Safety and efficacy of treatment in acute ischemic stroke secondary to tandem internal carotid artery/middle cerebral artery occlusion: a single-center experience.

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OBJECTIVES

Compared to single intracranial artery occlusion, treatment and recanalization of tandem internal carotid artery (ICA) / intracranial artery occlusion (TO) remain extremely challenging. After all recent results from endovascular stroke trials, although new perspectives have been opened for the treatment of severe stroke secondary to intracranial artery occlusion [1-2], to date there is not yet a standardized recommendation for emergent management of TO. According to the low recanalization rate and poor outcome reported after intravenous thrombolysis [3], more data on current reperfusion strategies in TO are needed. The aim of our study was to report on safety and efficacy outcomes of endovascular treatment (ET) in a large cohort of patients with anterior circulation acute ischemic stroke (AIS) due to TO occlusion. We further investigated clinical and imaging predictors of outcome.

METHODS

Subjects and study procedures: Patients with anterior circulation AIS who underwent ET between august 2009 and January 2016 were retrospectively collected. Patients were selected based on following criteria: 1. CT-angiography (CTA) documentation of TO (extracranial ICA plus terminal ICA or M1 or M2 or A1 or a combination of them); 2. onset to groin puncture (OGT) within 5 hours from symptom onset; 3. severe clinical deficit with NIHSS≥ 10; 4. premorbid mRS≤ 2; 5. available 3 months follow-up.



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NEUROSCIENZE E RIABILITAZIONE

Treatment: Intravenous thrombolysis (IVT) (rtPA 0.9 mg/kg, 10% as bolus and the remaining in 60 min) within 4.5 hours after symptoms onset and continued during endovascular procedure. In case of an unstable or subocclusive plaque or dissection in the extracranial ICA, a direct closed-cells stenting was placed (Wallstent, Boston Scientific, Natick, MA, USA). Post-stenting angioplasty (Falcon Grande, Medtronic Minneapolis, MN, USA) was performed if a suboptimal angiographic result was achieved (lumen stenosis >30%). Intracranial **thrombectomy**using a 6 Fr guiding catheter (Neuron 070, Penumbra Inc., Alameda, CA, USA or Envoy DA, Codman & Shurtleff Inc, Raynham, MA, USA). Over the years, due to technological progress, different devices have been used such as penumbra aspiration system (Penumbra, Alameda, California, USA), stent retriever systems such as Solitaire AB (ev3 Inc, Plymouth, MN, USA), Revive SE (Codman & Shurtleff, Inc, Raynham, MA, USA), Trevo Retriever (Stryker Neurovascular, Mountain View, CA, USA) and, more recently, new aspiration devices such as Penumbra MAX ACE aspiration system.

Figure. 61 y.o. patient presenting with right hemiplegia and aphasia (NIHSS:23). Digital subtraction angiography revealed tandem occlusion (a) treated by means of IVT followed by thromboaspiration by PS started 125 min after symptom onset: the separator advancing in the MCA is shown in b while ICA and MCA recanalization (TIMI 3) obtained 254 min after symptom onset is shown in c.

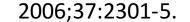
RESULTS										
Clinical and radiological characteristics		Univariable analysis of factors predicting 3-months outcome								
Baseline variables	N of patients=72	Variable	Good outcome (23)	Poor outcome (49)	p-value					
Age (ys) mean±SD	65.6±12.8	Age	59.43±11,72	68.57±12.41	0.004					
Male gender (%)	44(61.1)	Right hemisphere (%)	13(56.5)	24(49)	0.6					
NIHSS mean±SD	19±2.9	Single intracranial occlusion (%)	17(74)	28(57)	0.2					
Hypertension (%)	53(73.6)	IVT (%)	14(61)	25(51)	0.4					
Diabetes (%)	12(16.6)	GA (%)	20(87)	34(69.3)	0.1	Multivariable logistic	c regre	ession:	predictors of g	good outcome
Atrial fibrillation (%)	22(30.5)	CAS (%)	14(61)	21(43)	0.2				-	
Current Smoking (%)	3(4.1)	CAS patency (%)	14(100)	20(95)	1.00		OR	SE	95% CI	р
ASPECTS	6.9±2.28	Baseline ASPECTS (mean±SD)	8.22±1.81	6.34±2.15	0.0005	Age	0.92	0.03	0.87 - 0.98	0.013
Extracranial ICA	66(91.6)	Acute none APT (%)	8(34.5)	26(53)	0.2		4 00	0.07	0.00 4.00	0.440
subocclusion/occlusion		Thromboaspiration device (%)	15(65)	32(65.3)	1.00	ASPECTS	1.33	0.27	0.90 - 1.98	0.149
Site of intracranial occlusion (%)		Rescue device (%)	6(26)	17(34.7)	0.5	TICI 2b-3	6.40	6.39	0.90 - 45.36	0.063
M1	36(50)	OGP (min, mean±SD)	240.23±124.72	229.51±76.67	0.66	Device passages	0.93	0.22	0.58 - 1.47	0.750
M2	8(11.1)	OTR (min, <i>mean</i> ±SD)	316.95±56.34	328.8±83.15	0.63					
t-ICA	3(4.1)	GPR (min, mean±SD)	78.09±35.3	97.06±52.26	0.12	Fair Collaterals	5.18	4.09	1.10 - 24.33	0.037
A1 + M1	3(4.1)	TICI≥2b (%)	21(91.3)	25(51)	0.001	ICH	0.21	0.17	0.04 - 1.08	0.62
t-ICA + M1	22(30.5)	Device passages (mean±SD)	2.30±1.58	3.57±1.91	0.007		atorolo		ignificant and	_ independent
Collateral adequacy (%)		Fair Collateral flow (%)	18(78)	13(26.5)	0.0001	Age and good colla				independent
Fair	31(43)	Onset NIHSS (mean±SD)	18.17±3	19.47±2.82	0.07	pre	dictor	s of goo	od outcome	
Poor	41(57)	ICH (%)	6(26)	28(57)	0.02					
M1: proximal portion of the middle cerebral artery; M2: mid portion of the middle cerebral artery; A1: proximal portion of anterior cerebral artery; t-ICA: terminal intracranial internal carotid artery.		Symptomatic ICH (%)	0	9(18.3)	0.04					
		Procedural IV Heparin (%)	11(48)	29(59)	0.4					
		Follow-up ASPECTS (mean±SD)	4.35±2.38	1.84±1.91	<0.0001					
Treatment characteristics		Univariable analysis of risk factors for sICH								
Extracranial CAS (%)	35(48.6)	Variable	sICH (9)	No sICH(63)	p-value	Multivariable lo	aistic	rogross	ion : predictor	s of sICH
IVT (%)	37(51.3)	IV Heparin (%)	7(77)	35(55.5)	0.28		gistic	i egi ess		
APT (%)		IV Thrombolysis (%)	4(44)	33(52)	0.73		OR	SE	95% CI	р
Dual	12(16.6)	<i>IV</i> Lysine acetylsalicylate (%)	3(33)	5(8%)	0.05	IV Lysine ASA	4.32	5.83	0.30 - 60.97	0.278
Single	26(36.1)	Double APT (%)	1(11)	11(17.5)	1.00	ASPECTS	0 00	0 10		0 111
None	34(47.2)	Baseline ASPECTS (mean±SD)	5.1±1.7	7.2±2.1	0.006	ASPECTS	0.83	0.18	0.54 - 1.29	0.414
Intravenous heparin (%)	40(55.5)	Fair collateral flow (%)	0	31(49)	0.008	Fair Collaterals	0.09	0.11	0.01 - 1.09	0.059
OTR, min (mean±SD)	325.1±95	Onset NIHSS (mean±SD)	19.6±2.6	18.9±2.9	0.5	Rescue device	10.74	11.28	1.37 - 84.13	0.024
	3.2±1.9	CAS %	4(44)	31(49)	1.00					
Device attempts (n)		Thromboaspiration device (%)	6(66.5)	41(65)	1.00	GPR	1.01	0.01	0.99 - 1.04	0.253
CAS: carotid artery stenting; IVT: intravenous thrombolysis; APT: antiplatelet therapy; OTG: onset to groin puncture time; OTR: onset to reperfusion time; GTR: groin puncture to reperfusion time		Rescue device (%)	7(77)	16(25.5)	0.003	Device passages	0.89	0.28	0.47 - 1.67	0.723
		TICl≥2b (%)	4(44)	42(66.5)	0.2					
		OTR (min. <i>mean</i> ±SD)	275±69.7	318.7±97	0.13	The only independent predictor of sICH was the use of an				
REFERENCES		GPR (min, <i>mean</i> ±SD)	129.66±78.22	85.47±40.12	0.008	addition	nal dev	vice for	thrombectomy	/.
 Berkhemer OA, Fransen PS, Beumer D, et al. A randomized trial of intraarterial treatment for acute ischemic stroke. N Engl J Med 2015;372:11-20. Goyal M, Demchuk AM, Menon BK, et al. Randomized assessment of rapid endovascular treatment of ischemic stroke. N Engl J Med. 2015:272:1010.1020 			4.4±2.7	2.9±1.7	0.03					
				CON	CLUSIO	NS				
		1. ET for AIS secondary to TO can be performed with reasonable safety and efficacy.								

3) Rubiera M, Ribo M, Delgado-Mederos R, et al. Tandem internal

1. ET for AIS secondary to TO can be performed with reasonable safety and efficacy.

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predictor of poor outcome after systemic thrombolysis. Stroke



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