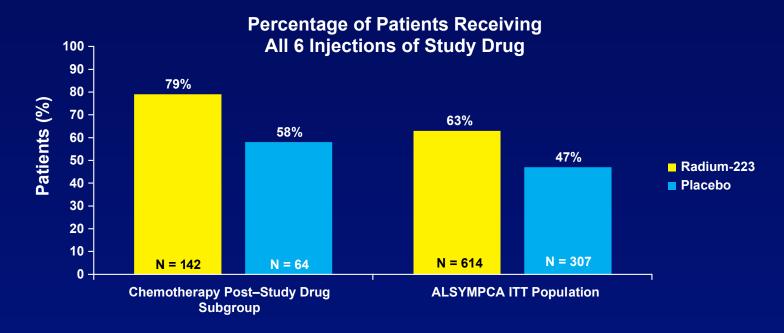
3-Year Follow-up of Chemotherapy Following Radium-223 Dichloride in Castration-Resistant Prostate Cancer Patients With Symptomatic Bone Metastases From ALSYMPCA

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### **Patients**

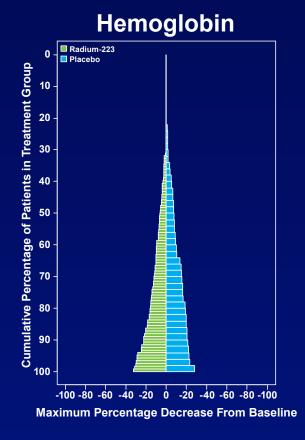
- Of 921 ALSYMPCA patients, 206 (22%) received chemotherapy post–study drug (radium-223: n = 142; placebo: n = 64)
- Docetaxel was the most common chemotherapy;
   70% post-radium-223 and 72% post-placebo
- Compared with the ALSYMPCA ITT population, chemotherapy post-study drug patients were younger, had better ECOG PS, fewer metastases, and lower tALP and PSA

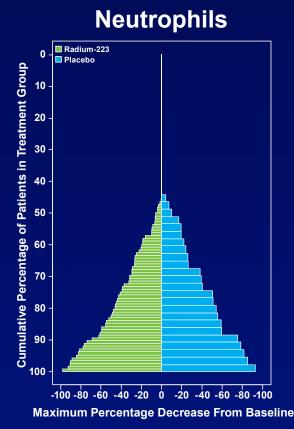


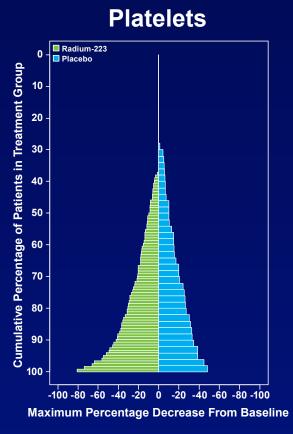
## **Safety**

 Maximum percentage decrease from baseline in hemoglobin and neutrophils was similar across treatment arms, but trended greater for platelets in radium-223 arm

#### **Maximum Percentage Decrease from Baseline**







# **Safety**

- Hematologic parameters were constant from last measurement pre-chemotherapy to 18 months post-chemotherapy in both treatment arms; no apparent impact of prior docetaxel
- Percentages of patients with hematologic values corresponding to grade 3 or 4 adverse events were generally low, trending higher among patients in the radium-223 than the placebo arm

Percentage of Patients in Chemotherapy Post–Study Drug Subgroup With Hematologic Laboratory Values Corresponding to Grade 3 or 4 Adverse Events

|  | Radium-223<br>N = 142 |         | Placebo<br>N = 64 |       | <i>P</i> Value |
|--|-----------------------|---------|-------------------|-------|----------------|
|  | n                     | n (%)   | n                 | n (%) |                |
| Hemoglobin, < 8.0-6.5 g/dL or < 6.5 g/dL                             | 116                   | 9 (8)   | 49                | 2 (4) | 0.509          |
| Neutrophils, < 1-0.5 x $10^9$ /L or < 0.5 x $10^9$ /L                | 114                   | 11 (10) | 48                | 1 (2) | 0.112          |
| Platelets, < 50-25 x 10 <sup>9</sup> /L or < 25 x 10 <sup>9</sup> /L | 116                   | 7 (6)   | 49                | 1 (2) | 0.438          |

### **Overall Survival**

- Similar proportions of radium-223 (41/142 [29%]) and placebo (21/64 [33%]) patients died during and within 30 days of completing chemotherapy
- Median OS from start of chemotherapy was 16.0 months postradium-223 and 15.8 months post-placebo (HR = 1.23; 95% CI, 0.86-1.75)

### **Conclusions**

- Safety of chemotherapy post–radium-223 assessed in post hoc analysis showed no detrimental effects on hematology or OS
- Findings indicate that chemotherapy can be administered safely after radium-223