

**Additional File 1**

François Piette et al. Masitinib as an adjunct therapy for mild-to-moderate Alzheimer's disease: a randomised, placebo-controlled phase 2 trial

**Table S1.** Alternative sensitivity analyses for primary efficacy outcome ADAS-Cog at weeks 12 and 24 in the intent-to-treat population. For comparison against Table 2 - Summary of efficacy outcomes at weeks 12 and 24.

<b>Sensitivity Analysis Method</b>	<b>Week 12</b>			<b>Week 24</b>		
	<b>Masitinib</b>	<b>Placebo</b>	<b>P-value</b>	<b>Masitinib</b>	<b>Placebo</b>	<b>P-value</b>
<b>Observed Cases<sup>a</sup></b>						
Evaluable patients ( <i>n</i> )	17	6		9	5	
Improvement (decrease $\geq 4$ )	7 (41%)	1 (17%)	0.369	3 (33%)	1 (20%)	1.0
Worsening (increase $\geq 4$ )	1 (6%)	3 (50%)	0.040	1 (11%)	3 (60%)	0.095
Mean absolute change $\pm$ SD <sup>b</sup>	-2.6 $\pm$ 3.6	4.2 $\pm$ 6.6	0.016	-0.7 $\pm$ 7.8	6.5 $\pm$ 8.6	0.182
Mean treatment effect <sup>c</sup>	<b>6.8</b>			<b>7.2</b>		
<b>Last Observation Carried Forward</b>						
Evaluable patients ( <i>n</i> )	24	7		24	7	
Improvement (decrease $\geq 4$ )	9 (38%)	2 (29%)	1.0	8 (33%)	2 (29%)	1.0
Worsening (increase $\geq 4$ )	4 (17%)	3 (43%)	0.302	4 (17%)	3 (43%)	0.302
Mean absolute change $\pm$ SD	-1.0 $\pm$ 6.2	2.3 $\pm$ 7.9	0.247	-0.4 $\pm$ 7.2	3.6 $\pm$ 9.1	0.238
Mean treatment effect	<b>3.3</b>			<b>4.0</b>		
<b>Missing Data As Non Responders<sup>d</sup></b>						
Evaluable patients ( <i>n</i> )	26	8		26	8	
Improvement (decrease $\geq 4$ )	7 (27%)	1 (13%)	0.645	6 (23%)	1 (13%)	1.0
Worsening (increase $\geq 4$ )	10 (39%)	5 (63%)	0.417	11 (42%)	5 (63%)	0.429

<sup>a</sup>For the observed cases dataset no data imputation has been performed for the closed study centre. <sup>b</sup>Negative change reflects improvement.

<sup>c</sup>Treatment effect is difference between treatment and placebo mean absolute change in ADAS-Cog score relative to baseline. Positive change reflects a positive treatment effect. <sup>d</sup>For the missing data considered as non responders dataset, week 12 data for closed study centre (*n*=8 patients) was imputed using last observation carried forward for week 24.