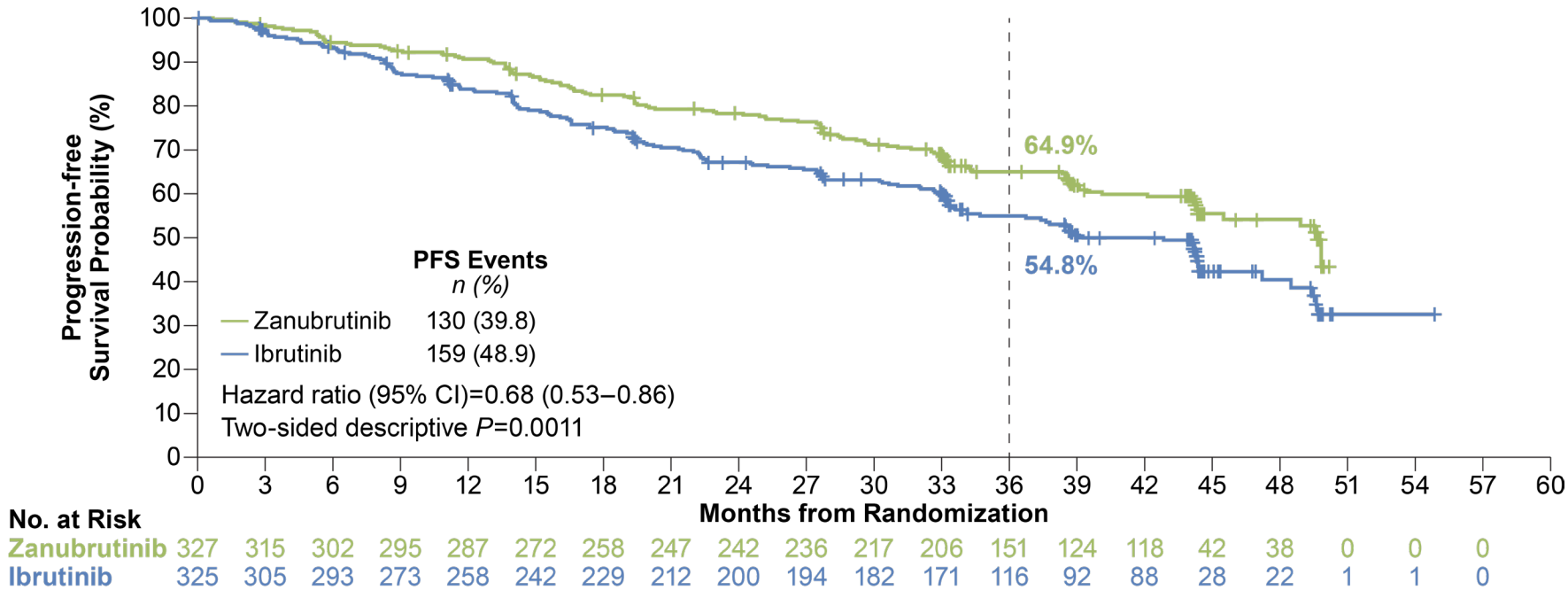


# Extended Follow-Up of ALPINE Randomized Phase 3 Study Confirms Sustained Superior Progression-Free Survival With Zanubrutinib vs Ibrutinib for Treatment of R/R CLL/SLL

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# Zanubrutinib demonstrated sustained PFS benefit over ibrutinib in patients with R/R CLL/SLL with a median follow-up of 39 months



- Durable PFS benefits were seen across major subgroups, including the del(17p)/TP53<sup>mut</sup> population

## Zanubrutinib continues to demonstrate a more favorable safety and tolerability profile compared with ibrutinib

	Zanubrutinib (n=324)	Ibrutinib (n=324)
<b>Median treatment duration, median (range), months</b>	<b>38.3 (0.4, 54.9)</b>	<b>35.0 (0.1, 58.4)</b>
<b>Any grade adverse events, n (%)</b>	<b>320 (98.8)</b>	<b>323 (99.7)</b>
Grade 3-5	235 (72.5)	251 (77.5)
Grade 5	41 (12.7)	40 (12.3)
<b>Serious adverse events, n (%)</b>	<b>165 (50.9)</b>	<b>191 (59.0)</b>
<b>Adverse events leading to, n (%)</b>		
Dose reduction	47 (14.5)	59 (18.2)
Dose interruption	196 (60.5)	201 (62.0)
Treatment discontinuation	64 (19.8)	85 (26.2)
Hospitalization	150 (46.3)	180 (55.6)
<b>Cardiac adverse events, n (%)</b>	<b>80 (24.7)</b>	<b>112 (34.6)</b>
<b>Serious cardiac adverse events, n (%)</b>	<b>11 (3.4)</b>	<b>31 (9.6)</b>

- No fatal cardiac events occurred with zanubrutinib treatment, and 6 fatal cardiac events occurred with ibrutinib
- Significantly fewer atrial fibrillation/flutter events occurred with zanubrutinib than with ibrutinib (6.8% vs 16.4%;  $P=.0001$ )

**With over 3 years of follow-up, these data reconfirm that zanubrutinib has improved efficacy over ibrutinib and a more favorable safety profile in patients with R/R CLL/SLL**