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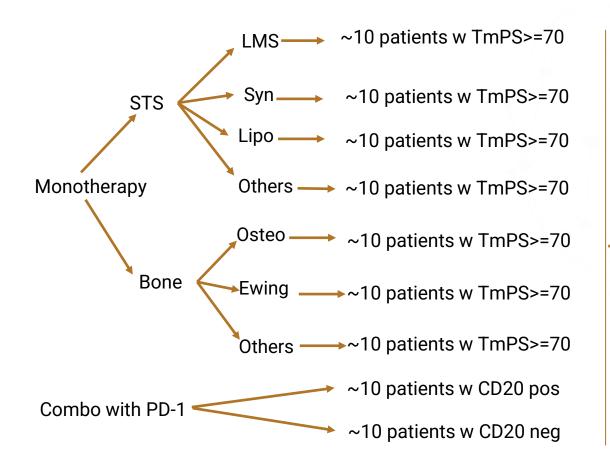
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## Sarcoma Phase 2 Part 1 Design: Predefined Interim Analysis Decision Tree



PHASE 2 - SARCOMA

Mecbotamab vedotin – BA3011



For each Subgroup up to 10 patients

No Go
if 0 CR/PR and
PFR12 week < 40%

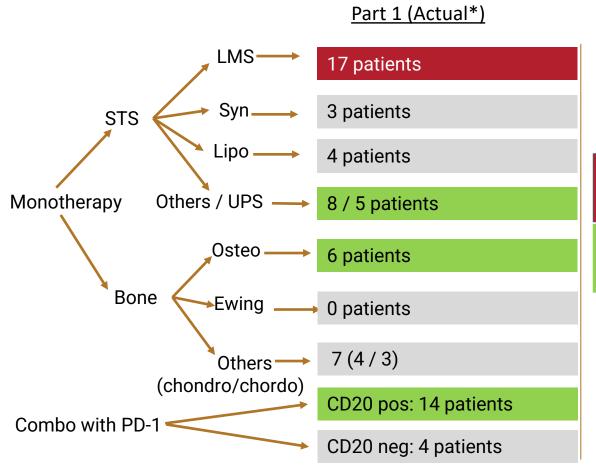
Go
if ≥ 1 CR/PR or
PFR12 week ≥ 40%

## Sarcoma: Phase 2 Part 1 Topline Interim Analysis Results



PHASE 2 - SARCOMA

Mecbotamab vedotin – BA3011



#### Predefined Criteria

For each Subgroup up to 10 patients

No Go if 0 CR/PR **and** PFS 3months < 40%

 $\frac{Go}{if ≥ 1 CR/PR or}$ PFS 3months ≥ 40%

## <u>Interim Results</u>

- LMS not moving forward
  - PFS = 27% (pending confirmation)
- UPS moving forward
  - $\circ$  PR = 2/6
- Osteosarcoma moving forward
  - o PFS = 67%
- Combo w PD-1 moving forward
  - $\circ$  PR = 1 / 1 (UPS)\*
  - PFS (combo overall) = 51%
- All other cohorts (in grey) continue enrollment

<sup>\*</sup>Included in UPS cohort

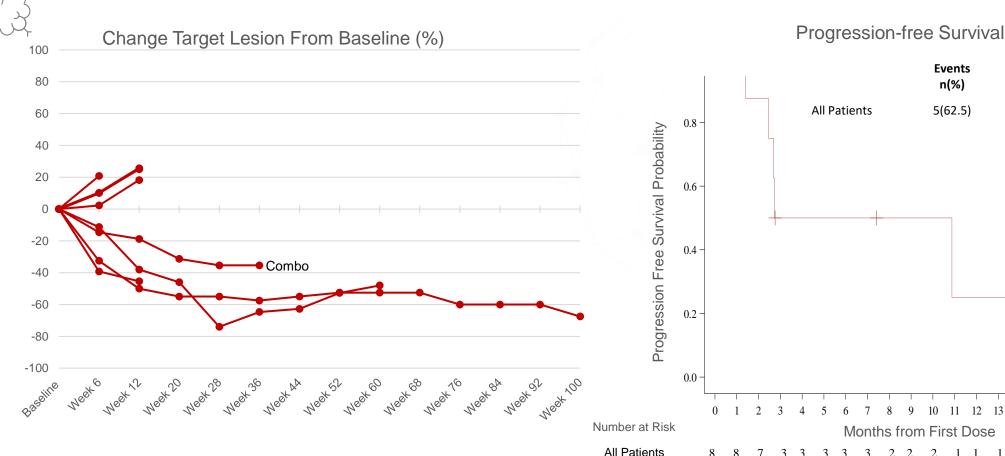
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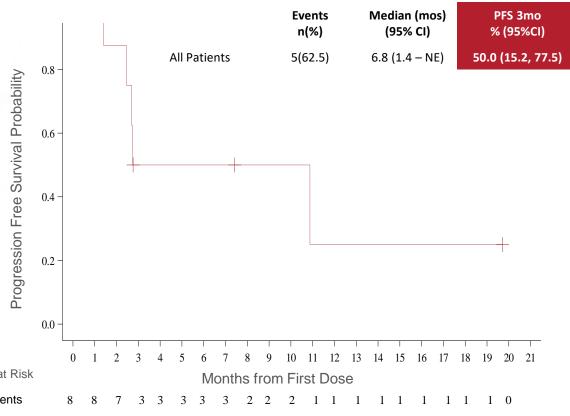
## Undifferentiated Pleomorphic Sarcoma (UPS): Phase 1 & 2 (1.8mg/kg; n=8) Change in Target Lesion and Progression Free Survival











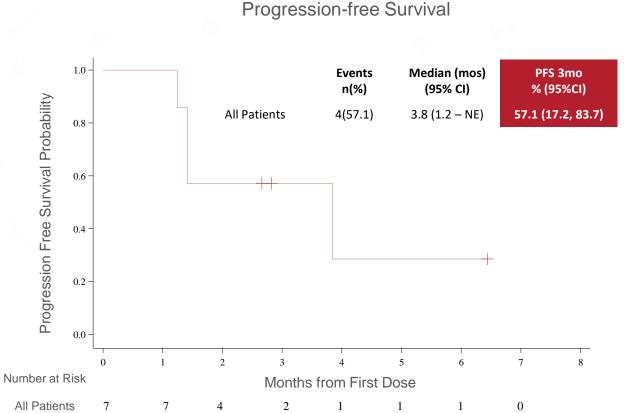
# Osteosarcoma: Phase 1 & 2 (1.8mg/kg; n=7) Change in Target Lesion and Progression Free Survival



PHASE 1 / 2 - Osteosarcoma







Recent independent phase 2 study demonstrated placebo PFS rate at 2 months for 1st and 2nd line metastatic osteosarcoma of ~0% (www.thelancet.com/oncology Vol 20 January 2019)

## Stage IV Multi-Refractory Melanoma: Phase 1 & 2 Complete Response Observed in Two of Two ROR2+ Patients



Phase 1 / 2 - MELANOMA

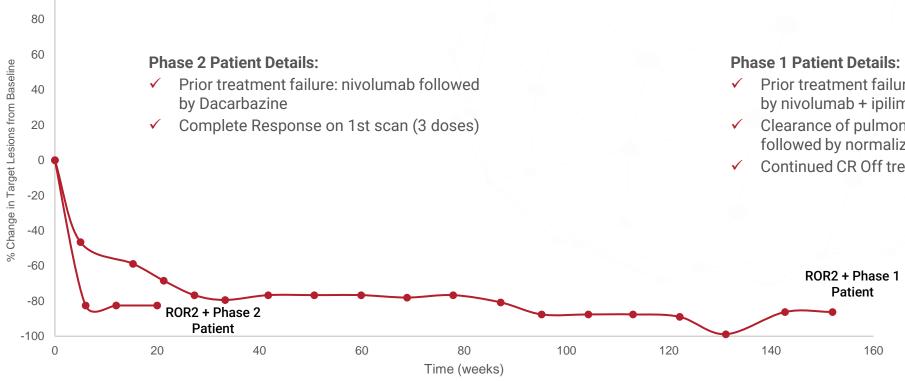


Ozuriftamab vedotin - BA3021



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- Prior treatment failure: nivolumab followed by nivolumab + ipilimumab combination
- Clearance of pulmonary metastases followed by normalization of adenopathy
- Continued CR Off treatment for over 2 yrs





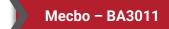
## Continued Promising Safety and Tolerability Profile in Phase 2 at the RP2D



#### PHASE 2 AEs

#### Overview

- AEs consistent with MMAE-based toxicity, including:
  - o Reversible myelosuppression
  - Transient liver enzyme elevation
  - Metabolic disturbances
- Few related SAEs
- Few related AEs leading to treatment discontinuation



CAB AXL-ADC Dosing 1.8mg/kg Q2W (safety population Phase 2)

Characteristic	BA3011 (N=68)
Any Adverse Events (AEs)	61 ( 90%)
Related AEs with CTCAE <sup>1</sup> Grade 3 or 4 <sup>2</sup>	16 ( 23%)
Any related serious AEs <sup>2</sup>	6 ( 9%)
Related AEs leading to death <sup>2</sup>	0
Related AEs leading to treatment discontinuation <sup>2</sup>	2 (3%)§

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Constipation	Grade 1-2 (18%)		
	Grade 3-4 (0%)		
Peripheral Neuropathy	All Grade 1-2 (23%)		
Diarrhea	Grade 1-2 (19%)		
	Grade 3-4 (0%)		

Constipation is believed to be an on-target mediated effect

- · No clinically meaningful on-target toxicity observed over background
- Differentiated profile due to advantageous pharmacokinetic characteristics of CAB ADCs

# Significant Unmet Need and Commercial Opportunity Across Multiple Solid Tumors Types



#### Sarcoma

- ~17K newly diagnosed metastatic cases / yr (US)
- 5-year survival rate (~20% for advanced metastatic disease)
- UPS is one of the largest and most aggressive sarcoma subtypes
- Osteosarcoma represents nearly 1/3 of all bone sarcoma cases – aggressive tumors – high recurrence – with limited treatment options

### **NSCLC**

- ~200K newly diagnosed NSCLC patients / yr (US)
- 5-year survival rate (~25% for advanced metastatic disease)
- While relatively crowded market, there is a lack of efficacious therapies for refractory patients

## Melanoma

- ~200K newly diagnosed cases / yr (US) – 50% invasive
- 5-year survival rate (~25% for advanced metastatic disease)
- ~1.3mm people in the US living with melanoma

## Head & Neck

- ~66K newly diagnosed cases / yr (US)
- 5-year survival rate (~40% for recurrent or advanced metastatic disease)
- Deficit of 2<sup>nd</sup> line treatments