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EXTENDED REPORT

A phase 2 dose-ranging study of subcutaneous tabalumab for the treatment of patients with active rheumatoid arthritis and an inadequate response to methotrexate

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ABSTRACT

Objectives To assess the dose-response relationship, efficacy and safety of tabalumab, a human monoclonal antibody that neutralises membrane-bound and soluble B-cell activating factor (BAFF), in patients with rheumatoid arthritis (RA) with inadequate response to methotrexate (MTX).

Methods In this phase 2, 24-week, double-blind, placebo-controlled, dose-ranging study, patients with RA (N=158) on stable MTX were randomised by Bayesian-adaptive method to receive 1, 3, 10, 30, 60, or 120 mg tabalumab or placebo subcutaneously every 4 weeks for 24 weeks. The primary objective was to test for a significant dose-response relationship using a statistical model of the proportion of patients having ≥50% improvement in American College of Rheumatology (ACR) criteria (ACR50) at week 24 (prespecified α=0.10).

Results At week 24, a significant dose-response relationship was observed using ACR50 (p=0.059) and ACR20 (p=0.044) response rates. Using model-estimated data, only 120 mg had significantly higher ACR50 and ACR20 response rates versus placebo (p<0.05). Observed response rates were significantly higher for 120 mg versus placebo as measured by ACR50 at weeks 12 (p=0.039) and 20 (p=0.018), but not week 24, and by ACR20 at weeks 12 (p=0.011) and 24 (p=0.039). Mean DAS28 C-reactive protein improved with 120 mg at week 24 (p=0.048). Frequency of TEAEs was similar across groups (range 50–69%, p=0.884). Ten (8.2%) tabalumab and 5 (13.9%) placebo patients reported a serious adverse event (SAE). Infections occurred more frequently in patients exposed to tabalumab (30.3% vs 19.4%). Serious infections were reported in 3 (2.5%) tabalumab-treated patients only.

Conclusions A dose-response relationship was detected with monthly subcutaneous tabalumab. A significant effect was detected with the 120 mg dose with no unexpected safety signals.

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INTRODUCTION

Rheumatoid arthritis (RA) affects approximately 1% of the population[1] and is characterised by joint inflammation that can lead to joint destruction and systemic complications.[2] Currently available biologic therapies selectively target key molecules associated with joint inflammation, but approximately 30% of patients will remain unresponsive to these treatments.[3]

B-cell activating factor (BAFF) is a tumour necrosis factor (TNF) family ligand that is increased in the sera and synovial fluid of patients with RA,[4–6] and is required for B-cell survival.[7] BAFF has two biologically active forms, a soluble and membrane-bound form,[8] and induces polyclonal maturation of immature and mature B cells involved in RA pathogenesis.[9 10]

Tabalumab is a human monoclonal antibody that neutralises soluble and membrane-bound BAFF.[11] In a previous study, intravenous tabalumab (30, 60 and 160 mg) reduced RA signs and symptoms in patients with an inadequate response to methotrexate (MTX-IR); although all doses were effective, no dose-response relationship in American College of Rheumatology (ACR) scores was observed.[12]

The current trial, which used Bayesian-adaptive randomisation, explored the dose-response relationship of tabalumab given subcutaneously once every 4 weeks (Q4W) to patients with active RA receiving stable doses of MTX.

METHODS

Patients

Patients were recruited from 64 centres in 12 countries (Argentina, Australia, Chile, Germany, Hungary, India, Mexico, Poland, Romania, Slovakia, Ukraine and the USA). All patients provided voluntary written informed consent. The study was approved by local Institutional Review Boards in accordance with the Declaration of Helsinki and applicable laws and regulations.

Patients were aged between 18 and 75 years, were taking MTX (10–25 mg/week) for ≥16 weeks, and met ACR (1987 revised) criteria for RA.[13] Major inclusion criteria included ≥5/28 swollen and ≥5/28 tender joints; ACR functional class I, II, or III; a history of, or a current, positive rheumatoid factor (RF+) test; a C-reactive protein (CRP) ≥1.2 times upper limit of normal (ULN; 1.0 mg/dl); and the absence of pregnancy or breast feeding.

Major exclusion criteria included use of any parenteral or oral corticosteroid at >10 mg/day of prednisone or its equivalent within 4 weeks of

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baseline; use of any B-cell biotherapies at any time; insufficient response to a TNF inhibitor (TNF-IR; patients who stopped for reasons other than lack of efficacy were eligible); presence of other autoimmune disorders; a positive protein derivative test for tuberculosis; or serious bacterial infections within 6 months of enrolment. Study participants must have discontinued etanercept ≥28 days before baseline, and infliximab, adalimumab, or other biologic TNF inhibitors ≥56 days before baseline.

Study design

This was a phase 2, 24-week, double-blind, placebo-controlled, dose-ranging study. Following two screening visits, patients were randomised by Bayesian-adaptive method[14] to receive placebo or tabalumab (1, 3, 10, 30, 60, or 120 mg) subcutaneously Q4W for 24 weeks (figure 1A). Patients maintained their prestudy stable dose of MTX and received their other usual medical treatments for RA or concomitant diseases as allowed by protocol.

Bayesian-adaptive randomisation used accumulating data from the ongoing trial to make progressive adjustments in dose-group assignments. Over time, these adjustments were expected to provide a more precise estimate of the dose-response relationship. Computer-generated random treatment assignments were made using an interactive voice-response system (IVRS). The first 35 patients were randomised to 1 of 7 treatment groups in equal number. Once 35 patients were randomised, there was a constant 20% chance of randomisation to placebo and an 80% chance of randomisation to one of the tabalumab doses. A contract research organisation periodically updated a non-informative, normal, dynamic linear model prior distribution, from which the posterior distribution of a 24-week, treatment-response model was derived. The posterior distribution was used to adjust the probability of randomisation to each tabalumab dose. As the study progressed, the probability of assignment to higher doses increased. Enrolment and randomisation ended when approximately 150 patients had been randomised. These 150 patients provided 80% power to detect a difference based on simulations using varying responses-to-dose assumptions as well as varying patient accrual and dropout rates.

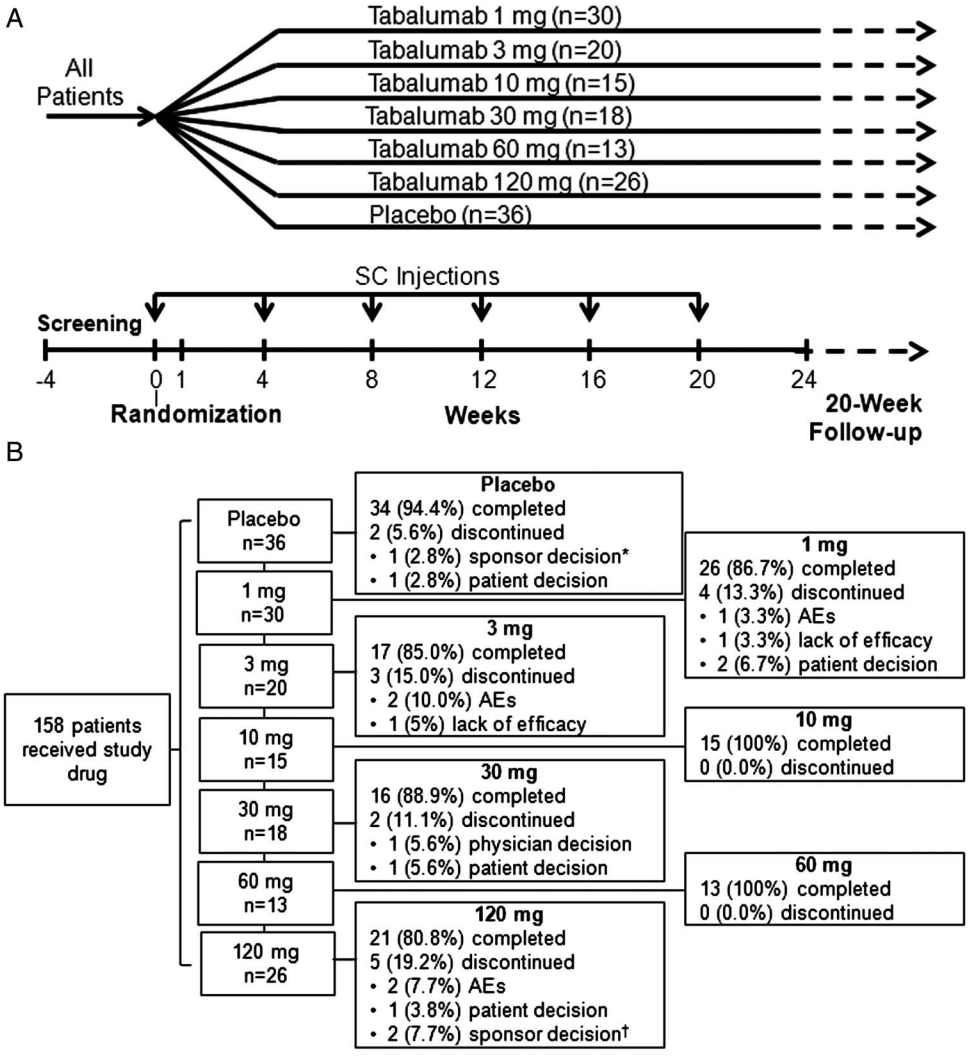

Figure 1 Study design and patient disposition. (A) Patients were adaptively randomised by Bayesian method. Over time, the probability of randomisation to more effective doses increased, resulting in unequal distribution of patients across groups. Study drug was administered six times by SC injection for 24 weeks. (B) Disposition of 158 randomised and treated patients through week 24. The percentage of patients who withdrew from the study and reasons for discontinuation are also shown for each group. *One patient in the placebo group was discontinued due to elevated liver enzymes at the sponsor's request. †Two patients in the 120 mg group were excluded from primary efficacy analyses due sponsor decision to withdraw a site for violation of good clinical practices. AEs, adverse events, SC, subcutaneous.

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Endpoints

The primary endpoint was to test for a significant dose-response relationship based on a statistical model of the proportion of patients having ≥50% improvement in ACR criteria (ACR50) at week 24. Key secondary efficacy endpoints included the dose-response relationship based on modelled ACR20 response rates at week 24; ACR20 and ACR50 response rates, change in DAS28, and percent change in CRP at each visit (weeks 1, 4, 8, 12, 16, 20 and 24); and pharmacokinetic parameters. Safety endpoints included incidences of adverse events (AEs) and serious adverse events (SAEs), and clinical laboratory test and immunogenicity results. Pharmacodynamic endpoints included changes over time in serum immunoglobulins (IgM, IgG, IgA), total B cells (CD20), and B-cell subsets (mature naïve (CD19, IgD+, CD27–), memory (CD19, IgD–, CD27)).

All efficacy measures used 28-joint counts and CRP for response calculations. Blood samples were collected at baseline and subsequent visits for determining serum tabalumab concentrations (weeks 1, 4, 8, 12, 16, 20 and 24), serum immunoglobulins (IgG, IgM and IgA) (weeks 1, 4, 16 and 24), and antitabalumab antibodies (weeks 4, 8 and 24). Serum tabalumab and immunoglobulin concentrations were assayed using validated methods. B cells were assayed by flow cytometry.[15]

Standard laboratory tests, including chemistry, haematology, urinalysis panels and ECGs were obtained at regular intervals. Vital signs were taken, and AEs and SAEs were recorded and summarised at each visit.

Statistical methods

All analyses were performed with the intent-to-treat population (all randomised patients who received ≥1 dose of study drug). Two 120 mg patients were excluded from primary efficacy analyses (site withdrawn due to good clinical practices violation). For ACR analyses, non-responder imputations (NRIs) were used for patients who discontinued early. For all other efficacy analyses, a last-observation-carried-forward (LOCF) approach was used.

The primary analysis tested for a significant ACR50 dose-related response over doses ranging from placebo to 120 mg tabalumab at week 24. The dose-response relationship was tested with a joint test of linear and quadratic dose response (regression model included terms for dose and dose^2) from the likelihood ratio test (α=0.10). The smallest dose achieving ≥95% of maximal efficacy (ED_{95}) and ACR50 response rates corresponding to each dose were estimated from the logistic regression model. This analysis was repeated for ACR20. The dose-response relationship for DAS28 was performed by Spearman non-parametric correlations with dose (1-sided, α=0.05).

ACR20 and ACR50 responses were summarised, and Fisher exact tests compared tabalumab observed response rates with placebo at all timepoints. Modelled response rates were compared using a 1-sided, z test with SE estimated using the delta method. For DAS28, pairwise comparisons of tabalumab doses versus placebo were performed using contrast statements within an analysis of covariance (ANCOVA) model with treatment as the fixed factor and baseline as a covariate. Pairwise comparisons were 1-sided (α=0.05).

Pharmacodynamic analyses were performed by 2-sided comparisons of all tabalumab doses combined versus placebo, using ranked ANCOVA with the standardised rank outcome variable, treatment as the fixed factor, and the standardised rank baseline value as a covariate (α=0.10). For CRP and serum immunoglobulins, 2-sided pairwise comparisons of tabalumab dose versus placebo were performed using contrast statements within ANCOVA (α=0.10).

Tabalumab pharmacokinetic parameters were analysed using a population approach implemented with NONMEM (ICON Development Solutions, Ellicott City, Maryland, USA). Clearance and distribution volume were characterised as a function of dose and treatment duration.

Safety data were descriptively summarised by treatment. Placebo and tabalumab doses were compared using a χ^2 test, or Fisher exact test if χ^2 assumptions were violated, for any event.

RESULTS

Patient disposition and characteristics

A total of 142/158 patients (89.9%) completed: 34/36 (94.4%) in the placebo group and 108/122 (88.5%) in all tabalumab groups combined (figure 1B). The most common reasons for early withdrawal were AEs and patient decision (figure 1B).

Overall, baseline characteristics were similar across groups with a few exceptions (table 1). In the 60 mg group, a smaller percentage were Caucasian (39%; range 53–73%, p=0.049 vs placebo). The placebo group had shorter disease duration than the 30 mg group (p=0.001). Baseline disease activity parameters and average MTX and prednisone doses were comparable across groups. Eleven patients had prior TNF-inhibitor exposure: 7 placebo (infliximab, n=2; adalimumab, n=2; investigational drug, n=1; etanercept, n=2), one 1 mg (infliximab), one 30 mg (etanercept), one 60 mg (infliximab), and one 120 mg tabalumab (investigational drug).

Clinical response

A significant (prespecified α=0.10) dose-response relationship was detected for ACR50 (p=0.059) and ACR20 (p=0.044) at week 24 using a regression model. The ED_{95} was 119.0 mg and 118.5 mg for ACR50 and ACR20, respectively.

Modelled and observed results for ACR50 and ACR20 responders at week 24 are presented in table 2. Using model-estimated data at week 24, only the 120 mg dose had significantly higher ACR50 and ACR20 response rates versus placebo (table 2).

The observed ACR50 response rate (NRI) was significantly higher with only the 120 mg dose versus placebo at week 12 (33.3% (8/24) vs 11.1% (4/36); p=0.039) and week 20 (33.3% (8/24) vs 8.3% (3/36); p=0.018), but not at week 24 (table 2, figure 2A). The ACR20 response rate (NRI) was significantly higher with 120 mg versus placebo at week 12 (66.7% (16/24) vs 33.3% (12/36); p=0.011), and week 24 (table 2, figure 2B). No other dose was significantly different from placebo at any timepoint for ACR20, except 60 mg at week 4 (38.5% (5/13) vs 11.1% (4/36); p=0.043) (figure 2B).

At baseline, DAS28 scores were similar across groups (table 1). DAS28 score significantly improved from baseline with 120 mg versus placebo at week 24 (table 2); this improvement was also observed at earlier timepoints (figure 2C). No other dose showed a significant DAS28 improvement.

At week 24, tender joint counts, swollen joint counts, and patient's assessments of disease activity and pain were similar across groups, whereas physician's assessment of disease activity was significantly reduced with 120 mg (table 3). Mean CRP was similar between placebo and tabalumab groups at week 24. In some patients, elevated CRP at screening was no longer elevated at baseline. A posthoc analysis of patients with baseline CRP>ULN was conducted by treatment group (placebo, 1, 3, 10, 30, 60 and 120 mg). For these groups, baseline CRP was 38.1, 27.0, 45.0, 38.4, 30.5, 37.4 and 30.2 mg/dl, and mean CRP

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Table 1 Baseline demographics and clinical parameters (ITT population)

		Tabalumab					
	Placebo (N=36)	1 mg (N=30)	3 mg (N=20)	10 mg (N=15)	30 mg (N=18)	60 (mg N=13)	120 mg (N=26)
Gender, n (%)							
Male	6 (16.7)	4 (13.3)	6 (30.0)	3 (20.0)	3 (16.7)	1 (7.7)	8 (30.8)
Female	30 (83.3)	26 (86.7)	14 (70.0)	12 (80.0)	15 (83.3)	12 (92.3)	18 (69.2)
Age, years	50.6±11.7	54.6±11.7	53.4±10.8	51.2±13.8	54.5±11.8	44.4±13.8	50.7±12.0
Race/ethnicity, n (%)							
Caucasian	25 (69.4)	22 (73.3)	13 (65.0)	8 (53.3)	11 (61.1)	5 (38.5)*	17 (65.4)
Hispanic	8 (22.2)	5 (16.7)	6 (30.0)	6 (40.0)	6 (33.3)	5 (38.5)	8 (30.8)
East Asian	2 (5.6)	2 (6.7)	1 (5.0)	0	1 (5.6)	3 (23.1)	1 (3.8)
West Asian	0	0	0	1 (6.7)	0	0	0
African	1 (2.8)	1 (3.3)	0	0	0	0	0
Disease duration, years	6.2±7.2	9.3±7.9	8.4±7.0	9.0±6.7	11.7±7.0*	10.1±9.3	9.9±11.6
CRP above ULN at baseline, n (%)	28 (77.8)	21 (70.0)	15 (75.0)	12 (80.0)	15 (83.3)	12 (92.3)	19 (73.1)
HAQ-DI	1.8±0.5	1.7±0.5	1.8±0.5	1.7±0.6	1.5±0.9	1.8±0.6	1.7±0.5
DAS28-CRP	6.2±1.0	6.4±0.8	6.1±1.0	6.2±1.0	6.1±0.8	6.1±0.9	5.9±0.7
MTX dose, mg/week	16.8±4.5	15.7±4.6	14.9±4.2	16.0±4.4	14.7±3.3	16.2±3.9	14.8±3.5
Prednisone dose, mg/day	6.6±2.2	7.6±2.6	8.1±2.1	7.1±2.0	6.8±2.2	6.0±2.4	5.6±2.0
CD20 B-cell counts, cells/μl	193.4±130.3	220.8±133.2	208.1±171.8	189.5±64.1	220.1±181.6	212.5±122.1	187.4±110.0

All values are mean±SD, unless otherwise noted.
*p<0.05 versus placebo.
CRP, C reactive protein; CRP; HAQ-DI, Health Assessment Questionnaire-Disability Index; DAS28-CRP, Disease Activity Score based on 28-joint count; ITT, Intent-to-treat population; MTX, methotrexate; ULN, upper limit of normal.

(LOCF) at week 24 was 16.8, 8.1, −8.8, 21.8, 46.2, 38.9 and 38.5 mg/dl, respectively.

Pharmacokinetics

Serum tabalumab concentrations demonstrated non-linear elimination. The time to reach maximum concentration following subcutaneous injection at steady state ($T_{max,ss}$) and the half-life over the 4-week dosing interval ($t_{1/2,tau}$) increased with increasing dose. At 120 mg, $T_{max,ss}$ and $t_{1/2,tau}$ were 5.0 and 21.6 days, respectively.

Biologic activity

CD20 B-cell counts across all tabalumab doses initially increased at week 1 (mean change in all doses combined vs placebo: 67.75 vs 14.64 cells/μl, p<0.001) and subsequently decreased back to baseline or below it starting from week 4 (15.03 vs 6.91 cells/μl, p=0.371) and continuing through week 24 (−34.56 vs 17.63 cells/μl, p=0.008). Of the CD19 cell subsets, only mature, naïve cell counts showed a pattern similar to CD20 cells (week 1: 38.17 vs 15.70 cells/μl, p=0.024; week 24: −45.63 vs 18.06 cells/μl, p<0.001). Increases in memory cell counts from baseline were observed across groups over 24 weeks, with the largest increases in the higher dose groups. At week 24, only 30 mg (p=0.006), 60 mg (p=0.003) and 120 mg (p<0.001) demonstrated statistically significant increases in memory B cells versus placebo.

Mean IgM and IgA levels tended to be lower than baseline for all tabalumab groups at weeks 16 and 24. For IgM, this difference was statistically significant at week 16 with the 30 mg (−0.15±0.40 g/l, p=0.016), 60 mg (−0.24±0.27 g/l, p=0.028), and 120 mg (−0.17±0.32 g/l, p=0.005) versus placebo (0.02±0.39 g/l), and at week 24 with 30 mg (−0.08±0.53 g/l, p=0.023), 60 mg (−0.30±0.30 g/l, p=0.004), and 120 mg (−0.09±0.38 g/l, p=0.017) versus placebo (0.02±0.32 g/l). For

Table 2 Summary of efficacy endpoints at week 24

		Tabalumab						
	Placebo (N=36)	1 mg (N=30)	3 mg (N=20)	10 mg (N=15)	30 mg (N=18)	60 mg (N=13)	120 mg (N=24)	p Value
ACR50 obs, n (%) (p)*†	8 (22.2)	3 (10.0) (0.896)	2 (10.0) (0.940)	5 (33.3) (0.311)	2 (11.1) (0.919)	1 (7.7) (0.954)	9 (37.5) (0.255)	NA
ACR50 model, % (p) *†	18.1	17.7 (0.827)	17.0 (0.828)	15.0 (0.833)	11.8 (0.834)	11.8 (0.767)	37.0 (0.037)	(0.059)‡
ACR20 obs, n (%) (p) *†	16 (44.4)	12 (40.0) (0.730)	9 (45.0) (0.594)	7 (46.7) (0.563)	11 (61.1) (0.193)	7 (53.8) (0.397)	17 (70.8) (0.039)	NA
ACR20 model, % (p) *†	43.2	43.6 (0.171)	44.3 (0.170)	46.9 (0.162)	53.5 (0.132)	61.5 (0.070)	70.1 (0.005)	(0.044)‡
DAS28, mean±SD change from baseline (p)§¶	−1.5±1.3	−1.5±1.3 (0.457)	−1.0±1.1 (0.874)	−1.7±1.0 (0.278)	−1.5±1.3 (0.357)	−1.6±1.2 (0.271)	−1.9±1.2 (0.048)	(0.071)**

*1-sided Fisher exact test that the tabalumab group has more responders than placebo.
†Imputed by non-responder imputation.
‡Likelihood ratio test of quadratic logistic regression model, testing the existence of a dose-response with a prespecified 2-sided type 1 error rate of 0.10.
§1-sided pairwise comparison using contrast statements with an analysis of covariance (ANCOVA) model with treatment as the fixed factor and baseline as the covariate.
¶Imputed by last-observation-carried-forward.
**Dose-response relationship from 1-sided Spearman non-parametric correlation analysis.
ACR50, American College of Rheumatology 50 responder index; ACR20, American College of Rheumatology 20 responder index; DAS28, Disease Activity Score based on 28-joint count; obs, actual observations; model, assessed using a statistical model; NA, not available; p, p Value.

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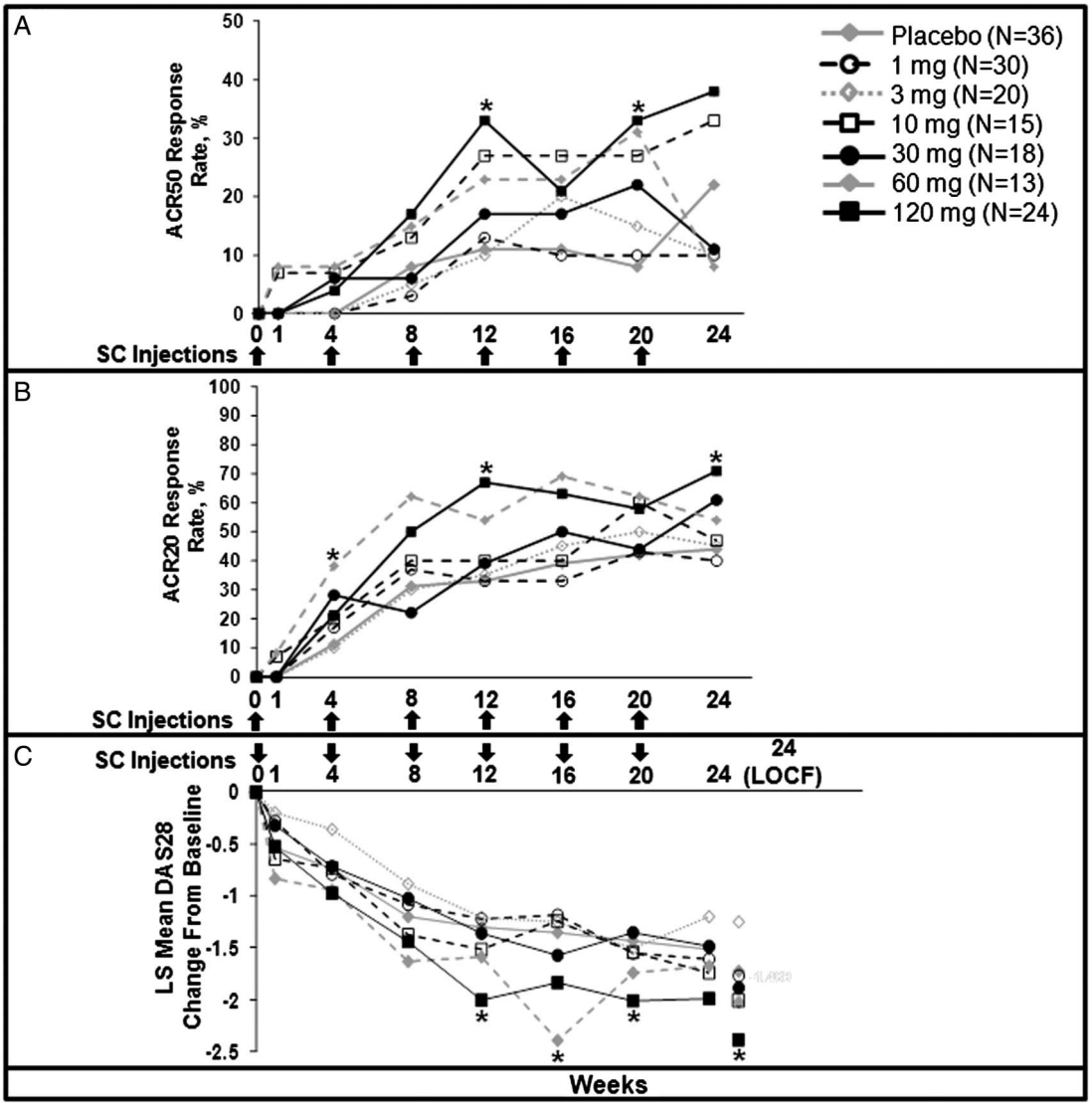

Figure 2 (A) Values are observed changes in the ACR50 response rate at each treatment visit with the tabalumab dose or placebo; non-responders imputed. p Values are based on 1-sided Fisher exact test that the tabalumab group has more responders than placebo. *p≤0.05 Versus placebo. (B) Values are observed changes in the ACR20 response rate at each treatment visit with the tabalumab dose or placebo; non-responders imputed. p Values are based on 1-sided Fisher exact test that the tabalumab group has more responders than placebo. *p≤0.05 Versus placebo. (C) Time course of LS mean change in DAS28 score at each treatment visit with the tabalumab dose or placebo. p Values are based on 1-sided pairwise comparison of the tabalumab dose level versus placebo using contrast statements within an ANCOVA model with treatment as the fixed factor and the baseline value as a covariate. *p≤0.05 Versus placebo. ACR20, proportion of responders having ≥20% improvement according to American College of Rheumatology (ACR) criteria; ACR50, proportion of patients having ≥50% improvement according to ACR criteria; ANCOVA, analysis of covariance; DAS28, Disease Activity Score based on 28-joint count; LOCF, last-observation-carried-forward, LS mean, least squares mean; SC, subcutaneous.

IgA, this difference was statistically significant at week 16, with 120 mg versus placebo (−0.32±0.42 vs −0.12±0.50 g/l, p=0.040), and at week 24 with 60 mg (−0.42±0.45 g/l, p=0.025) and 120 mg (−0.25±0.63 g/l, p=0.030) versus placebo (−0.12±0.48 g/l). Mean IgG levels were not significantly different from placebo with any tabalumab dose at any timepoint. There was no association between changes in serum immunoglobulin levels and occurrence of infection or other AEs. There was no correlation between clinical response, B-cell counts, or serum immunoglobulin levels.

Safety

The frequency of treatment-emergent adverse events (TEAEs) was similar across tabalumab doses (range 50–69%, p=0.884) (table 4). The majority of TEAEs were mild or moderate in severity with no obvious trends in the nature or frequency by dose. The most frequently reported TEAEs were injection-site pain and upper respiratory tract infection (table 4). Five patients discontinued due to an AE, all of whom were treated with tabalumab 1 mg (hemiplegia), 3 mg (prolonged QT and RA worsening (exacerbation/flare)), or 120 mg (diverticulitis and RA worsening). The incidence of SAEs was 13.9% (5/36 patients) with placebo and ranged from 3.8% (1/26 patients) with 120 mg to 16.7% (3/18 patients) with 30 mg (table 4). RA worsening was the only SAE reported in >1 patient (n=2, placebo). No patients died during the study.

Infection was reported at a higher incidence with tabalumab (30.3% (37/122 patients)) than placebo (19.4% (7/36 patients)), but did not increase with higher doses. The lowest incidence of infection was observed with 120 mg (11.5%). Serious infections

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Table 3 Composite score components at baseline and at week 24 (LOCF)

		Tabalumab					
	Placebo (N=36)	1 mg (N=30)	3 mg (N=20)	10 mg (N=15)	30 mg (N=18)	60 mg (N=13)	120 mg (N=24)
Swollen joint count (28)							
Baseline	12.0±5.5	13.4±5.8	11.2±5.0	15.4±6.3	12.7±5.6	13.1±5.4	11.9±6.1
Week 24	6.3±4.7	6.5±5.5	5.5±5.8	6.5±5.3	7.3±7.6	6.9±4.8	4.3±3.5
Tender joint count (28)							
Baseline	16.0±6.5	17.8±7.2	16.5±6.3	16.5±8.3	15.4±5.8	15.2±6.3	13.7±6.3
Week 24	8.9±6.7	10.5±9.9	12.0±9.6	7.8±5.9	7.7±6.9	7.8±7.4	6.2±5.9
Physician's global assessment of disease activity							
Baseline	61.8±16.6	61.5±20.2	57.9±16.5	67.2±14.6	59.9±16.8	53.0±13.7	61.6±14.8
Week 24	39.0±23.7	38.7±24.3	31.7±22.5	37.0±22.3	31.6±21.7	31.6±20.8	25.8±14.2*
Patient's global assessment of disease activity							
Baseline	70.3±19.4	71.6±17.5	66.8±18.3	61.6±20.8	64.5±24.3	60.9±20.1	69.4±15.0
Week 24	46.1±26.2	50.9±27.5	50.8±21.2	39.1±28.4	45.9±23.7	45.5±17.3	35.6±20.2
Patient's global assessment of pain							
Baseline	66.3±23.7	66.4±20.2	63.1±18.4	61.5±23.4	59.5±23.7	64.4±16.1	66.2±15.3
Week 24	46.8±25.6	48.6±26.1	50.4±20.8	40.3±28.6	46.4±25.3	46.8±16.9	36.4±21.0
CRP, mg/dl							
Baseline	3.1±2.5	2.1±1.5	3.5±3.8	3.3±2.2	2.7±2.2	3.5±3.2	2.5±2.9
Week 24	2.2±1.8	1.8±1.4	3.5±3.1	2.7±1.3	1.8±1.9	1.5±0.9	1.7±1.3

All values are mean±SD. Analyses are 2-sided pairwise comparisons of tabalumab dose versus placebo were performed using contrast statements within ANCOVA.
*p<0.05 versus placebo.
CRP, C-reactive protein; LOCF, last-observation-carried-forward; p, p Value.

were reported in three (2.5%) tabalumab-treated patients. One patient discontinued due to a serious infectious event of H1N1 influenza pneumonia 24 days after a single 30 mg dose.

No clinically relevant differences or trends were identified in vital signs, ECG, or chemistry, haematology and urinalysis panels. The percentage of patients with abnormal laboratory values was comparable across groups, with no apparent trends.

Immunogenicity

At 24 weeks, two tabalumab-treated patients had treatment-emergent antitabalumab antibodies (TEAb (fourfold increase from baseline); 1 mg and 120 mg group). One patient seroconverted at week 4, the other at week 8, and TEAb persisted to week 24. None were neutralising. Two additional patients (1 mg and 10 mg group) and 2 placebo patients had transient TEAb detected at a single sampling. One placebo patient had neutralising antibodies at week 24. The presence of TEAb did not appear to have an effect on ACR response. None of the tabalumab-treated patients who tested positive for antibodies experienced an SAE or serious infection, nor did they discontinue due to lack of efficacy. There was no discernible reduction in tabalumab serum concentrations in the presence of antitabalumab antibodies and no dose-related trends.

Table 4 Overview of adverse events at week 24

		Tabalumab						
Number of patients (%)	Placebo (N=36)	1 mg (N=30)	3 mg (N=20)	10 mg (N=15)	30 mg (N=18)	60 mg (N=13)	120 mg (N=26)	All tabalumab doses combined (N=122)
Deaths	0	0	0	0	0	0	0	0
SAEs	5 (13.9)	4 (13.3)	0	0	3 (16.7)	2 (15.4)	1 (3.8)	10 (8.2)
Patients who discontinued due to an AE	0	1 (3.3)	2 (10.0)	0	0	0	2 (7.7)	5 (4.1)
TEAEs	22 (61.1)	20 (66.7)	12 (60)	9 (60)	12 (66.7)	9 (69.2)	13 (50)	75 (61.5)
TEAEs that occurred in ≥3% of patients in the combined tabalumab group								
Injection-site pain	0	1 (3.3)	0	4 (26.7)	1 (5.6)	1 (7.7)	2 (7.7)	9 (7.4)
Upper respiratory tract infection	1 (2.8)	3 (10.0)	0	3 (20.0)	0	2 (15.4)	1 (3.8)	9 (7.4)
RA worsening	7 (19.4)	3 (10.0)	2 (10.0)	0	0	0	1 (3.8)	6 (4.9)
Hypertension	0	1 (3.3)	1 (5.0)	0	1 (5.6)	0	2 (7.7)	5 (4.1)
Anemia	0	1 (3.3)	2 (10.0)	1 (6.7)	0	1 (7.7)	0	5 (4.1)
Pharyngitis	0	0	1 (5.0)	1 (6.7)	1 (5.6)	0	1 (3.8)	4 (3.3)
Pyrexia	1 (2.8)	2 (6.7)	1 (5.0)	0	0	0	1 (3.8)	4 (3.3)
Urinary tract infection	1 (2.8)	1 (3.3)	2 (10.0)	1 (6.7)	0	0	0	4 (3.3)
Weight increased	0	1 (3.3)	1 (5.0)	0	1 (5.6)	1 (7.7)	0	4 (3.3)
Respiratory tract infection	0	1 (3.3)	2 (10.0)	0	0	1 (7.7)	0	4 (3.3)
Nasopharyngitis	2 (5.6)	2 (6.7)	0	0	2 (11.1)	0	0	4 (3.3)

AE, adverse event; RA, rheumatoid arthritis; SAEs, serious adverse events; TEAEs, treatment-emergent adverse events.
Percentages are based on the number of patients in each column.

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DISCUSSION

In the current study, which used Bayesian-adaptive randomisation, a statistically significant dose-response relationship was detected using regression models of ACR50 and ACR20 response rates at 24 weeks for tabalumab, administered monthly by subcutaneous injection over a 1 mg to 120 mg dose range. Significant efficacy versus placebo was detected with the 120 mg dose for ACR50 (by regression analysis only), ACR20, and DAS28 at week 24.

The principles and potential utility of B-cell depletion in RA treatment have been recognised since 2001. [16] Subsequent work resulted in the approval of rituximab for RA treatment in TNF-IR patients.[17] The concept of effective RA treatment through B-cell inhibition (without profound depletion) has been explored with limited success. Belimumab, a monoclonal antibody that neutralises soluble BAFF, was studied but not developed as an RA treatment after modest phase 2 results.[18 19] Atacicept, a fusion protein targeting BAFF and APRIL (a proliferation–inducing ligand), failed to achieve clinical benefits in two RA trials.[20 21]

Tabalumab neutralises membrane-bound and soluble BAFF and, thus, may have different biologic action and clinical outcomes than belimumab, atacicept and briobacept (a recombinant BAFF receptor immunoglobulin fusion protein). In an earlier study, intravenous tabalumab (30, 60 and 160 mg) administered every 3 weeks (Q3W) for 6 weeks demonstrated clinical efficacy in biologic-naïve patients with RA.[12] The present trial evaluated subcutaneous tabalumab across a wider range of doses given Q4W for 6 months. Adjusting for bioavailability and differences in dosing frequency, a subcutaneous Q4W 120 mg dose provides exposure equivalent to an intravenous Q3W 45 mg dose (data not shown).

Overall, no unexpected safety signal was detected in this relatively small study; tabalumab had a consistent safety profile across dose groups and placebo. Infectious events were more frequent with tabalumab than placebo, although the frequency did not increase with higher doses. Mean IgM and IgA levels tended to be lower than baseline in all tabalumab groups in the later weeks of the study; changes from baseline were not associated with increases in AEs or infectious AEs.

Within 1 week of the first injection, CD20 B cells and CD19 mature, naïve cells transiently increased with all tabalumab doses before decreasing back to baseline levels or below without profound reductions. Early increases in B cells with subsequent decreases have been observed with other BAFF-targeted therapies, such as briobacept[22] and atacicept.[23] No correlations were observed between B-cell changes and clinical efficacy or safety with tabalumab.

The following limitations of this study should be considered. Bayesian-adaptive randomisation was intended to estimate the dose-response relationship more precisely and to allocate patients to more effective dosing. However, patients were enrolled faster than planned, and updates to randomisation probabilities were not frequent enough. As a result, a higher percentage of patients were assigned to placebo or very low dose treatments. Despite this, a dose-response relationship was detected. Additionally, this was a short-term study with small treatment arms, and ACR50, rather than ACR20, was the primary endpoint. This study enrolled patients with active RA who were taking MTX, but not patients who previously failed TNF inhibitors or other biologic RA treatments, and only included RF+ patients. These findings cannot be generalised to patients on other background RA medications, with a history of exposure to a larger repertoire of agents, or who are seronegative. Despite these limitations, tabalumab demonstrated efficacy in MTX-IR patients.

In the present study, subcutaneous 120 mg tabalumab appeared to reduce RA signs and symptoms in patients taking concomitant MTX. Overall, based on a limited number of exposures in this phase 2 study, tabalumab had no unexpected safety signals. After this study was completed, phase 3 clinical trials were undertaken using tabalumab in patients with RA. These trials were recently discontinued after interim analyses provided results that did not meet efficacy expectations.[24 25] No safety concerns were noted.

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