rates increased from week 4 to 24, with 84.3% of patients achieving a good or moderate response by week 24. A total of 1161 (15.7%) patients had no response to etanercept at week 24 (Fig. 1a). Remission was reported in 1395 patients (18.9%) at week 24, with significant improvement noted during the observation period ($p < 0.001$ for trend; Fig. 1b). Low, moderate, and high disease activity were reported in 1179 (15.9%), 3730 (50.4%), and 1091 (14.8%) patients, respectively, at week 24 (Fig. 1b).

Factors that affected the odds of achieving RA remission are shown in Table 5. Compared with the referent group in model 1, male sex, younger age (<65 years), concomitant methotrexate use, lower baseline DAS28 (moderate

Table 3 Hazard ratios for serious infection (adverse events)

Variable	HR	95% CI	<i>p</i> value
Model 1^a			
Sex (women vs. men)	0.63	0.50–0.81	<0.001
Age (≥ 65 vs. <65 years of age)	1.66	1.33–2.07	<0.001
History of infectious disease (yes vs. no)	2.26	1.38–3.70	0.001
History of tuberculosis (yes vs. no)	1.24	0.85–1.80	0.274
Presence of any comorbidities (yes vs. no)	2.72	2.02–3.66	<0.001
Steinbrocker functional class (4 vs. 1 + 2 + 3)	2.54	1.73–3.71	<0.001
Duration of RA (years)			
≥ 5 and <10 vs. <5	1.20	0.89–1.61	0.237
≥ 10 and <15 vs. <5	1.01	0.72–1.41	0.971
≥ 15 vs. <5	1.04	0.78–1.40	0.774
Concomitant use of MTX (yes vs. no)	0.59	0.47–0.74	<0.001
Concomitant use of glucocorticoids (yes vs. no)	2.03	1.46–2.84	<0.001
Model 2^b			
Presence of combined risk factors ^c			
1 vs. 0	1.96	1.30–2.96	0.001
2 vs. 0	4.24	2.83–6.34	<0.001
3 vs. 0	9.91	5.48–17.94	<0.001

HR hazard ratio, CI confidence interval, MTX methotrexate

^a Cox proportional hazard model (for all cases, $n = 12,620$ and for serious infection cases, $n = 330$)

^b Results were adjusted for sex, age, history of infectious disease, history of tuberculosis, duration of rheumatoid arthritis (RA), and concomitant use of glucocorticoid

^c Combined factors: Steinbrocker functional class = 4, concomitant use of MTX = no, any comorbidities = yes; $p < 0.001$ for linear trend using the Wald test

Table 4 Measures of effectiveness

Measure	Baseline, mean (SD)	Patients evaluated at baseline (<i>n</i>)	Week 4, mean (SD) ^a	Patients evaluated at week 4 (<i>n</i>)	Week 24, mean (SD) ^a	Patients evaluated at week 24 (<i>n</i>)
DAS28	5.9 (1.2)	8902	4.3 (1.3)	4754	3.8 (1.3)	8137
Duration of morning stiffness (min)	111.3 (181.3)	5858	44.2 (114.0)	3201	31.1 (90.1)	5058
Tender joints (<i>n</i>)	9.2 (7.0)	12727	4.2 (4.8)	7873	3.0 (4.1)	12321
Swollen joints (<i>n</i>)	8.6 (6.2)	12727	4.1 (4.3)	7871	2.8 (3.6)	12319
General health status (patient visual analog scale) (mm)	60.1 (22.7)	11535	36.6 (22.5)	6702	30.7 (22.1)	10616
ESR (mm/h)	58.7 (33.0)	9719	38.1 (27.7)	5732	34.8 (27.5)	9484
CRP (mg/dL)	3.6 (3.3)	12693	1.4 (2.1)	8298	1.2 (2.0)	12770

CRP C-reactive protein, DAS28 modified disease activity score including a 28-joint count, ESR erythrocyte sedimentation rate

^a All comparisons differed significantly ($p < 0.01$ vs. baseline), by *t*-test

disease), and better Steinbrocker functional class (1–3) significantly improved the odds of achieving remission. Shorter duration of RA also significantly improved the chances of achieving remission compared with those with ≥ 15 years' duration (odds ratio [OR] 1.62; 95% CI 1.36–1.92). Model 2 showed the combined effects of multiple risk factors on the likelihood of achieving remission. Compared with the groups with zero risk factors, the

odds ratio of achieving remission was 6.30 (95% CI 4.83–8.21; $p < 0.001$) for patients who had 4 risk factors, 3.20 (95% CI 2.64–3.88; $P < 0.001$) for patients who had 3 risk factors, and 1.87 (95% CI 1.55–2.26; $p < 0.001$) for patients who had 2 risk factors. Highly significant linear association ($p < 0.001$) was evident for increasing number of combined risk factors. When we added 79 patients with lower disease activity (baseline DAS28, ≥ 2.6 and ≤ 3.2)

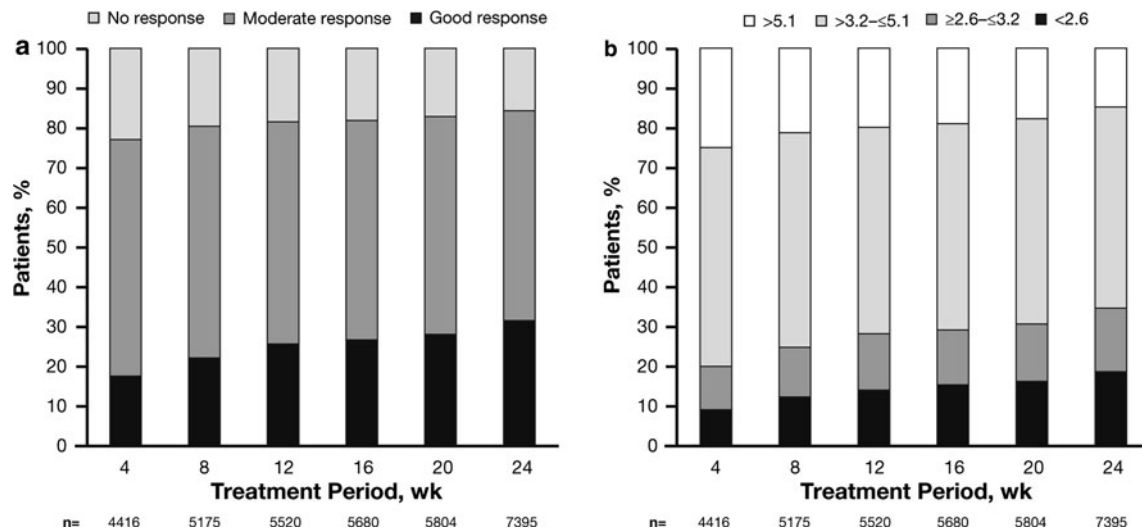


Fig. 1 Time course of **a** EULAR response and **b** DAS28 from week 4 to week 24. *DAS28* modified disease activity score including a 28-joint count, *EULAR* European League Against Rheumatism.

Cochran-Armitage tests were used to examine evidence of trends in response rates; trends for good response rate and remission rate were statistically significant ($p < 0.001$). *wk* Week

Table 5 Odds ratios for achieving remission

Variable	OR	95% CI	<i>p</i> value
Model 1			
Sex (men vs. women)	1.30	1.10–1.53	0.002
Age (<65 vs. ≥65 years of age)	1.43	1.23–1.65	<0.001
Steinbrocker functional class (1–3 vs. 4)	2.41	1.30–4.49	0.006
Duration of RA (years)			
<5 vs. ≥15	1.62	1.36–1.92	<0.001
5–10 vs. ≥15	1.17	0.97–1.42	0.098
10–15 vs. ≥15	1.04	0.84–1.28	0.733
DAS28 at baseline (moderate vs. high)	2.95	2.59–3.37	<0.001
Concomitant use of MTX			
<8 mg/week vs. none	1.30	1.13–1.49	<0.001
≥8 mg/week vs. none	1.74	1.32–2.28	<0.001
Previous treatment with infliximab (yes vs. no)	0.65	0.53–0.81	<0.001
Model 2^a			
Presence of combined risk factors ^b			
4 vs. 0–1	6.30	4.83–8.21	<0.001
3 vs. 0–1	3.20	2.64–3.88	<0.001
2 vs. 0–1	1.87	1.55–2.26	<0.001

Multiple logistic regression models [for all cases, $n = 6763$ (79 patients with low disease activity were excluded) and for remission cases, $n = 1234$]

DAS28 modified disease activity score including a 28-joint count, *MTX* methotrexate, *OR* odds ratio, *RA* rheumatoid arthritis

^a Results were adjusted for sex, age, and previous treatment with infliximab

^b Combined factors: Steinbrocker functional class = 1–3; *MTX* = yes; baseline *DAS28* = >3.2 and ≤5.1 (moderate disease); duration of *RA* <5 years; $p < 0.001$ for linear trend using the Wald test

into the lower baseline *DAS28* group to examine the odds ratio of achieving remission, a similar pattern was seen (data not shown).

Discussion

The current trial was one of the largest surveillance studies of biologic use in the rheumatology area, with nearly 14,000 patients registered. Mandatory registration for all patients treated with etanercept occurred at participating sites in Japan during the 2-year study period. This PMS study provided a unique opportunity to capture real-world safety and effectiveness data for a large patient population in an Asian country. Although there are published accounts of other PMS studies with *RA* biologic treatments [23–25], the large number of patients registered in the current study allowed for the safety and effectiveness of etanercept to be documented in a real-world clinical setting in Japan. The advantage of conducting a study like this is a high precision of AE incidence, remission rate, and other important data.

In the current PMS study, AEs and SAEs were reported in 31.2 and 6.2% of patients, respectively. A good or moderate response (i.e., effective treatment) occurred in a very high percentage of patients from week 4 to 24. These data are consistent with those reported in the interim analysis from this study [17]. No new signaling risk factors for serious infection were observed in this PMS study.

Real-world efficacy data were also reported from the Rheumatoid Arthritis DMARD Intervention and Utilization Study (RADIUS), a 5-year, multicenter, observational

registry of patients treated with etanercept and other RA therapies in the United States ($N = 10,061$ in 2 cohorts) [26, 27]. Although safety data have not yet been reported, patients treated with etanercept, with ($p < 0.01$) or without ($p < 0.05$) methotrexate, were significantly more likely to have a modified ACR 20% response at 12 months compared with those receiving only methotrexate. In another PMS study involving TNF- α antagonists, Feltelius et al. [23] collected safety and effectiveness data from 1999 to 2003 in the cohort of all etanercept-treated Swedish patients with RA ($N = 1073$). The 24-month incidence rates of ADRs and SAEs of 27 and 7%, respectively, in that study are both similar to those observed in the current PMS study. A good or moderate EULAR response rate was observed in 86% of 517 patients, which is similar to the rate reported in our analysis (84.3%) at week 24.

Because the registered patients in the current PMS study had varied backgrounds, multivariate analyses were conducted to ascertain risk factors related to safety and effectiveness in a real-world setting. The current PMS study identified several factors that improved the safety and effectiveness outcomes of patients treated with etanercept. Patients were more likely to achieve DAS28 remission if they had moderate disease activity, better Steinbrocker functional class, shorter disease duration, and received concomitant methotrexate treatment. Importantly, patients who had a combination of these factors showed a higher probability of achieving remission.

Cox proportional hazard model results also demonstrated several risk factors for serious infection. A combination of Steinbrocker functional class 4, no use of concomitant methotrexate, and the presence of any comorbidities significantly increased the risk for developing serious infection. Thus, the combined use of etanercept and methotrexate in patients with early moderate RA with less comorbidity and better physical function appears to provide patient benefit for the achievement of remission and lowering of SAE occurrences.

Interpretation of these data is somewhat limited by the fact that no control arm was included in this large PMS study. This makes it difficult to distinguish outcomes relating to etanercept treatment from those caused by other factors (e.g., patient expectations, natural history of the disease, or concomitant treatments). The study length (6 months) allowed for the collection of important safety and effectiveness data, but longer-term studies would also be useful. Additionally, evaluations of effectiveness did not include radiographic analysis to confirm the effectiveness of treatment.

This PMS study collected safety and effectiveness data for every Japanese patient with RA receiving etanercept at the participating study sites for a 2-year period. With nearly 14,000 patients registered, this represents one of the largest observational surveillance studies conducted to date

in RA patients treated with biologics. The safety and effectiveness data reported here support data from previous clinical trials with etanercept and are also consistent with the data from the interim analysis of this study. Additional subgroup analyses from this study may enable the identification of important factors affecting the safety and effectiveness of etanercept so that treatment decisions can be further optimized.

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Conflict of interest Doctors T. Koike, M. Harigai, S. Inokuma, N. Ishiguro, J. Ryu, T. Takeuchi, Y. Tanaka, and H. Yamanaka are all members of the Etanercept Postmarketing Surveillance Committee of the Japan College of Rheumatology. It is the belief of the first author that this position does not constitute a Conflict of Interest. The doctors participated in the review and analysis of the PMS data in their capacity as Committee members and are so listed. The financial relationships of the authors with all manufacturers of biological products used in the management of RA are as follows. #1 is a research grant to the institute to which they are affiliated, #2 is a consulting fee, #3 is membership of a speakers' bureau, #4 is a full-time employee, and #5 is a previous employee of Pfizer. T. Koike, Abbott Japan, 1; Bristol-Myers Squibb, 1; Chugai Pharmaceutical Co. Ltd, 1; Eisai Co. Ltd, 1; Mitsubishi Tanabe Pharma, 1; Takeda Pharmaceutical Co. Ltd, 1; Wyeth KK, 1; Otsuka Pharmaceutical Co. Ltd, 2; Abbott Japan, 3; Bristol-Myers Squibb, 3; Chugai Pharmaceutical Co. Ltd, 3; Eisai Co. Ltd, 3; Mitsubishi Tanabe Pharma, 3; Takeda Pharmaceutical Co. Ltd, 3; Wyeth KK, 3; M. Harigai, Abbott Japan, 1; Bristol-Myers Squibb, 1; Chugai Pharmaceutical Co. Ltd, 1; Eisai Co. Ltd, 1; Mitsubishi Tanabe Pharma, 1; Pfizer Japan Inc., 1; Takeda Pharmaceutical Co. Ltd, 1; Abbott Japan, 2; Chugai Pharmaceutical Co. Ltd, 2; Mitsubishi Tanabe Pharma, 2; Abbott Japan, 3; Bristol-Myers Squibb, 3; Chugai Pharmaceutical Co. Ltd, 3; Eisai Co. Ltd, 3; Mitsubishi Tanabe Pharma, 3; Pfizer Japan Inc., 3; Takeda Pharmaceutical Co. Ltd, 3; S. Inokuma, None; N. Ishiguro, Abbott, 1; Chugai Pharmaceutical Co. Ltd, 1; Daiichi-Sankyo Pharmaceutical Co. Ltd, 1; Eisai Co. Ltd, 1; Mitsubishi Tanabe Pharma, 1; Takeda Pharmaceutical Co. Ltd, 1; Wyeth KK, 1; Abbott, 3; Bristol-Myers Squibb, 3; Chugai Pharmaceutical Co. Ltd, 3; Daiichi-Sankyo Pharmaceutical Co. Ltd, 3; Eisai Co. Ltd, 3; Mitsubishi Tanabe Pharma, 3; Takeda Pharmaceutical Co. Ltd, 3; Wyeth KK, 3; J. Ryu, None; T. Takeuchi, Bristol-Myers Squibb, 2; Mitsubishi Tanabe Pharma, 2; Novartis, 2; Abbott, 3; Chugai Pharmaceutical Co. Ltd, 3; Eisai Pharma, 3; Mitsubishi Tanabe Pharma, 3; Takeda Pharmaceutical Co. Ltd, 3; Y. Tanaka, Abbott, 1; Astellas Pharma Inc., 1; Chugai Pharmaceutical Co. Ltd, 1; Eisai Co. Ltd, 1; Mitsubishi Tanabe Pharma, 1; MSD KK, 1; Pfizer Inc., 1; Takeda Pharmaceutical Co. Ltd, 1; Mitsubishi Tanabe Pharma, 2; Abbott, 3; Astellas Pharma Inc., 3; Chugai Pharmaceutical Co. Ltd, 3; Eisai Co. Ltd, 3; Mitsubishi

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