

ORIGINAL ARTICLE

Ligelizumab for Chronic Spontaneous Urticaria

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ABSTRACT

BACKGROUND

In the majority of patients with chronic spontaneous urticaria, most currently available therapies do not result in complete symptom control. Ligelizumab is a next-generation high-affinity humanized monoclonal anti-IgE antibody. Data are limited regarding the dose–response relationship of ligelizumab and the efficacy and safety of ligelizumab as compared with omalizumab and placebo in patients who have moderate-to-severe chronic spontaneous urticaria that is inadequately controlled with H₁-antihistamines at approved or increased doses, alone or in combination with H₂-antihistamines or leukotriene-receptor antagonists.

METHODS

In a phase 2b dose-finding trial, we randomly assigned patients to receive ligelizumab at a dose of 24 mg, 72 mg, or 240 mg, omalizumab at a dose of 300 mg, or placebo, administered subcutaneously every 4 weeks for a period of 20 weeks, or a single 120-mg dose of ligelizumab. Disease symptoms of hives, itch, and angioedema were monitored by means of weekly activity scores. The main objective was to determine a dose–response relationship for the complete control of hives (indicated by a weekly hives-severity score of 0, on a scale from 0 to 21, with higher scores indicating greater severity); the primary end point of this response was assessed at week 12. Complete symptom control was indicated by a weekly urticaria activity score of 0 (on a scale from 0 to 42, with higher scores indicating greater severity). Safety was analyzed throughout the trial.

RESULTS

A total of 382 patients underwent randomization. At week 12, a total of 30%, 51%, and 42% of the patients treated with 24 mg, 72 mg, and 240 mg, respectively, of ligelizumab had complete control of hives, as compared with 26% of the patients in the omalizumab group and no patients in the placebo group. A dose–response relationship was established. At week 12, a total of 30%, 44%, and 40% of the patients treated with 24 mg, 72 mg, and 240 mg, respectively, of ligelizumab had complete control of symptoms, as compared with 26% of the patients in the omalizumab group and no patients in the placebo group. In this small and short trial, no safety concerns regarding ligelizumab or omalizumab emerged.

CONCLUSIONS

A higher percentage of patients had complete control of symptoms of chronic spontaneous urticaria with ligelizumab therapy of 72 mg or 240 mg than with omalizumab or placebo. (Funded by Novartis Pharma; ClinicalTrials.gov number, NCT02477332.)

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CHRONIC SPONTANEOUS URTICARIA IS A skin disorder that is characterized by the occurrence of itchy wheals (hives), angioedema, or both for 6 weeks or more in the absence of specific external stimuli.^{1,2} Studies have indicated that chronic spontaneous urticaria has a prevalence of approximately 1 case per 200 persons in adult European populations.³⁻⁵ Evidence suggests that chronic spontaneous urticaria has a long duration^{6,7} and can have a negative effect on quality of life.^{6,8} The pathogenesis of chronic spontaneous urticaria is not fully clear, but it is known to involve autoimmune mechanisms in most patients by means of IgG autoantibodies against the high-affinity receptor for the Fc region of IgE (FcεRI) or less frequently against IgE and IgE autoantigens.⁹⁻¹³

Standard first-line treatment of chronic spontaneous urticaria consists of nonsedating (second-generation) H₁-antihistamines at locally approved doses, with escalation up to four times the approved dose used off-label as second-line treatment; even so, the disease is still uncontrolled in some patients.^{1,14,15} The revised guidelines regarding the treatment of urticaria from the European Academy of Allergology and Clinical Immunology (EAACI), the Global Allergy and Asthma European Network (GA²LEN), the European Dermatology Forum (EDF) and the World Allergy Organization (WAO)¹ recommend the use of omalizumab as add-on third-line therapy to H₁-antihistamines. The goal of therapy, as noted in the current treatment guidelines, is complete control of the disease, yet some patients have symptoms that remain uncontrolled with the current standard of care.¹ To date, add-on therapy with omalizumab has been the most effective treatment in patients who continue to have symptoms while they are taking H₁-antihistamines,¹⁶⁻¹⁸ but some patients do not have a response, and better treatment options are needed.¹⁹⁻²²

Ligelizumab is a new high-affinity humanized monoclonal anti-IgE antibody that has previously shown dose-dependent and time-dependent suppression of free IgE, basophil FcεRI, basophil surface IgE, and skin-prick test responses to allergen that was superior in extent and duration to that observed with omalizumab.²³ In a phase 2b trial, we examined the efficacy and safety of ligelizumab as compared with omalizumab and

placebo in patients with chronic spontaneous urticaria that was inadequately controlled with standard-of-care therapy including H₁-antihistamines.

METHODS

TRIAL DESIGN AND OVERSIGHT

This phase 2b, dose-finding, multicenter, randomized, double-blind, active-controlled and placebo-controlled trial was designed to establish a dose–response relationship for ligelizumab and to evaluate its efficacy and safety as compared with omalizumab and placebo (Fig. 1). The trial consisted of a screening period (days –14 to 1), a treatment period (days 1 to 140 [20 weeks]), and a follow-up period after the cessation of the trial regimens (days 141 to 309 [24 weeks]). Patients who remained in the follow-up period for at least 12 weeks and had active disease (defined as a mean weekly urticaria activity score [measured twice daily] of ≥12; scores range from 0 to 42, with higher scores indicating greater severity; minimally important difference [MID], 9.5 to 10.0 points)²⁴⁻²⁷ were eligible to enter an extension study (ClinicalTrials.gov number, NCT02649218) from week 32 onward. The trial protocol, with the statistical analysis plan, is available with the full text of this article at NEJM.org.

This trial was designed and sponsored by Novartis Pharma. The institutional review board at each participating center approved the protocol. Patients provided written informed consent before any assessment was performed. Data were collected by the trial investigators according to Good Clinical Practice guidelines and were analyzed by the sponsor. The first draft of the manuscript was written by a medical writer paid by the sponsor, with critical input and approval from all authors. All the authors critically reviewed each manuscript draft, provided substantial input on the content, and made the decision to submit the manuscript for publication. The authors vouch for the accuracy and completeness of the data and for the fidelity of the trial to the protocol.

PATIENTS

Eligible patients were 18 to 75 years of age and had chronic spontaneous urticaria that was inadequately controlled with H₁-antihistamines at

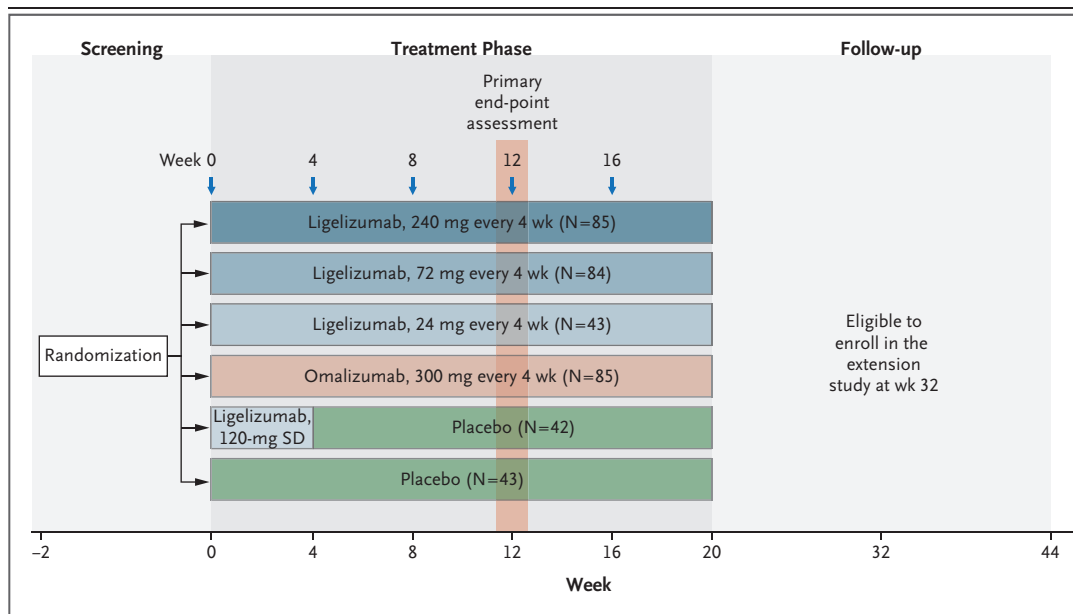


Figure 1. Trial Design.

The single 120-mg dose of ligelizumab was chosen in order to characterize the pharmacokinetics and pharmacodynamics of ligelizumab. Data from this group were used to assess the duration of response and to correlate this information with the concentration of drug in the serum at the time when symptoms reappeared. Patients who remained in the follow-up period for at least 12 weeks and had active disease (weekly urticaria activity score of ≥ 12 on a scale from 0 to 42, with higher scores indicating greater severity) could enter an extension study from week 32 onward. SD denotes single dose.

approved or increased doses alone or in combination with H₂-antihistamines or leukotriene-receptor antagonists. Key inclusion criteria were the following: moderate-to-severe chronic spontaneous urticaria, defined as a weekly urticaria activity score of at least 16 (disease activity categories are as follows: 0, symptom free; 1 to 6, well-controlled urticaria; 7 to 15, mild urticaria; 16 to 27, moderate urticaria; and 28 to 42, severe urticaria²⁶); a weekly hives-severity score of at least 8 (scores range from 0 to 21, with higher values indicating greater severity; MID, 5.0 to 5.5 points)²⁷ during the 7 days before randomization (day 1); and an in-clinic urticaria activity score of at least 4 (scores range from 1 to 6, with higher values indicating greater severity; MID, not available) on at least one of the screening visit days. Key exclusion criteria were the following: previous exposure to omalizumab or ligelizumab, any other skin disease that is associated with chronic itching that might confound the trial evaluations and results, and a clearly defined underlying cause of chronic urticaria other than

chronic spontaneous urticaria (e.g., inducible urticaria).

The demographic and clinical characteristics of the patients and the disease activity were elicited or measured at baseline. The Chronic Urticaria Index (CU Index; Viracor-IBT Laboratories), an *in vitro* basophil histamine-release assay in which a serum sample obtained from a patient is mixed with donor basophils and the released histamine levels are measured by a quantitative enzyme immunoassay, was performed at baseline. A positive CU Index (scores range from 1 to 50, with scores ≥ 10 representing a positive result) indicates that a patient has either an autoimmune basis for the urticaria or an alternative histamine-releasing factor that has been associated with greater disease severity than that in patients with a negative CU Index.^{28,29} The serum samples for this analysis were obtained centrally and analyzed by Viracor-IBT Laboratories. From the time that samples were obtained until analysis, they were kept frozen to ensure sample integrity.

TRIAL PROCEDURES

Patients were randomly assigned, in a 2:2:2:1:1:1 ratio, to receive one of the following trial regimens: 240 mg of ligelizumab every 4 weeks, 72 mg of ligelizumab every 4 weeks, 300 mg of omalizumab every 4 weeks, 24 mg of ligelizumab every 4 weeks, placebo every 4 weeks, or a single 120-mg dose of ligelizumab at week 0 followed by placebo every 4 weeks; the treatment period was 20 weeks (Fig. 1). The single 120-mg dose of ligelizumab was used to gain blinded washout information, to determine the level of ligelizumab in serum that was associated with the return of itch and hives symptoms, and to identify an appropriate administration interval for phase 3 trials.

Adjustments to the doses of trial agents were not permitted. Nonsedating H₁-antihistamines were used as background medication and as rescue medication as needed during the screening, treatment, and follow-up periods. This trial required concurrent use of H₁-antihistamines (at locally approved doses or at increased doses up to four times the locally approved dose) alone or in combination with H₂-antihistamines or leukotriene-receptor antagonists (montelukast, zafirlukast, or pranlukast) as background medication. It was recommended that patients continue taking a stable background medication throughout the trial, and patients were not permitted to switch their rescue medications.

END-POINT MEASURES

The main objective of the trial was to establish a dose–response relationship with respect to the achievement of complete hives response (weekly hives-severity score of 0) at week 12. Complete hives response was chosen as the primary end point because hives were considered to be a more objective symptom than itch and are specific to chronic spontaneous urticaria; itch is subjective and nonspecific and can be of different origins. We chose week 12 as the time point for the assessment of the primary end point because this was also the time point for the primary end point in phase 3 trials of omalizumab.^{19,20}

Secondary end points included the following: the efficacy of specific ligelizumab doses of 24 mg, 72 mg, and 240 mg as compared with omalizumab at a dose of 300 mg with respect to the achievement of complete hives response at weeks 12 and 20; the efficacy of ligelizumab at doses

of 24 mg, 72 mg, and 240 mg as compared with placebo and with omalizumab at a dose of 300 mg in the mean change from baseline in the weekly hives-severity score, in the weekly itch-severity score (scores range from 0 to 21, with higher scores indicating greater severity; MID, 4.5 to 5.0 points), in the weekly urticaria activity score, and in the weekly angioedema activity score (scores range from 0 to 105, with higher scores indicating greater severity; MID, not available); and the safety of ligelizumab as compared with placebo and with omalizumab at a dose of 300 mg during the 20 weeks of the treatment phase and 24 weeks of follow-up.

STATISTICAL ANALYSIS

The null hypothesis of a constant dose–response curve for the primary efficacy end point was tested at a significance level of 5% against the one-sided alternative hypothesis of a nonconstant dose–response curve with the use of the multiple comparison procedure and modeling method, which indicates the potential dose–response relationship.^{30,31} The dose–response curve was modeled as a weighted average of prespecified model candidates with the use of bootstrap sampling. The dose–response model was derived from testing a range of ligelizumab doses, starting at 24 mg and then 72 mg and 240 mg, with placebo considered to be the zero dose. For the primary variable, patients who had data missing because of discontinuation before the week 12 assessment were considered not to have had a response, regardless of their last available assessment of the weekly hives-severity score.

Since the statistical analysis plan did not include a provision for correcting for multiplicity when tests were conducted for secondary or other outcomes, results are reported as point estimates with 95% confidence intervals. This was a small and clinically nondirective trial. The widths of the confidence intervals have not been adjusted for multiplicity, so the intervals should not be used to infer any effects for secondary outcomes.

RESULTS**PARTICIPANTS**

A total of 574 patients were screened, of whom 382 were randomly assigned to a trial group. A

total of 338 patients (88%) completed the treatment phase of the trial (Fig. S1 in the Supplementary Appendix, available at NEJM.org). There were no notable imbalances among the trial groups regarding the demographic and clinical characteristics of the patients at baseline (Table 1).

PRIMARY END POINT

The main objective of the trial was achieved, with ligelizumab showing a dose–response relationship with respect to the achievement of complete hives response (weekly hives-severity score of 0) at week 12 (primary end point) in patients with chronic spontaneous urticaria. The dose–response curve showed a steep dose–response relationship with a plateau starting close to the 72-mg dose of ligelizumab (Fig. 2A); no further improvement in response was noted with the 240-mg dose.

SECONDARY END POINTS

At week 12, a total of 30%, 51%, and 42% of the patients treated with 24 mg, 72 mg, and 240 mg, respectively, of ligelizumab had a complete hives response, as compared with 26% of the patients in the omalizumab group (Fig. 2B) and no patients in the placebo group. The response with the 72-mg dose of ligelizumab was maintained at week 20, with 51% of the patients having a response, as compared with 34% of those in the omalizumab group. A total of 45% of the patients in the group that received the 240-mg dose of ligelizumab had a response. The response regarding itch severity (weekly itch-severity score of 0) showed a pattern similar to that seen with the weekly hives-severity score. Details are provided in Figure S2 in the Supplementary Appendix.

At week 12, a total of 30%, 44%, and 40% of patients treated with 24 mg, 72 mg, and 240 mg, respectively, of ligelizumab had complete control of symptoms (weekly urticaria activity score of 0), as compared with 26% of the patients in the omalizumab group and none of the patients in the placebo group (Fig. S2 in the Supplementary Appendix). At week 20, a response was observed in 39% of the patients who received the 72-mg dose of ligelizumab and in 40% of those who received the 240-mg dose, as compared with 31% of those in the omalizumab group and 5% of those in the placebo group. After the end of the treatment phase, the median time to loss of response in patients who had had a response

at week 20 was 3 weeks, 4 weeks, and 10.5 weeks among patients who received 24 mg, 72 mg, and 240 mg, respectively, of ligelizumab, as compared with 4 weeks among patients in the omalizumab group and 1 week among those in the placebo group.

The median (with interquartile ranges) and mean weekly hives-severity scores are shown in Figure 3. Patients in the groups that received the 72-mg or 240-mg doses of ligelizumab had median weekly hives-severity scores close to 0 from weeks 12 to 20. The mean changes from baseline to week 32 in the weekly hives-severity score, itch-severity score, and urticaria activity score are shown in Figure S3 in the Supplementary Appendix. The single 120-mg dose of ligelizumab resulted in suppression of symptoms that was similar to that seen with the 72-mg and 240-mg doses at week 4 and that lasted until week 8. At week 12, the mean changes from baseline in the angioedema activity score were -21.1 , -37.6 , and -27.3 among patients treated with 24 mg, 72 mg, and 240 mg, respectively, of ligelizumab, as compared with -23.1 in the omalizumab group and -23.6 in the placebo group (Fig. S3 in the Supplementary Appendix).

SAFETY

A higher proportion of adverse events that were considered by the investigators to be related to treatment was seen in the groups that received the 72-mg and 240-mg doses of ligelizumab than in the other groups (Table 2). These results were driven by mild or moderate injection-site reactions (in 4% of the patients in the 72-mg group and 7% of those in the 240-mg group) and by mild injection-site erythema (in 2% and 6%, respectively). All other adverse events that were considered to be possibly related to treatment during the trial were reported in three or fewer patients per group, with no meaningful differences among the trial groups. Serious adverse events were reported in 7%, 2%, and 2% of the patients treated with 24 mg, 72 mg, and 240 mg, respectively, of ligelizumab, as compared with 4% of the patients treated with omalizumab and 9% of those who received placebo.

The most frequently reported adverse events (occurring in $\geq 10\%$ of the patients overall) were viral upper respiratory tract infection (in 20%), upper respiratory tract infection (in 13%), and

Table 1. Demographic and Clinical Characteristics of Patients with Chronic Spontaneous Urticaria.*

Characteristic	24 mg (N=43)	Ligeluzumab, Every 4 Wk 72 mg (N=84)	Omalizumab, 300 mg Every 4 Wk (N=85)	Placebo (N=43)	Ligeluzumab, 120-mg Single Dose (N=42)	Total (N=382)
Age — yr	44.1±14.4	44.3±12.4	41.8±13.1	45.4±11.2	42.4±14.5	43.3±12.5
Female sex — no. (%)	31 (72)	61 (73)	66 (78)	31 (72)	30 (71)	286 (75)
Body-mass index†	27.3±6.3	28.5±7.1	28.1±6.4	27.4±6.5	27.5±6.9	27.9±6.5
Race — no. (%)‡						
Native American	0	1 (1)	0	0	0	1 (<1)
Asian	8 (19)	20 (24)	12 (14)	9 (21)	8 (19)	76 (20)
Black	0	2 (2)	4 (5)	0	2 (5)	8 (2)
White	32 (74)	57 (68)	67 (79)	31 (72)	31 (74)	283 (74)
Other	3 (7)	3 (3.6)	1 (1)	3 (7)	1 (2)	12 (3)
Time since diagnosis of chronic spontaneous urticaria — yr	3.4±4.0	3.9±5.4	5.1±7.5	3.6±3.5	5.7±7.5	4.3±6.0
IgE level — IU/ml						
Median	88.2	101.0	74.1	111.5	61.2	87.2
Range	0–5050.0	0–942.0	0–3480.0	2.2–870.0	0–889.0	0–14,100.0
Weekly itch-severity score§	12.5±3.4	13.6±4.1	12.7±4.4	13.6±4.1	13.6±4.0	13.1±4.1
Weekly hives-severity score¶	16.1±4.5	18.1±4.3	17.3±4.1	17.6±4.1	17.8±4.1	17.3±4.4
Weekly urticaria activity score¶¶	28.6±7.1	31.7±7.3	30.3±7.3	31.1±6.8	31.3±6.9	30.4±7.4
Presence of angioedema — no. (%)	17 (39.5)	39 (46)	44 (52)	25 (58)	18 (43)	182 (48)
Weekly angioedema activity score	28.0±16.3	42.2±25.0	32.8±28.1	39.6±24.9	32.2±26.3	34.7±24.9
Positive Chronic Urticaria Index — no. (%)***	16 (37)	32 (38)	33 (39)	14 (33)	15 (36)	145 (38)
Background medication						
Locally approved dose of H ₁ -antihistamine	19 (44)	35 (42)	37 (44)	19 (44)	18 (43)	164 (43)
Escalated dose of locally approved H ₁ -antihistamine	24 (56)	49 (58)	48 (56)	24 (56)	24 (57)	218 (57)

* Plus-minus values are means ±SD. There were no notable imbalances among the trial groups regarding the demographic and clinical characteristics of the patients at baseline.

† The body-mass index is the weight in kilograms divided by the square of the height in meters.

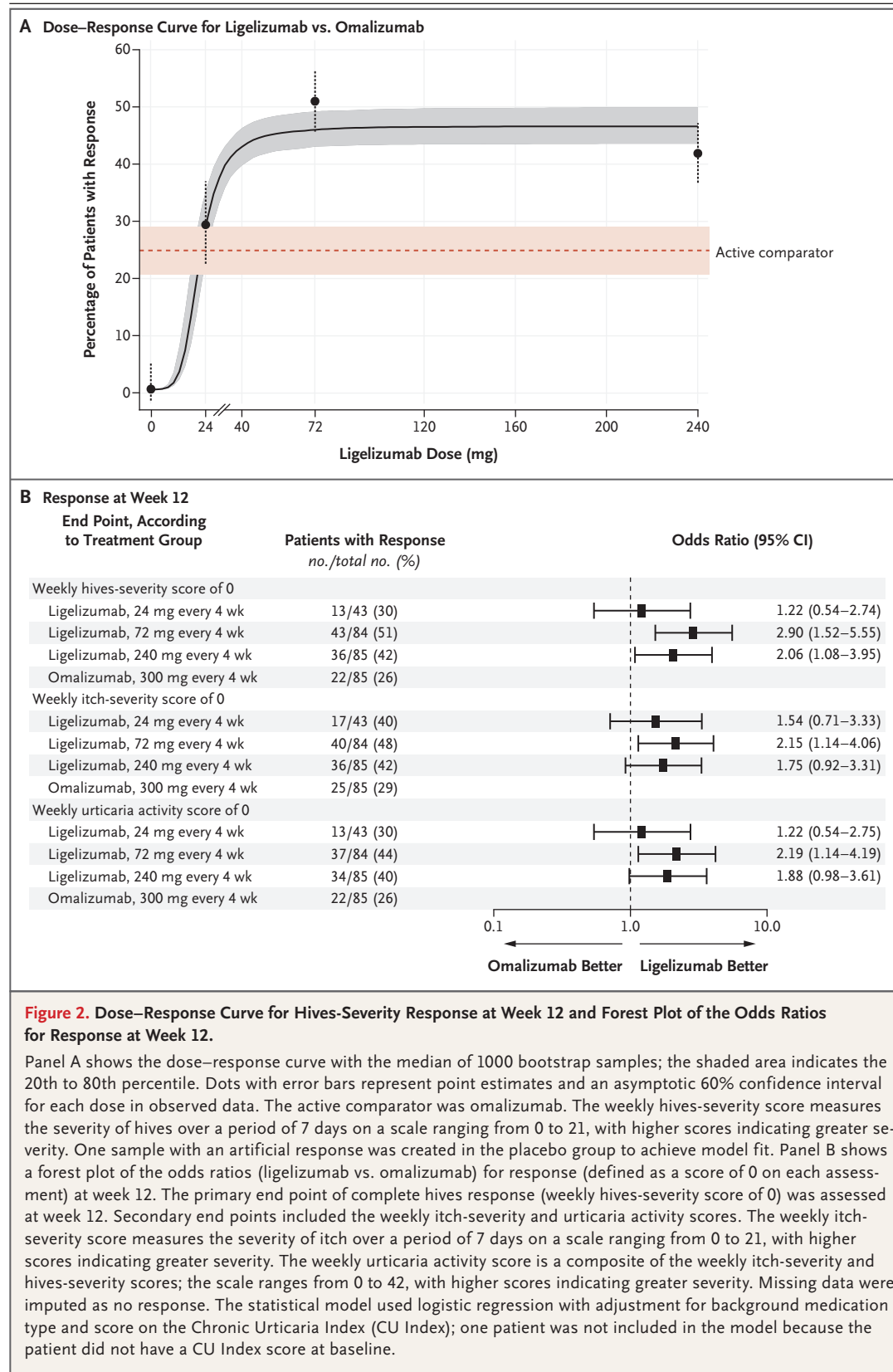
‡ Race was reported by the patient or determined by the investigator.

§ The weekly itch-severity and hives-severity scores measure the severity of itch and hives, respectively, over a period of 7 days on scales ranging from 0 to 21, with higher scores indicating greater severity.

¶ The weekly urticaria activity score is a composite of the weekly itch-severity and hives-severity scores. Scores range from 0 to 42, with higher scores indicating greater severity.

¶¶ Presence of angioedema at baseline for the previous week was determined according to both the Urticaria Patient Daily Diary and the weekly angioedema activity score, which measures angioedema activity over a period of 7 days on a scale ranging 0 to 105, with higher scores indicating worse angioedema.

*** A positive Chronic Urticaria (CU) Index (scores range from 1 to 50, with scores ≥10 representing a positive result) indicates that the patient has either an autoimmune basis for the urticaria or an alternative histamine-releasing factor that has been associated with greater disease severity than that in patients with a negative CU Index.



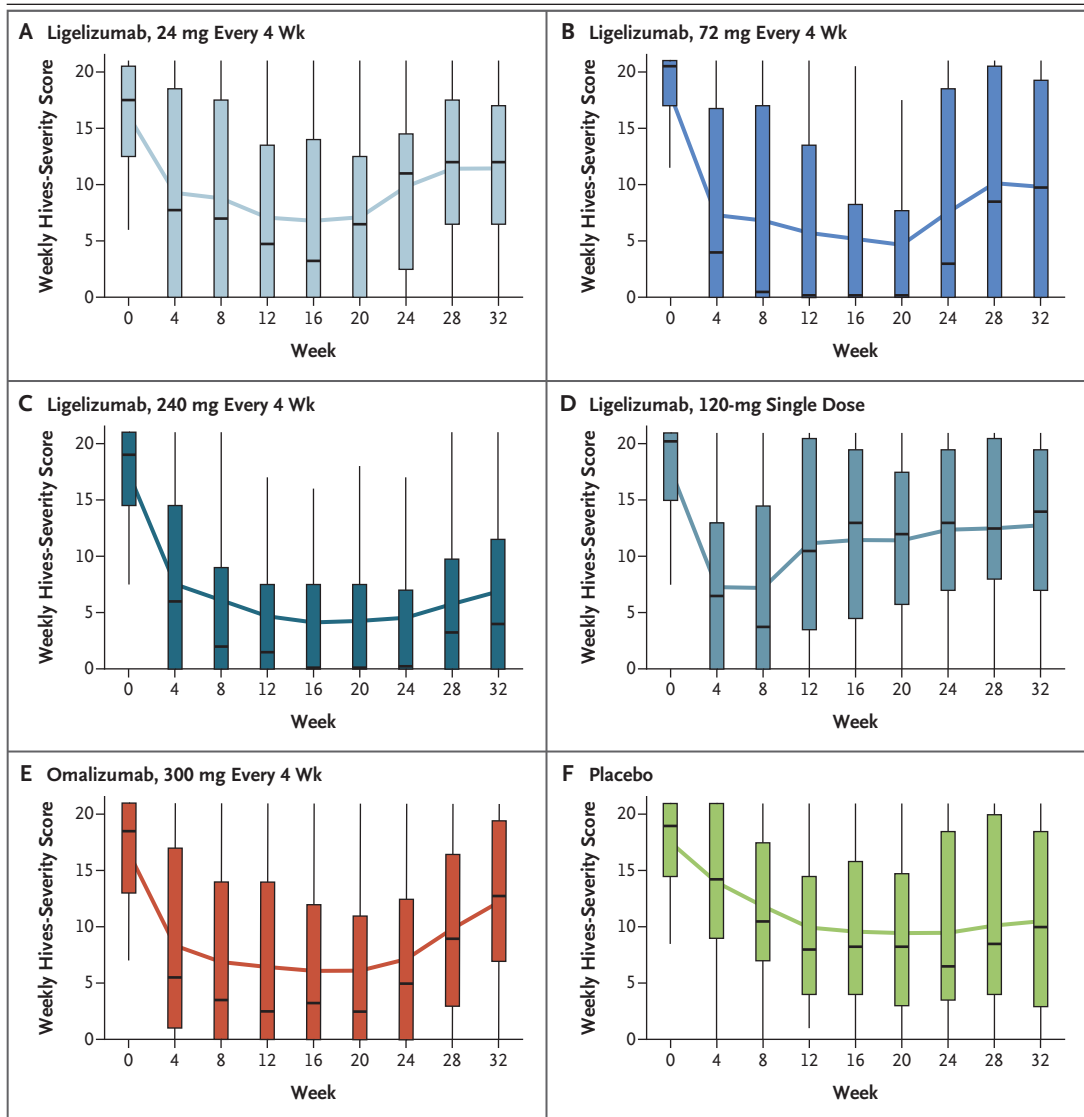


Figure 3. Hives-Severity Scores from Baseline to Week 32, According to Trial Group.

The horizontal lines within the boxes represent median values, the boxes represent interquartile ranges, and the vertical lines represent the data ranges. Colored lines represent mean values. The scale for the weekly hives-severity score ranges from 0 to 21, with higher scores indicating greater severity.

headache (in 11%). Viral upper respiratory tract infection was reported more frequently in the placebo group (30%) than in the four groups that received ligelizumab (15 to 24%) or in the omalizumab group (20%). No deaths or anaphylaxis events were reported in any of the trial groups.

DISCUSSION

Most currently available therapies for chronic spontaneous urticaria do not result in complete

control of symptoms in the majority of patients. The main objective of this trial was achieved, with a clear dose–response relationship being observed with ligelizumab in the achievement of complete hives response (weekly hives-severity score of 0) at week 12 (primary end point) in patients with chronic spontaneous urticaria. Our trial showed that 72 mg of ligelizumab, administered subcutaneously every 4 weeks, resulted in complete hives response in 51% of the patients, whereas 26% of the patients treated with omaliz-

Table 2. Adverse Events.

Variable	Ligelizumab, Every 4 Wk			Omalizumab, 300 mg Every 4 Wk (N = 85)	Placebo (N = 43)	Ligelizumab, 120-mg Single Dose (N = 42)	Total (N = 382)
	24 mg (N = 43)	72 mg (N = 84)	240 mg (N = 85)				
	<i>number of patients (percent)</i>						
At least one adverse event	36 (84)	63 (75)	63 (74)	62 (73)	34 (79)	37 (88)	295 (77)
Mild	16 (37)	31 (37)	32 (38)	36 (42)	15 (35)	22 (52)	152 (40)
Moderate	16 (37)	27 (32)	28 (33)	21 (25)	12 (28)	13 (31)	117 (31)
Severe	4 (9)	5 (6)	3 (4)	5 (6)	7 (16)	2 (5)	26 (7)
At least one serious adverse event	3 (7)	2 (2)	2 (2)	3 (4)	4 (9)	4 (10)	18 (5)
Adverse event leading to discontinuation	0	1 (1)	1 (1)	2 (2)	2 (5)	2 (5)	8 (2)
At least one adverse event possibly related to treatment	5 (12)	18 (21)	24 (28)	7 (8)	12 (28)	6 (14)	72 (19)
General disorder or administration-site condition	8 (19)	13 (15)	19 (22)	9 (11)	4 (9)	10 (24)	63 (16)
Injection-site reaction	0	3 (4)	6 (7)	0	1 (2)	0	10 (3)
Injection-site erythema	0	2 (2)	5 (6)	0	0	1 (2)	8 (2)
Most frequent adverse events*							
Viral upper respiratory tract infection	7 (16)	13 (15)	17 (20)	17 (20)	13 (30)	10 (24)	77 (20)
Upper respiratory tract infection	7 (16)	7 (8)	10 (12)	10 (12)	6 (14)	9 (21)	49 (13)
Headache	7 (16)	9 (11)	7 (8)	12 (14)	7 (16)	1 (2)	43 (11)

* The most frequent adverse events were defined as those that occurred in at least 10% of the patients in any group.

umab at a dose of 300 mg, which is the dose that has been considered to be effective in other studies,¹⁹⁻²¹ had this response.

The frequency of response according to the weekly urticaria activity score in the omalizumab group at week 12 was lower than in previous studies (34 to 44%).¹⁹⁻²¹ An explanation for this finding may be that a higher percentage of patients in our trial had a positive CU Index (indicating an autoimmune type IIb component to their chronic spontaneous urticaria),²⁸ had angioedema (Table 1), and had received treatment with elevated doses of H₁-antihistamines or combination therapies. Previous studies have shown that a positive CU Index was associated with increased severity of chronic urticaria and that significantly more patients with refractory chronic urticaria had a positive CU Index than those who had had a response to H₁-antihistamines.^{28,29}

In the current trial, responses according to changes from baseline in the weekly hives-severity score, itch-severity score, and urticaria activity score were observed as early as week 4 after receipt of the 72-mg, 120-mg, and 240-mg doses of ligelizumab, thus indicating an onset of action within this time frame for ligelizumab. The suppression of symptoms after the single 120-mg dose of ligelizumab was maintained until week 8, after which symptoms returned to the level observed with placebo. In contrast, a partial relapse of symptoms was noted with the 72-mg dose of ligelizumab toward the end of the 4-week administration interval. These data suggest that although the 72-mg dose of ligelizumab led to rapid control of symptoms, a dose higher than 72 mg could potentially provide enough drug effect throughout the administration interval to minimize relapse of symptoms and offer sustained control of symptoms throughout the 4-week administration interval. In support of this sustained treatment effect, the median time to loss of complete response in patients who had a weekly urticaria activity score of 0 at week 20 (end of the treatment phase) was greatest in the group that was treated with 240 mg of ligelizumab (10.5 weeks) and was similar in the groups that received 72 mg of ligelizumab or 300 mg of omalizumab (4 weeks). Previous stud-

ies of omalizumab have indicated that some patients have a return of symptoms within 4 weeks (i.e., before the administration of the next dose).^{32,33} Future studies will be needed to determine the dose of ligelizumab that results in a sustained clinical response throughout the administration interval.

The percentage of patients who had at least one adverse event was similar among the ligelizumab dose groups, the omalizumab group, and the placebo group. Adverse events were reported in 84%, 75%, and 74% of the patients who were treated with 24 mg, 72 mg, and 240 mg, respectively, of ligelizumab every 4 weeks and in 88% of the patients who received the single 120-mg dose of ligelizumab, 73% of those in the omalizumab group, and 79% of those in the placebo group. The higher incidence of injection-site reactions and injection-site erythema observed with ligelizumab at the doses of 72 mg and 240 mg than in the other groups may have been associated with differences in the administration volumes and compositions. The administration of the 240-mg dose of ligelizumab involved two injections of active drug (1.0 ml each), as compared with injections of one active drug (0.6 ml) and one placebo (0.6 ml) that were used for the 72-mg dose. No cases of anaphylaxis were reported in this trial.

In conclusion, in patients with moderate-to-severe chronic spontaneous urticaria, ligelizumab showed a clear dose-response relationship with regard to complete hives response (weekly hives-severity score of 0) at week 12. Ligelizumab resulted in rapid and sustained symptom control in patients with chronic spontaneous urticaria. In this small trial, side effects or laboratory abnormalities were not dose-limiting. Larger and longer trials are needed to establish the clinical efficacy of ligelizumab in patients with chronic spontaneous urticaria and its comparative profile with that of omalizumab.

A data sharing statement provided by the authors is available with the full text of this article at NEJM.org.

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APPENDIX

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