

Long-term outcomes from the phase II L-MIND study of tafasitamab (MOR208) plus lenalidomide in patients with relapsed or refractory diffuse large B-cell lymphoma

Johannes Duell,¹ Kami J. Maddocks,² Eva González-Barca,³ Wojciech Jurczak,⁴ Anna Marina Liberati,⁵ Sven de Vos,⁶ Zsolt Nagy,⁷ Aleš Obr,⁸ Gianluca Gaidano,⁹ Pau Abrisqueta,¹⁰ Nagesh Kalakonda,¹¹ Marc André,¹² Martin Dreyling,¹³ Tobias Menne,¹⁴ Olivier Tournilhac,¹⁵ Marinela Augustin,¹⁶ Andreas Rosenwald,¹⁷ Maren Dirnberger-Hertweck,¹⁸ Johannes Weirather,¹⁸ Sumeet Ambarkhane¹⁸ and Gilles Salles^{19*}

¹Medizinische Klinik und Poliklinik II, Universitätsklinik Würzburg, Würzburg, Germany; ²Department of Internal Medicine, Arthur G James Comprehensive Cancer Center, Ohio State University Wexner Medical Center, Columbus, OH, USA; ³Department of Hematology, Institut Catalá d'Oncologia (ICO), Hospital Duran i Reynals, Universitat de Barcelona, Barcelona, Spain; ⁴Maria Skłodowska-Curie National Research Institute of Oncology, Kraków, Poland; ⁵Università degli Studi di Perugia, Azienda Ospedaliera Santa Maria di Terni, Terni, Italy; ⁶Department of Medicine, Ronald Reagan UCLA Medical Center, Santa Monica, CA, USA; ⁷1st Department of Internal Medicine, Semmelweis University, Budapest, Hungary; ⁸Department of Hemato-Oncology, Palacký University and University Hospital, Olomouc, Czech Republic; ⁹Division of Hematology, Department of Translational Medicine, University of Eastern Piedmont, Novara, Italy; ¹⁰Department of Hematology, Vall d'Hebron Institute of Oncology (VHIO), Vall d'Hebron University Hospital, Barcelona, Spain; ¹¹Molecular and Clinical Cancer Medicine, University of Liverpool and The Clatterbridge Cancer Centre, Liverpool, UK; ¹²Department of Haematology, Université Catholique de Louvain, CHU UCL Namur, Yvoir, Belgium; ¹³Department of Medicine III, LMU University Hospital, Munich, Germany; ¹⁴Department of Haematology, Freeman Hospital, Newcastle Upon Tyne Hospitals NHS Foundation Trust, Newcastle Upon Tyne, UK; ¹⁵Service d'Hématologie Clinique et de Thérapie Cellulaire, CHU Estaing, Clermont-Ferrand, France; ¹⁶Department of Hematology and Oncology, Paracelsus Medical University, Klinikum Nürnberg, Nürnberg, Germany; ¹⁷Institute of Pathology, University of Würzburg, Würzburg, Germany; ¹⁸MorphoSys AG, Planegg, Germany and ¹⁹Hématologie, Hospices Civils de Lyon and Université de Lyon, Lyon, France.

*Current address: Lymphoma Service, Memorial Sloan Kettering Cancer Center, New York, NY, USA.

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An error in the original manuscript (2020.275958) is present in the paper that appeared as Early view on July 1, 2021.

In Table 2, 'Efficacy outcomes in primary and follow-up analysis' The complete response for 'Last therapy refractory' should be 14 (40.0) and not 14 (14.0).

Correction

The corrected datum, a complete response rate of 40.0% in last-therapy-refractory patients receiving tafasitamab plus lenalidomide for relapsed/refractory diffuse large B-cell lymphoma, is shown in the Table below.

Table 2. Efficacy outcomes in the primary and follow-up analyses.

	Tafasitamab plus lenalidomide (N=80) [†]		Clinically relevant subgroups (follow-up analysis)		
	Primary analysis (data cut-off: Nov 30, 2018) [§]	Follow-up analysis (data cut-off: Oct 30, 2020)	Primary refractory disease (n=15)	Rituximab-refractory disease (n=33)	Last-therapy- refractory (n=35)
Best objective response, n (%)					
Complete response	34 (42.5)	32 (40.0)	5 (33.3)	13 (39.4)	14 (40.0)
Partial response	14 (17.5)	14 (17.5)	3 (20.0)	5 (15.2)	7 (20.0)
Stable disease	11 (13.8)	13 (16.3)	2 (13.3)	4 (12.1)	3 (8.6)
Progressive disease	13 (16.3)	13 (16.3)	3 (20.0)	7 (21.2)	7 (20.0)
Not evaluable*	8 (10.0)	8 (10.0)	2 (13.3)	4 (12.1)	4 (11.4)
ORR (CR + PR), n (%) [95% CI] [†]	48 (60.0) [48.4-70.9]	46 (57.5) [45.9-68.5]	8 (53.3) [26.6-78.7]	18 (54.5) [36.4-71.9]	21 (60.0) [42.1-76.1]
Median DoR (IRC), months (95% CI)	21.7 (21.7-NR)	43.9 (26.1-NR)	NR (1.8-NR)	NR (5.8-NR)	NR (5.8-NR)
Median PFS (IRC), months (95% CI)	12.1 (5.7-NR)	11.6 (6.3-45.7)	5.3 (0.9-NR)	7.6 (2.7-NR)	7.6 (2.7-NR)
Median OS, months (95% CI)	NR (18.3-NR)	33.5 (18.3-NR)	13.8 (1.3-NR)	15.5 (8.6-NR)	15.5 (8.6-NR)

*Non-evaluable patients had no valid post-baseline response assessments. [†]Using the two-sided 95% Clopper-Pearson exact method based on a binomial distribution. [‡]One patient received tafasitamab only. ORR: objective response rate; CR: complete response; PR: partial response; 95% CI: 95% confidence interval; DoR: duration of response; IRC: independent review committee; PFS: progression-free survival; OS: overall survival; NR: not reached.