

**IMpower-131**

# IMpower-131

(PD1/PDL1-未)

Co-Primary Endpoint: OS(ITT), PFS(investigator)

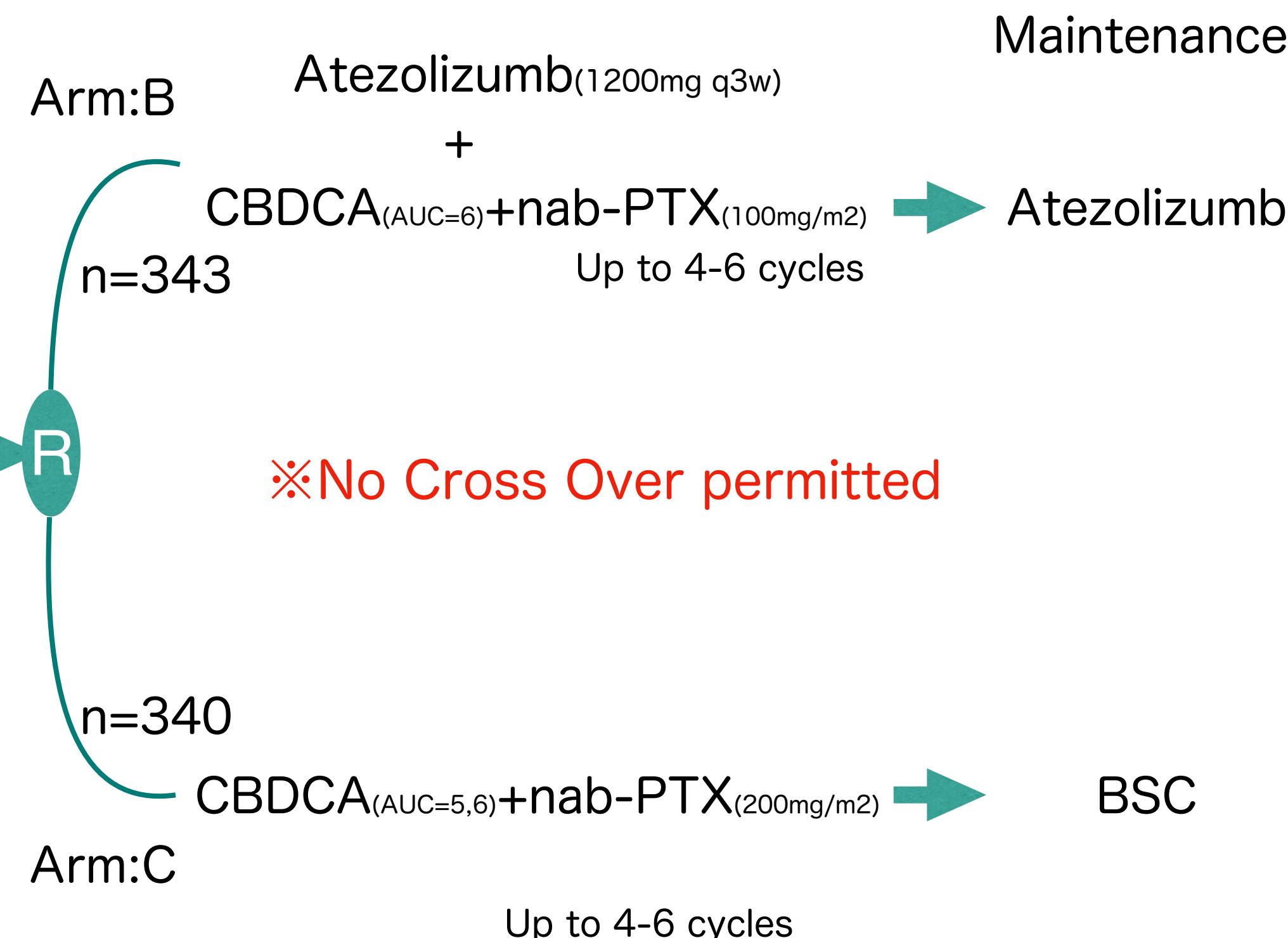
Secondary Endpoint: PFS and OS in PD-L1 subgroups

Primary

Subgroups

Sq

化学療法未治療の非小細胞  
肺癌患者（1次治療）  
・PD-L1 発現 Any



investigator  
PFS

OS

PD-L1 High    PD-L1 Low    PD-L1 Negative  
TC3 or IC3    TC1/2 or IC1/2    TC0 or IC0

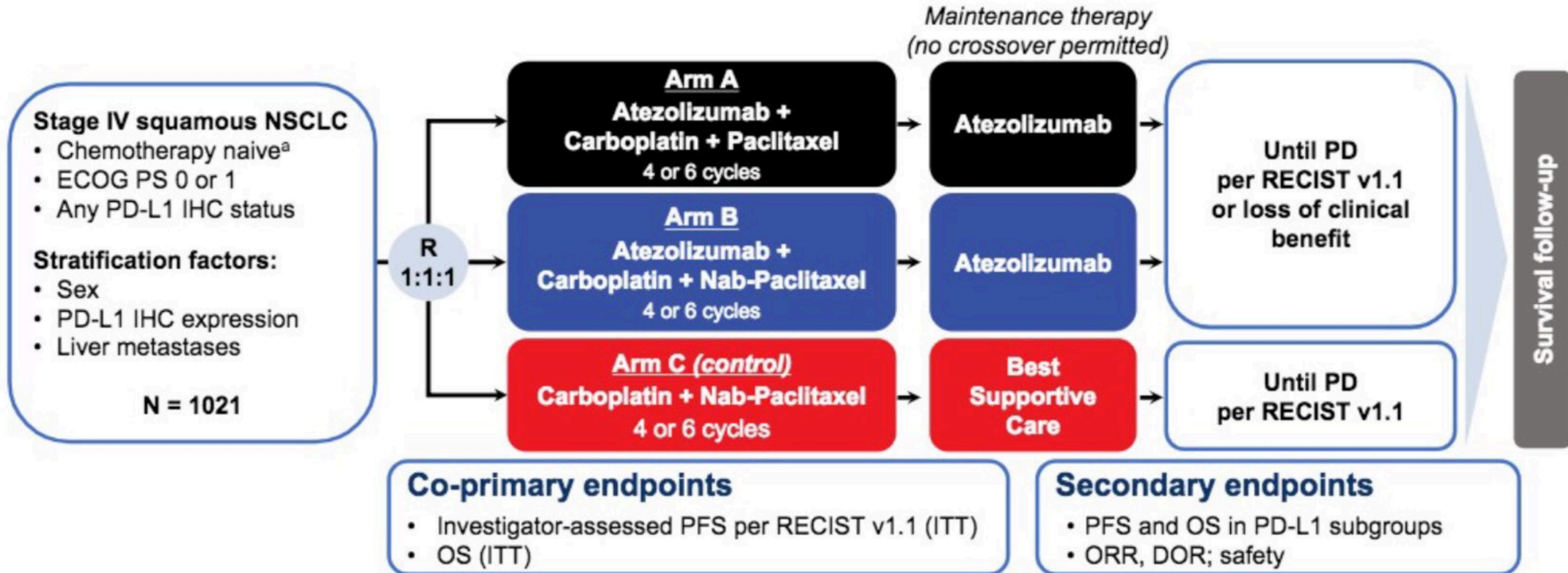
6.3ヶ月    14.0ヶ月    23.6ヶ月    12.4ヶ月    13.8ヶ月

HR=0.715    HR=0.96  
p=0.0001    p=0.6931    HR=0.56    HR=1.34    HR=0.86

5.6ヶ月    13.9ヶ月    14.1ヶ月    16.6ヶ月    12.5ヶ月

2019/01/14

# IMpower131: Study Design

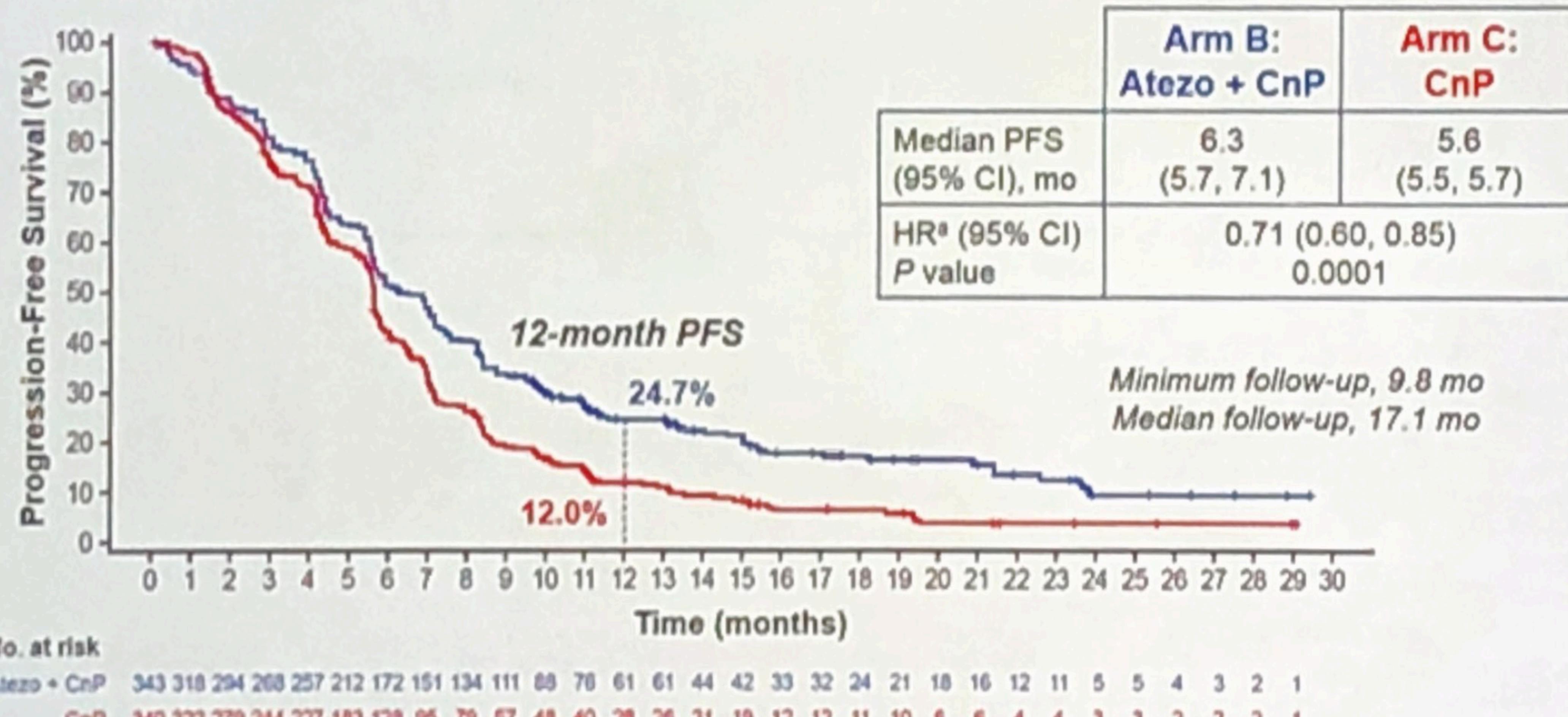


Atezolizumab 1200 mg IV q3w; carboplatin AUC 6 IV q3w; nab-paclitaxel 100 mg/m<sup>2</sup> IV qw; paclitaxel 200 mg/m<sup>2</sup> IV q3w.

<sup>a</sup> Patients with a sensitising EGFR mutation or ALK translocation must have disease progression or intolerance to treatment with ≥ 1 approved targeted therapies. Testing for EGFR mutation or ALK translocation was not mandatory.

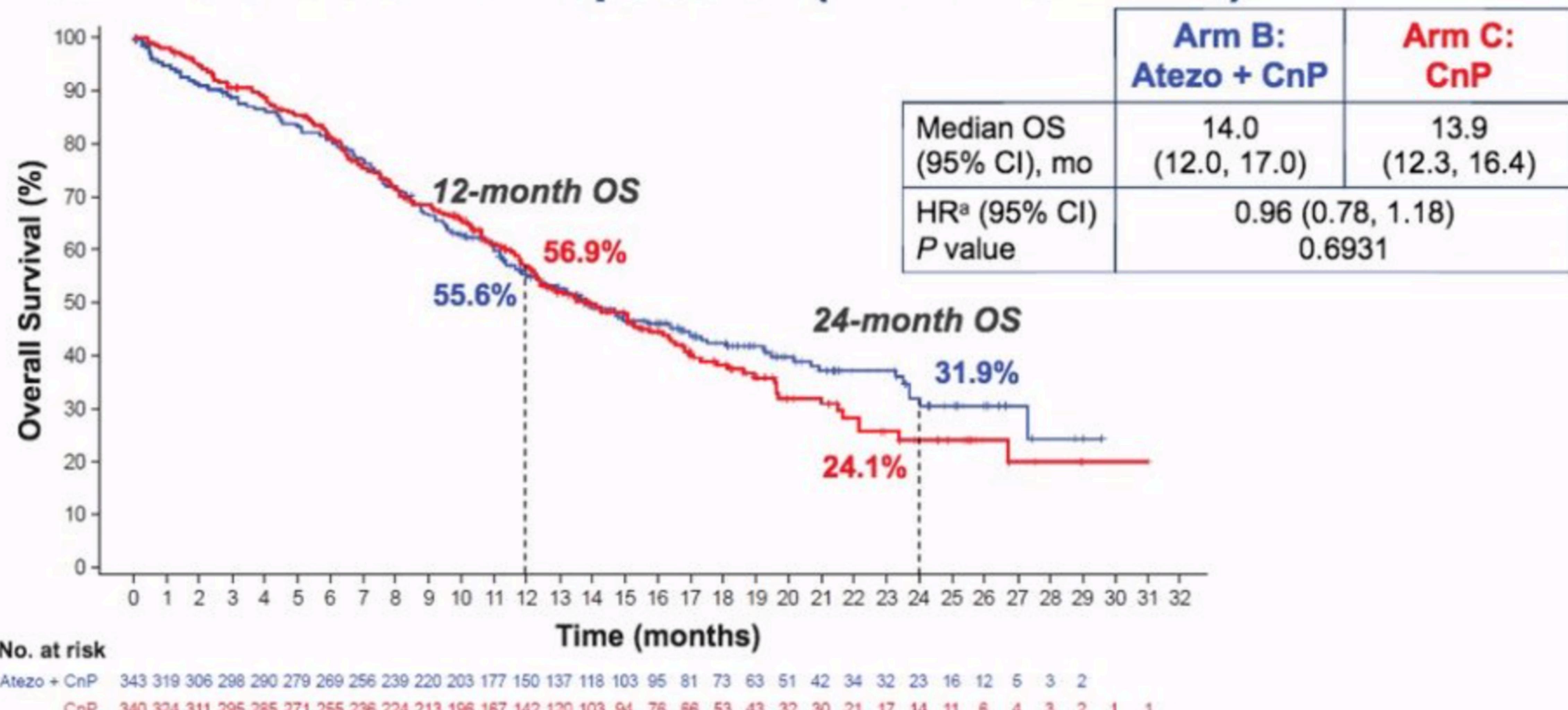
PD-L1 expression was evaluated using the VENTANA SP142 IHC assay.

# INV-Assessed PFS in the ITT Population (Arm B vs Arm C)



Data cutoff: January 22, 2018.  
INV, investigator; \*Stratified HR.

# First Interim OS in the ITT Population (Arm B vs Arm C)

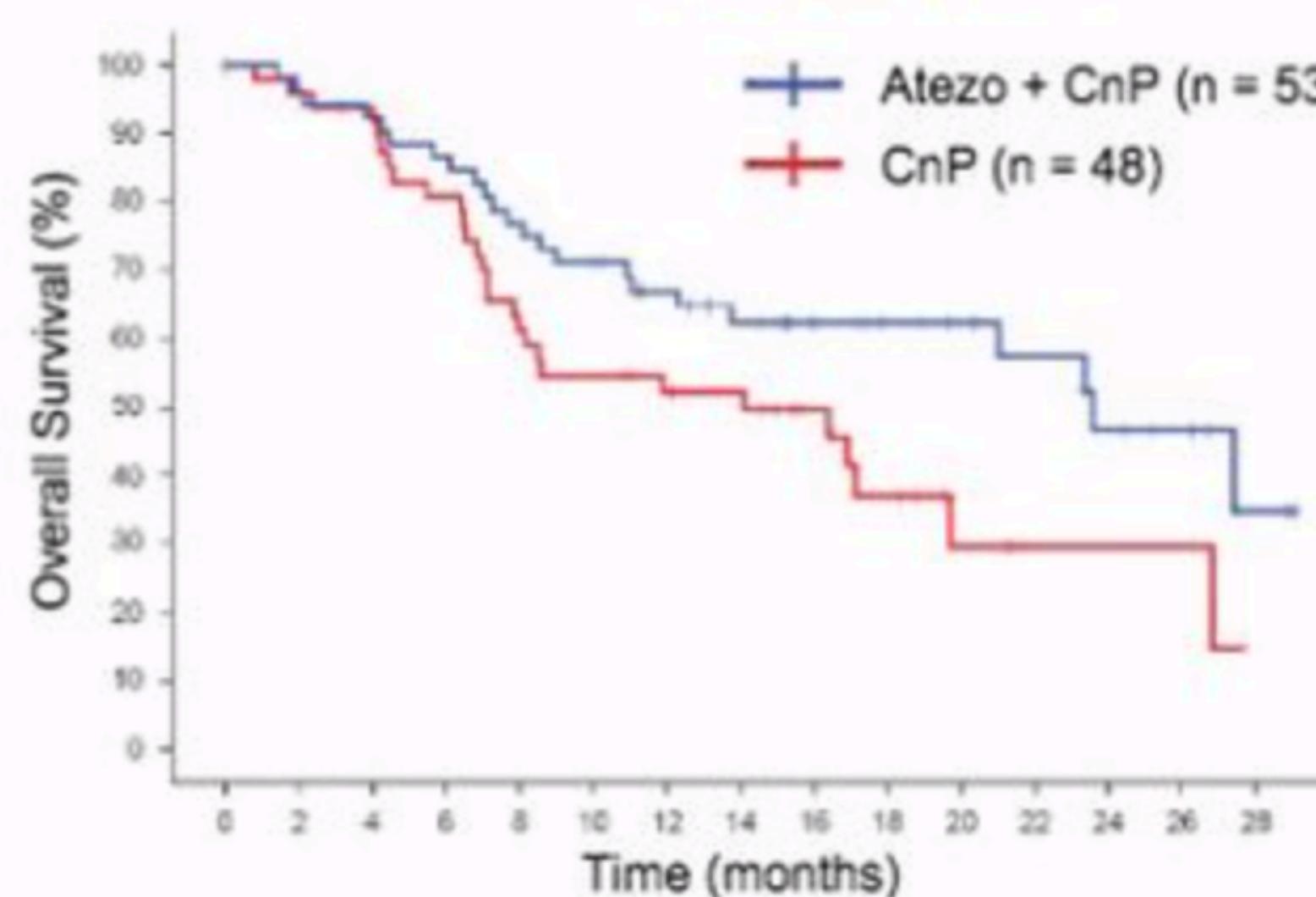


Data cutoff: January 22, 2018.

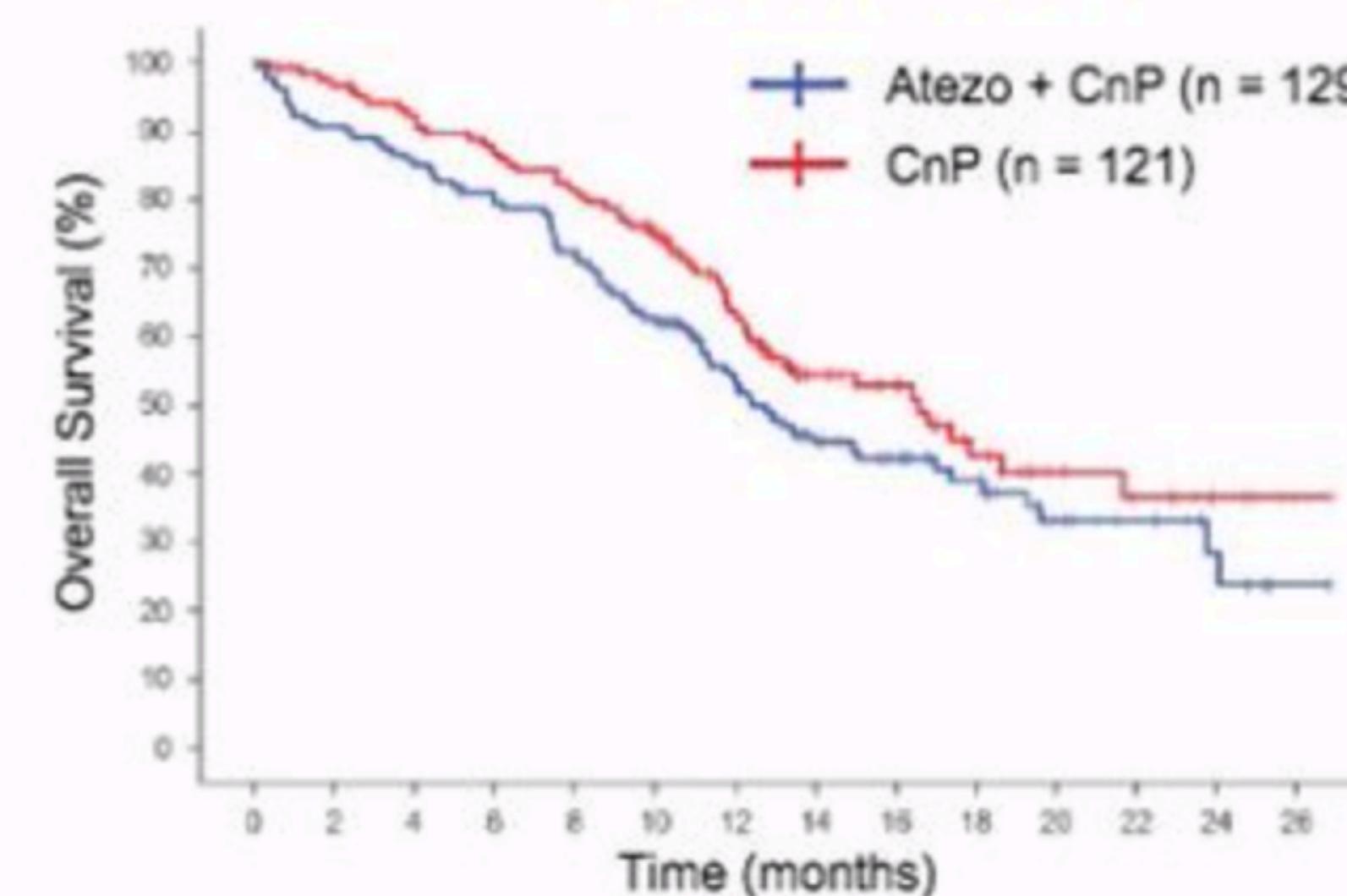
<sup>a</sup> Stratified HR.

# First Interim OS in PD-L1 Subgroups (Arm B vs Arm C)

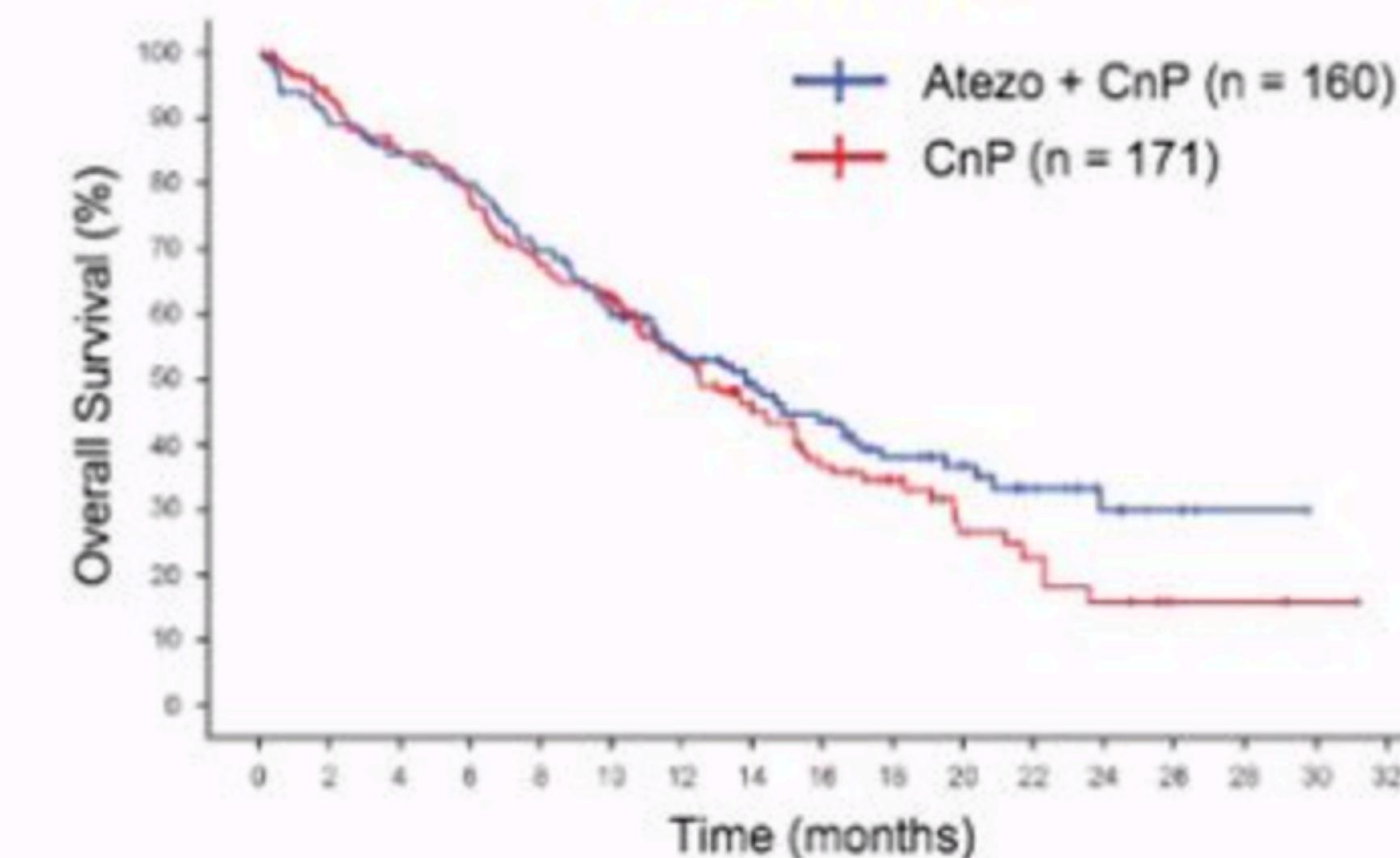
## PD-L1 High TC3 or IC3



## PD-L1 Low TC1/2 or IC1/2



## PD-L1 Negative TC0 and IC0



	Atezo + CnP	CnP	Atezo + CnP	CnP	Atezo + CnP	CnP
<b>12-month OS</b>	<b>67%</b>	<b>52%</b>	<b>54%</b>	<b>64%</b>	<b>53%</b>	<b>53%</b>
<b>24-month OS</b>	<b>47%</b>	<b>30%</b>	<b>28%</b>	<b>37%</b>	<b>30%</b>	<b>16%</b>
<b>Median OS, mo</b>	<b>23.6</b>	<b>14.1</b>	<b>12.4</b>	<b>16.6</b>	<b>13.8</b>	<b>12.5</b>
<b>HR<sup>a</sup> (95% CI)</b>	<b>0.56 (0.32, 0.99)</b>		<b>1.34 (0.95, 1.90)</b>		<b>0.86 (0.65, 1.15)</b>	

Data cutoff: January 22, 2018.

<sup>a</sup> Unstratified HR.