

## Erratum to: Safety of long-term denosumab therapy: results from the open label extension phase of two phase 3 studies in patients with metastatic breast and prostate cancer

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Table 3. The adverse event of ‘Anemia’ in the Denosumab/Denosumab column under the Breast Cancer Study is 53 (16.7 %).

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**Table 3** Adverse events during the open-label treatment phase

Event, <i>n</i> (%)	Breast cancer study		Prostate cancer study	
	Denosumab/Denosumab ( <i>N</i> =318) <sup>a</sup>	Zoledronic acid/Denosumab ( <i>N</i> =334) <sup>a</sup>	Denosumab/Denosumab ( <i>N</i> =147) <sup>a</sup>	Zoledronic acid/Denosumab ( <i>N</i> =118) <sup>a</sup>
All adverse events	283 (89.0)	303 (90.7)	138 (93.9)	105 (89.0)
Serious adverse events	126 (39.6)	133 (39.8)	78 (53.1)	63 (53.4)
Most common adverse events				
Nausea	72 (22.6)	77 (23.1)	20 (13.6)	16 (13.6)
Anemia	53 (16.7)	50 (15.0)	34 (23.1)	26 (22.0)
Fatigue	70 (22.0)	74 (22.2)	23 (15.6)	15 (12.7)
Back pain	66 (20.8)	56 (16.8)	29 (19.7)	19 (16.1)
Asthenia	40 (12.6)	48 (14.4)	29 (19.7)	11 (9.3)
Arthralgia	57 (17.9)	61 (18.3)	25 (17.0)	17 (14.4)
Adverse events of infection <sup>b</sup>	135 (42.5)	135 (40.4)	58 (39.5)	33 (28.0)
Osteonecrosis of the jaw (ONJ) <sup>c,d</sup>	20 (6.3)	18 (5.4)	12 (8.2)	7 (5.9)
CTCAE, v 3 grade 3	2 (0.6)	6 (1.8)	3 (2.0)	1 (0.8)
CTCAE, v 3 grade 4	0 (0.0)	1 (0.3)	0 (0.0)	1 (0.8)
Adverse events of new primary malignancy <sup>e</sup>	2 (0.6) <sup>f</sup>	1 (0.3) <sup>g</sup>	1 (0.7) <sup>h</sup>	0 (0.0)
Adverse events of hypocalcemia <sup>i</sup>	12 (3.8)	9 (2.7)	8 (5.4)	5 (4.2)
Serious	3 (0.9)	0 (0.0)	1 (0.7)	1 (0.8)

CTCAE common terminology criteria for adverse events; version 3

<sup>a</sup> Number of patients who received at least one dose of open-label denosumab

<sup>b</sup> System organ class: Infections and Infestations, Medical Dictionary for Regulatory Activities version 14.0 (breast cancer study), 14.1 (prostate cancer study)

<sup>c</sup> Positively adjudicated by blinded committee of experts. There were no grade 5 ONJ events

<sup>d</sup> Rates not adjusted for patient-years of investigational product exposure and patient follow-up

<sup>e</sup> No events were considered by the investigator to be related to denosumab treatment

<sup>f</sup> Renal cell cancer, squamous cell carcinoma of the skin

<sup>g</sup> Germ cell cancer

<sup>h</sup> Bladder cancer

<sup>i</sup> Includes the preferred terms hypocalcemia and blood calcium decreased