Supplementary materials

Table S1. Study Procedure of BRANT trial

Measurement	**Day 1**	**Day 2**	**Day 3**	**Day 7**	**Day 90**
Demographic characteristics	**X**				
Current medical history taking	**X**				
Body temperature measurement	**X**				
Physical examination	**X**			**X**	**X**
Past medical history	**X**				
Pre-randomization medication after onset	**X**				
Regular blood pressure monitoring	**X**	**X**			
NIHSS score	**X**			**X**	**X**
mRS score	**X**			**X**	**X**
Barthel index score	**X**				**X**
Magnetic resonance image	**X**				
Evaluation of Intracranial vessels	**X**				
Evaluation of extracranial vessels	**X**				
Laboratory tests*	**X**				
ECG*	**X**			**X**	
Verification of inclusion/exclusion criteria	**X**				
Signed informed consent	**X**				
Randomization	**X**				
Blood tests after enrollment		**X**	**X**		

Urine and fecal examination			**X**		
Compliance			**X**		
Concomitant medication				**X**	**X**
Early neurological deterioration				**X**	
Major bleeding				**X**	**X**
Adverse Events/ Serious Adverse Events				**X**	**X**
*Remarks: ECG and laboratory data performed within 48 hours of onset before signing the informed consent form can be used as trial data.					