## **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 (%) <sup>a</sup>	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) <sup>a</sup>	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.

<sup>a</sup> 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) <sup>a</sup>	-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) <sup>a</sup>	90 (80, 95)	90 (80, 95)	-	0.10
Barthel Index change from baseline to 3 months, median (IQR) <sup>a</sup>	15 (5, 35)	15 (5, 30)	-	0.01
Platelet counts and coagulation indicate	ors, median (IQR)			
Platelet counts, 10 <sup>9</sup> /L <sup>a</sup>	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 <sup>a</sup>	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.

<sup>a</sup> 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 (%) <sup>a</sup>	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) <sup>a</sup>	95 (90, 98)	95 (90, 96)	-	0.15
Barthel Index change from baseline to 12 months, median (IQR) <sup>a</sup>	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.

<sup>a</sup> 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.

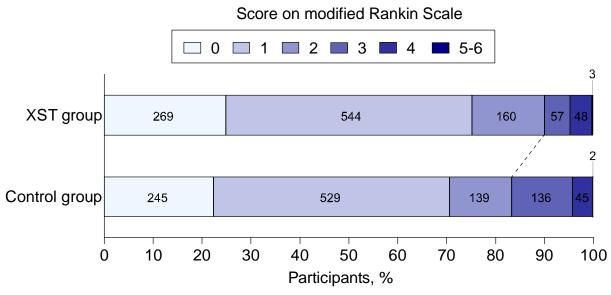
## eTable 5. Medications and Status at 1 month in the modified Intention-To-Treat Cohort

	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) <sup>a</sup>	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

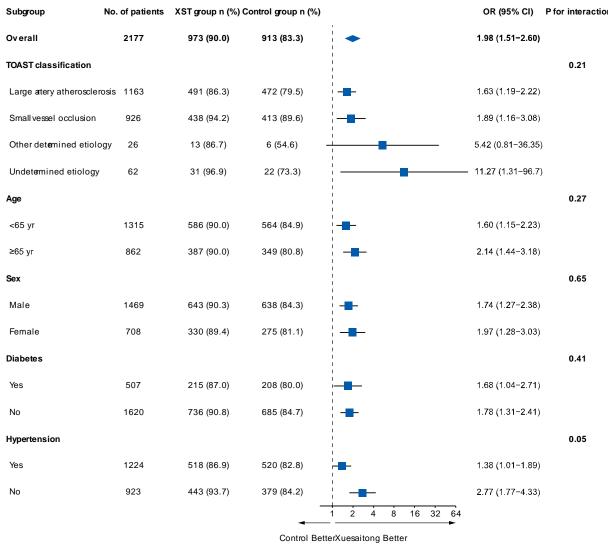
Stroke Scale.

<sup>a</sup> 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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#### **Executive Committee:**

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