MG68

Observed Efficacy of Efgartigimod in Generalized Myasthenia Gravis Across Patient Subgroups in the ADAPT-SC+ Study

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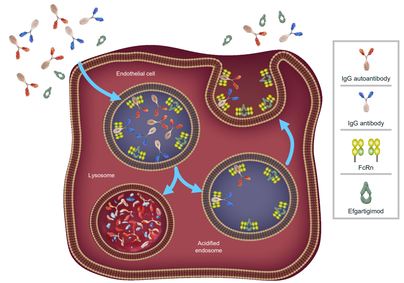
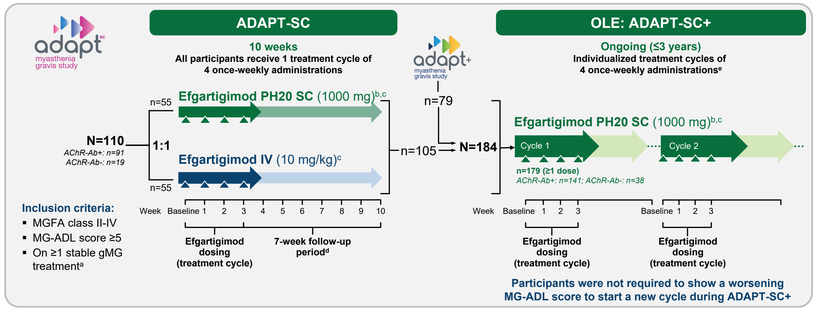
INTRODUCTION

Efgartigimod Mechanism of Action: Blocking FcRn



- FcRn recycles IgG antibodies and albumin. This recycling and salvage from lysosomal degradation results in IgG antibodies having the longest half-life and being the most abundant of all immunoglobulins[1-3]
- Efgartigimod is an IgG1 antibody Fc fragment that has been engineered for increased affinity to FcRn, and is uniquely composed of the only part of the IgG antibody that normally binds FcRn[1]
- Efgartigimod selectively reduces IgG by blocking FcRn-mediated IgG recycling without impacting antibody production, albumin levels, or other parts of the immune system[1,4,5]
- Efgartigimod PH20 SC is a coformulation of efgartigimod and recombinant human hyaluronidase PH20 (rHuPH20), which allows for rapid SC administration of larger volumes[6,7]
- PK/PD modeling and phase 3 data (ADAPT-SC) have demonstrated 4 once-weekly administrations of 1000 mg efgartigimod PH20 SC and 10 mg/kg efgartigimod IV result in comparable decreases in IgG levels[6,8]

METHODS



[a]AChEIs, steroids, and/or NSISTs. [b]Coformulated with 2000 U/mL rHuPH20. [c]Arrows indicate efgartigimod administration. [d]Participants could not receive treatment in the 7-week follow-up period. [e]Participants who are not in need of retreatment at study entry will instead start with an intertreatment period.

SUMMARY

- **Efgartigimod PH20 SC resulted in consistent improvements in AChR-Ab+ participants across subgroups, including those only receiving AChEIs**
- **Clinical improvements across subgroups were similar to those seen in participants in the ADAPT trial,[3] reinforcing efgartigimod's efficacy across a broad gMG population**
- **Results in participants with short disease duration and limited/no exposure to prior treatments suggest efgartigimod may be an effective option early in their disease course**
- **The majority of TEAEs were mild or moderate with no new safety findings**
- **The ADAPT-SC+ study is currently ongoing**

RESULTS

Table 1. Participant Demographics and Baseline Characteristics

Overall and AChR-Ab+ Population

	Efgartigimod PH20 SC *Overall* (n=179)	Efgartigimod PH20 SC *AChR-Ab+* (n=141)
Age, years, mean (SD)	50.7 (15.5)	51.0 (15.9)
Sex, female, n (%)	119 (66.5)	90 (63.8)
Weight, kg, median (Q1-Q3)	76.9 (64.0-89.8)	77.0 (63.0-92.0)
AChR-Ab+, n (%)	141 (78.8)	141 (100)
Total MG-ADL score, mean (SD)	7.9 (3.4)	7.6 (3.4)
Total MG-QoL15r score, mean (SD)	13.6 (6.9)	13.1 (6.8)
MG therapy during the first year, n (%)		
Any steroid	128 (71.5)	103 (73.0)
Any NSIST	89 (49.7)	67 (47.5)
Any AChEI	150 (83.8)	122 (86.5)
Steroid + NSIST	69 (38.5)	53 (37.6)
AChEI only	29 (16.2)	23 (16.3)

- 184 participants rolled over from ADAPT-SC (n=105) and ADAPT+ (n=79)
- 179 participants (141 AChR-Ab+ and 38 AChR-Ab-) received ≥1 dose of efgartigimod PH20 SC in ADAPT-SC+ through December 2022, with a mean (SD), median, and maximum study duration for all participants of 412.9 (104.52), 451, and 585 days, respectively

Table 2. Summary of TEAEs

Overall Population

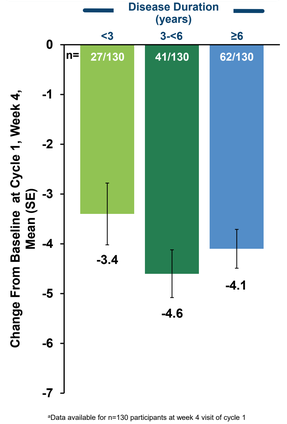
	Efgartigimod PH20 SC *(n=179; PYFU=193.4)*	
	ER[a]	n (%)
Any TEAE	9.0	152 (84.9)
Any TEAE grade ≥3	0.4	36 (20.1)
Any serious TEAE	0.3	33 (18.4)
Any injection site reaction	3.2	82 (45.8)
Fatal event[b]	<0.1	4 (2.2)
Discontinued study treatment owing to TEAEs[c]	<0.1	4 (2.2)
Most commonly observed TEAEs[d]		
Injection site erythema	1.7	52 (29.1)
COVID-19	0.2	40 (22.3)
Headache	0.6	36 (20.1)
Nasopharyngitis	0.2	28 (15.6)
Diarrhea	0.2	24 (13.4)
Injection site pain	0.2	21 (11.7)
Injection site pruritus	0.2	19 (10.6)
Injection site bruising	0.2	18 (10.1)

[a]Event rate was calculated as number of events per total PYFU. [b]Fatal events (metastatic renal cell cancer, cardiac arrest, pulmonary mass, and COVID-19/respiratory failure) were not related to efgartigimod PH20 SC treatment, as determined by investigators. [c]Treatment discontinuations were due to metastatic renal cell cancer (Cycle 1, death), cardiac arrest (Cycle 2, death), COVID-19/respiratory failure (Cycle 3, death), and MG crisis (Cycle 1). [d]Most frequent TEAEs occurring in >10% of participants receiving efgartigimod PH20 SC.

- Participants experiencing injection site reaction events decreased over subsequent cycles; from 34.6% (n=62/179) in Cycle 1 to 10.3% (n=7/68) in Cycle 9
- No injection site reactions were grade ≥3, serious, or resulted in treatment discontinuation

Figure 1. Week 4[a] MG-ADL Score Change from Baseline by Disease Duration

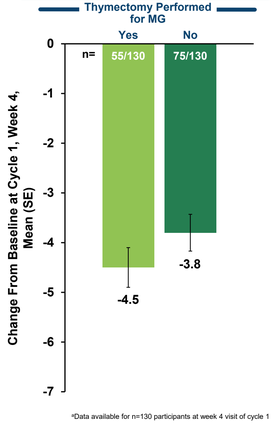
AChR-Ab+ Population



[a]Data available for n=130 participants at week 4 visit of cycle 1

Figure 2. Week 4[a] MG-ADL Score Change from Baseline by Thymectomy Status

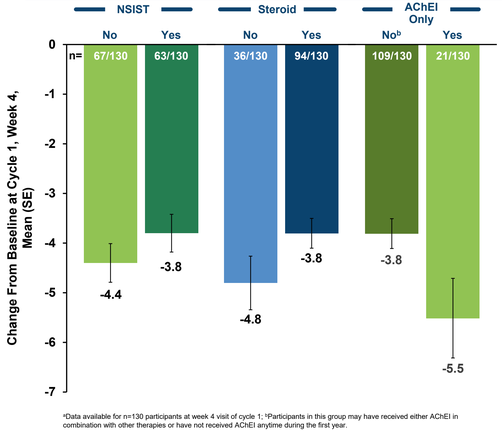
AChR-Ab+ Population



[a]Data available for n=130 participants at week 4 visit of cycle 1

Figure 3. Week 4[a] MG-ADL Score Change from Baseline by Concomitant Therapy

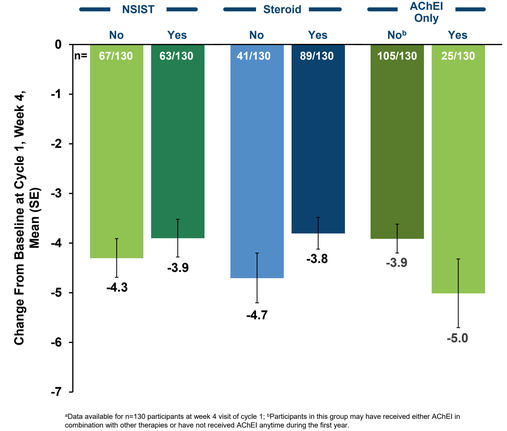
AChR-Ab+ Population



[a]Data available for n=130 participants at week 4 visit of cycle 1; [b]Participants in this group may have received either AChEI in combination with other therapies or have not received AChEI anytime during the first year.

Figure 4. Week 4[a] MG-ADL Score Change from Baseline by Prior Therapy

AChR-Ab+ Population



[a]Data available for n=130 participants at week 4 visit of cycle 1; [b]Participants in this group may have received either AChEI in combination with other therapies or have not received AChEI anytime during the first year.

ABBREVIATIONS

AChEI, acetylcholinesterase inhibitor; AChR-Ab, acetylcholine receptor antibody; ER, event rate; Fc, fragment crystallizable region; FcRn, neonatal Fc receptor; gMG, generalized myasthenia gravis; Ig, immunoglobulin; IV, intravenous; MG, myasthenia gravis; MG-ADL, Myasthenia Gravis Activities of Daily Living; MGFA, Myasthenia Gravis Foundation of America; MG-QoL15r, Myasthenia Gravis Quality of Life 15-Item Questionnaire, Revised; NSIST, nonsteroidal immunosuppressive therapy; OLE, open-label extension; PD, pharmacodynamic; PK, pharmacokinetic; PYFU, participant years of follow-up (sum of follow-up time of all participants expressed in years in the applicable period); rHuPH20, recombinant human hyaluronidase PH20; SC, subcutaneous; TEAE, treatment-emergent adverse event.

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SM: Alexion, argenx, UCB/Ra, Horizon. **TV:** Alexion, argenx, CSL Behring, Allergan/Abbvie, Alexion/AstraZeneca, Dianthus, Remegen, ImmunAbs, UCB, Amgen, Immunovant, Regeneron, Johnson & Johnson, Dianthus, Cartesians. **RB-S:** argenx and Sanofi. **EB, RK:** argenx. **KU:** UCB, Janssen, Horizon (Viela Bio), Chugai, Hanall BioPharma, Merck, Mitsubishi Tanabe, argenx, Alexion, Japan Blood Products Organization. **AM:** Alexion, argenx, Grifols, Hormosan, UCB, Janssen, Merck, Octapharma, German Myasthenia Gravis Society.
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