

Novartis Pharmaceuticals Corporation

Novartis Pharmaceuticals Corporation, East Hanover, NJ 07936

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Novartis Pharmaceuticals Corporation 70-90% 18^{3),4),5),6),7)}

References

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3) P. Krupp, P. Barnes. Clozapine-associated agranulocytosis risk and aetiology. Br J Psychiatry, 160, Suppl17, 38-40, 1992.

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	Novartis ^{a)}	Novartis	Novartis (%)
Novartis	77	2	2.60
Novartis	77	6	7.79
Novartis	77	2	2.60

a) Novartis 1301, 1201, 1202, 1203

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Table 1: Summary of Me dDRA PT results

Table 2: Summary of Me dDRA PT results

Me dDRA PT	Me dDRA PT	Concentration (mg/L)	Count (No)	Count (Yes)	Result	Count	Count
0/36	0/36	450	57	4	No	0/36	1
0/37	0/37	300	63	4	No	0/37	1
0/41	0/41	300	91	25	Yes	0/41	1,2
0/43	0/43	200	29	4	No	0/43	1
0/31	0/31	225	105	15	No	0/31	1
0/37	0/37	200	71	3	No	0/37	1
0/46	0/46	400	57	10	Yes	0/46	1,2
0/51	0/51	0 ^{a)}	113	2	No	0/51	1
0/28	0/28	450	1467	29	No	0/28	1
0/29	0/29	300	706	22	No	0/29	1

0/36 = 0/36, 0/37 = 0/37

a) 0/51 = 0/51 > 100mg/L

Table 3: Summary of Me dDRA PT results

Table 4: Summary of Me dDRA PT results

Table 5: Summary of Me dDRA PT results

Table 6: Summary of Me dDRA PT results

Me dDRA PT	Count ^{a)}	Count ^{b)}	Count (%)
0/36			
0/37	249,378	829	0.33
0/41	50,866	358	0.70
0/43	28,079	311	1.11
0/31	18,032	102	0.57
0/37	346,355	1,600	0.46
0/46			
0/51	249,378	1,652	0.66

Table 1. Summary of adverse events (AEs) in the placebo and active treatment groups.

Table 1. Summary of adverse events (AEs) in the placebo and active treatment groups.

	Total	Placebo	Active Treatment	Active Treatment	Active Treatment	Active Treatment	Active Treatment	Active Treatment	Active Treatment
	n	n (%) ^{a)}	n (%) ^{a)}	n (%) ^{a)}	n (%) ^{a)}	n (%) ^{a)}	n (%) ^{a)}	n (%) ^{a)}	n (%) ^{a)}
AEs	1,600	785 (49.1)	30 (1.9)	169 (10.6)	87 (5.4)	2 (0.1)	95 (5.9)	372 (23.2)	60 (3.8)
AEs	4,261	1,974 (46.3)	43 (1.0)	413 (9.7)	214 (5.0)	7 (0.1)	199 (4.7)	1,386 (32.5)	25 (0.6)
AEs	2,490	945 (38.0)	45 (1.8)	298 (12.0)	198 (8.0)	2 (<0.1)	100 (4.0)	848 (34.1)	54 (2.2)

a) Percentage of patients with AE. b) Percentage of patients with AE.

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0800-222-8814

9:00-17:30

Table 1. Summary of adverse events (AEs) in the placebo and active treatment groups.

0120-003-293

9:00-17:30

Table 1. Summary of adverse events (AEs) in the placebo and active treatment groups.

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