Supplementary Online Content

Wu L, Song H, Zhang C, et al. Efficacy and safety of Pannax notoginseng saponins in the treatment of adults with ischemic stroke in China. *JAMA Netw Open*. 2023;6(6):e2317574. doi:10.1001/jamanetworkopen.2023.17574

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This supplementary material has been provided by the authors to give readers additional information about their work.

eTable 1. Efficacy Outcomes at 12 Months in the Modified Intention-To-Treat Cohort

	XST group	Control group	OR/HR	Р	
	(n=1487)	(n=1479)	(95% CI)	value	
Secondary outcomes					
Recurrent stroke, n (%)	33 (2.2)	41 (2.8)	0.80 (0.51	0.34	
			to 1.26)		
Functional independence, n (%) ^a	1342 (92.4)	1324 (91.5)	1.13 (0.87	0.36	
			to 1.48)		
No or minimal disability, n (%)ª	1243 (85.6)	1191 (82.3)	1.27 (1.05	0.01	
			to 1.53)		
Composite cerebrovascular events, n	39 (2.6)	45 (3.0)	0.86 (0.56	0.49	
(%)			to 1.32)		
EQ-5D score, median (IQR) ^a	95 (90, 98)	94 (90, 96)	-	0.009	
Barthel Index change from baseline to	20 (5, 40)	20 (5, 40)	-	0.17	
12 months, median (IQR)ª	. ,				

XST, Xuesaitong soft capsule; OR, odds ratio; HR, hazard ratio; CI, confidence interval; EQ-5D, EuroQoL Group 5-Dimension; IQR, interquartile range.

^a Functional outcome data at 12 months were missing in 67 patients; Data of EQ-5D score at 12 months were missing in 84 patients; Data of Barthel Index change from baseline to 12 months were missing in 80 patients; The number of missing data was similar between the two groups.

eTable 2. Distribution of the modified Rankin Scale at 3 months in the modified Intention-To-Treat Cohort

The score of modified Rankin Scale, n (%)	XST group (n=1487)	Control group (n=1479)
0	348 (23.4)	320 (21.6)
1	754 (50.7)	705 (47.7)
2	226 (15.2)	193 (13.1)
3	89 (6.0)	192 (13.0)
4	65 (4.4)	65 (4.4)
5	4 (0.3)	2 (0.1)
6	1 (0.1)	2 (0.1)

XST, Xuesaitong soft capsule.

	XST group	Control group	OR/HR	Р
	(n=1081)	(n=1096)	(95% CI)	value
Primary outcome				
Functional independence, n (%)	973 (90.0)	913 (83.3)	1.98 (1.51 to 2.60)	<0.001
Secondary outcomes				
Recurrent stroke, n (%)	9 (0.8)	9 (0.8)	1.02 (0.40 to 2.56)	0.98
No or minimal disability, n (%)	813 (75.2)	774 (70.6)	1.26 (1.04 to 1.53)	0.02
NIHSS score change from baseline to 3 months, median (IQR) ^a	-4 (-5, -3)	-4 (-5, -3)	-	0.07
Composite cerebrovascular events, n (%)	9 (0.8)	10 (0.9)	0.91 (0.37 to 2.25)	0.84
EQ-5D score, median (IQR) ^a	90 (80, 95)	90 (80, 95)	-	0.10
Barthel Index change from baseline to 3 months, median (IQR) ^a	15 (5, 35)	15 (5, 30)	-	0.01
Platelet counts and coagulation indicate	ors, median (IQR)			
Platelet counts, 10 ⁹ /L ^a	215.5 (173, 255)	216 (178, 259)	-	0.14
PT, sª	11.5 (10.7, 12.5)	11.4 (10.6, 12.4)	-	0.63
APTT, s ^a	28.6 (24.8, 33.1)	28.1 (24.8, 33.1)	-	0.58

eTable 3. Efficacy Outcomes at 3 months in the Per-Protocol Cohort

XST, Xuesaitong soft capsule; OR, odds ratio; HR, hazard ratio; CI, confidence interval; NIHSS, National Institutes of Health Stroke Scale; IQR, interquartile range; EQ-5D, EuroQoL Group 5-Dimension; PT, prothrombin time; APTT, activated partial thromboplastin time.

^a Data of NIHSS score change from baseline to 3 months were missing in 40 patients; Data of EQ-5D score at 3 months were missing in 40 patients; Data of Barthel Index change from baseline to 3 months were missing in 40 patients; Data of platelet counts, PT, and APTT at 3 months were missing in 252, 287, and 297 patients, respectively. The number of missing data was similar between the two groups.

	XST group	Control group	OR/HR	Р
	(n=1081)	(n=1096)	(95% CI)	value
Secondary outcomes				
Recurrent stroke, n (%)	23 (2.1)	30 (2.7)	0.78 (0.45 to 1.34)	0.36
Functional independence at 12 months, n (%) ^a	982 (93.1)	990 (92.3)	1.13 (0.81 to 1.56)	0.47
No or minimal disability, n (%)ª	918 (87.0)	894 (83.3)	1.34 (1.06 to 1.71)	0.02
Composite cerebrovascular events, n (%)	26 (2.4)	32 (2.9)	0.82 (0.49 to 1.38)	0.46
EQ-5D score, median (IQR) ^a	95 (90, 98)	95 (90, 96)	-	0.15
Barthel Index change from baseline to 12 months, median (IQR) ^a	20 (5, 40)	20 (5, 40)	-	0.21

eTable 4. Efficacy Outcomes at 12 months in the Per-Protocol Cohort

XST, Xuesaitong soft capsule; OR, odds ratio; HR, hazard ratio; CI, confidence interval; EQ-5D, EuroQoL Group 5-Dimension; IQR, interquartile range.

^a Functional outcome data at 12 months was missing in 49 patients; Data of EQ-5D score at 12 months were missing in 56 patients; Data of Barthel Index change from baseline to 12 months were missing in 54 patients. The number of missing data was similar between the two groups.

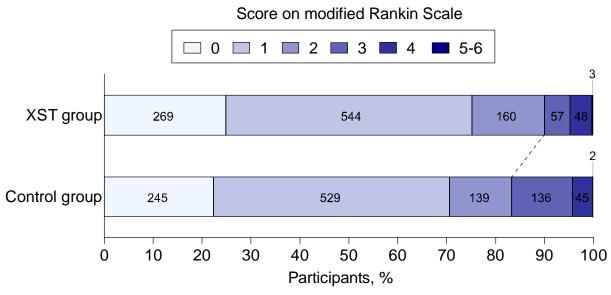
eTable 5. Medications and Status at 1 month in the modified Intention-To-Treat Cohort

	XST group (n=1487)	Control group (n=1479)	<i>P</i> value
Antithrombotic agents, n (%)	1434 (96.4)	1432 (96.8)	0.56
Antihypertensive agents, n (%)	637 (42.8)	600 (40.6)	0.21
Antidiabetic agents, n (%)	319 (21.5)	297 (20.1)	0.36
Anticholesteremic agents, n (%)	939 (63.1)	919 (62.1)	0.57
mRS score, median (IQR)	1 (1, 2)	1 (1, 2)	0.87
Functional independence, n (%)	1161 (78.1)	1134 (76.7)	0.36
NIHSS score, median (IQR) ^a	3 (2, 4)	3 (2, 4)	0.76
Rehabilitation therapy, n (%)	99 (6.7)	102 (6.9)	0.80

XST, Xuesaitong soft capsule; mRS, modified Rankin Scale; IQR, interquartile range; NIHSS, National Institutes of Health

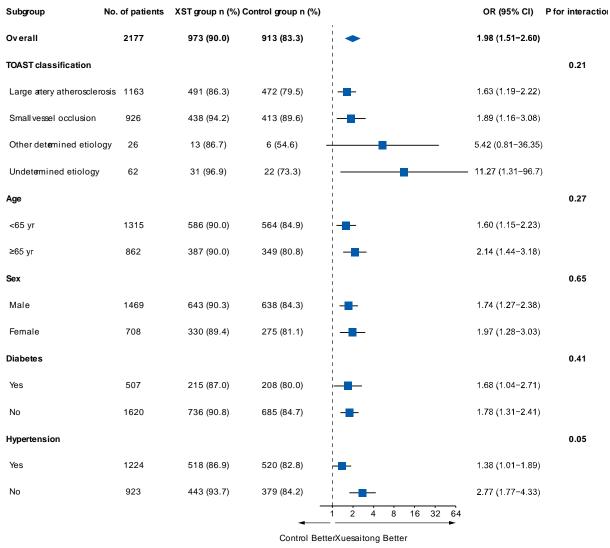
Stroke Scale.

^a Data of NIHSS score were missing in 2 patients in XST group.



eFigure 1. Distribution of the modified Rankin Scale at 3 months in the Per-Protocol Cohort

XST, Xuesaitong soft capsule.



eFigure 2. Subgroup Analyses in the Per-Protocol Cohort

XST, Xuesaitong soft capsule; OR, odds ratio; CI, confidence interval; TOAST, Trial of Org 10172 in Acute Stroke Treatment.

eAppendix. Trial Organization

Steering Committee:

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