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EDITED AND REVIEWED BY Sophia George, University of Miami, United States

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RECEIVED 31 May 2023 ACCEPTED 05 July 2023 PUBLISHED 07 August 2023

CITATION

Lorusso D, Danesi R, Locati LD, Masi G, De Giorgi U, Gadducci A, Pignata S, Sabbatini R, Savarese A, Valabrega G, Zamagni C and Colombo N (2023) Corrigendum: Optimizing the use of lenvatinib in combination with pembrolizumab in patients with advanced endometrial carcinoma. *Front. Oncol.* 13:1232476. doi: 10.3389/fonc.2023.1232476

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Corrigendum: Optimizing the use of lenvatinib in combination with pembrolizumab in patients with advanced endometrial carcinoma

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KEYWORDS

lenvatinib, pembrolizumab, endometrial cancer, tyrosine kinase inhibitor, immune response

A corrigendum on

Optimizing the use of lenvatinib in combination with pembrolizumab in patients with advanced endometrial carcinoma

by Lorusso D, Danesi R, Locati LD, Masi G, De Giorgi U, Gadducci A, Pignata S, Sabbatini R, Savarese A, Valabrega G, Zamagni C and Colombo N (2022) *Front. Oncol.* 12:979519. doi: 10.3389/fonc.2022.979519

In the published article, there was an error in Figure 1 as published. By mistake, we added the reference 18 for this figure. The corrected Figure 1 and its caption appear below.

In the published article, an author name was incorrectly written as Sabbatini Roberto. The correct name is Roberto Sabbatini.

The authors apologize for these errors and state that these do not change the scientific conclusions of the article in any way. The original article has been updated.

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Adverse Reaction	/	in contraction		in ooo	Control Ogo De Marine	mu 0000 - 000	Participant of the second	0 0 0 0 0 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22 23 24 25 26 27 28 29 30 50 50 50 50 50 50 50 50 50 5	aximu
Hypertension	61	× 65%			0%		0%	/0 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22 23 24 25 26 27 28 29 30 S ^m MiN: 0.1 / Q1: 1.4 / Median: 2.1 / Q3: 5.0 / MAX: 30.1 weeks	30.1
Musculoskeletal pain	61	65%	6%	6%	0%	2%	0%	MIN: 0.3 / Q1: 1.3 / Median: 2.4 / Q3: 9.1 / MAX: 31.3 weeks	31.3
Proteinuria	18	19%	4%	3%	0%	0%	0%	MIN: 1.1 / Q1: 2.7 / Median: 3.2 / Q3: 12.1 / MAX: 38.3 weeks	38.3
Fatigue	61	65%	16%	24%	1%	14%	0%	MIN: 0.1 / Q1: 0.9 / Median: 3.3 / Q3: 8.7 / MAX: 118.4 weeks	118.4
	45	48%	7%	9%	0%	3%	0%	MIN: 0.1 / Q1: 1.0 / Median: 4.7 / Q3: 11.4 / MAX: 143.1 weeks	143.1
Diarrhea	60	64%	14%	10%	1%	6%	0%	MIN: 0.1 / Q1: 1.0 / Median: 4.8 / Q3: 16.4 / MAX: 55.0 weeks	55.0
Decreased appetite	49	52%	5%	9%	0%	6%	0%	5.1 MIN: 0.1 / Q1: 2.0 / Median: 5.1 / Q3: 11.6 / MAX: 37.4 weeks	37.4
Stomatitis	40	43%	4%	5%	0%	1%	0%	5.5 MIN: 0.6 / Q1: 1.9 / Median: 5.5 / Q3: 13.3 / MAX: 29.1 weeks	29.1
Vomiting	37	39%	11%	6%	0%	4%	0%	MIN: 0.4 / Q1: 3.0 / Median: 5.9 / Q3: 11.7 / MAX: 96.6 weeks	96.6
Hypothyroidism	48	51%	2%	0%	0%	1%	0%	6.1 MIN: 1.0 / Q1: 5.9 / Median: 6.1 / Q3: 15.3 / MAX: 43.1 weeks	43.1
PPES	24	26%	5%	13%	0%	1%	0%	MIN: 1.1 / Q1: 4.2 / Median: 8.1 / Q3: 13.6 / MAX: 70.9 weeks	70.9
Weight decreased	34	36%	3%	3%	0%	4%	0%	9.1 MIN: 2.1 / Q1: 7.1 / Median: 9.1 / Q3: 15.1 / MAX: 124.3 weeks	124.3

Post hoc analysis of time to first onset of selected adverse reactions. Reproduced with permission from [19].