## Abstracts

Right internal iliac artery was then catheterized. Angiography and CBCT showed a less anastomotic right middle rectal artery involvement in hemorrhoid vascularization but confirmed the origin of the bleeding with contrast within the rectum. Hence, only one microcoil was used and some gelfoam pledget. Result(s): Angiographies and CBCT confirmed the origin of the bleeding and showed satisfying final result with complete exclusion of internal hemorrhoids. Stabilization of vital signs was perceived during the procedure. A three weeks clinical and endoscopic follow-up showed no recurrence of the bleeding. Conclusion(s): Emborrhoid technique can be used in urgent treatment of massive hemorrhoid bleeding even when the access to the inferior mesenteric artery is not possible by embolizing middle rectal artery as it is the main blood supply of hemorrhoid in this case. CBCT helps identifying the main feeder arteries and guides the embolization.

## P501

Effectiveness of ACE68 and ACE64 Catheters in Anterior Circulation Large Vessel Occlusion: Promise Study Subgroup Analysis by Occlusion Location

Rosario Papa, Peter Schramm<sup>1</sup>, Pedro Navia<sup>2</sup>, Joaquin Zamorra Parra<sup>3</sup>, Alejandro Tomasello Weitz<sup>4</sup>, Werner Weber<sup>5</sup>, Jens Fiehler<sup>6</sup>, Patrik Michel<sup>7</sup>, Vitor Pereira<sup>8</sup>, Timo Krings<sup>8</sup>, Laurent Pierot<sup>9</sup>

A.O.U Policlinico, Messina, Italy, Acireale, Italy, <sup>1</sup>Universitätsklinikum Schleswig-Holstein, Lübeck, <sup>5</sup>Universitätsklinikum Knappschaftskrankenhaus Bochum, Bochum, <sup>6</sup>Universitätsklinikum Hamburg-Eppendorf, Hamburg, Geramny, <sup>2</sup>Hospital Universitario Donostia, San Sebastian, <sup>3</sup>Hospital Clínico Universitario Virgen de la Arrixaca, Murcia, <sup>4</sup>Vall d'Hebron Hospital, Barcelona, Spain, <sup>7</sup>Centre Hospitalier Universitaire Vaudois, Lausanne, Switzerland, <sup>8</sup>University of Toronto, Toronto, Canada, <sup>9</sup>Hôpital Maison Blanche, Reims, France E-mail: medicalwriting@penumbrainc.com

Background: The PROMISE Study documented safety and efficacy of ACE68 and ACE64 Reperfusion Catheters in patients with acute ischemic stroke (AIS) from large vessel occlusion (LVO), treated with ADAPT (A Direct Aspiration First Pass Technique) as frontline treatment. This analysis examines the safety and efficacy by occlusion location. Method(s): PROMISE was a prospective, single-arm, multicenter study. Inclusion criteria were anterior circulation LVO within 6 hours of ictus; NIHSS  $\geq$  2; CT-ASPECTS  $\geq$  6; or MR-ASPECTS ≥ 5. Primary endpoints included successful angiographic revascularization (mTICI 2b-3), clinical independence (mRS 0-2) at 90 days. Secondary endpoints included safety events, functional improvement at 7-10 days. This subgroup analysis investigates these endpoints by occlusion location. Result(s): Across 20 European centers, 204 patients (median age 74 [IQR 65-80]) were enrolled. Primary occlusion locations were 21.1% (43/204) ICA/Carotid-T, 60.8% (124/204) M1, 18.1% (37/204) M2. Median baseline CT ASPECT score was 9 [IQR 8-10]. Median baseline NIHSS score was 16 [IQR 11-20]. Prior to procedure, 61.8% (126/204) patients had IV rtPA. Immediate post-procedural angiographic revascularization (mTICI 2b/3) rate was 93.1% (190/204), 90-day mRS 0-2 rate was 61.0% (122/200). Subgroup analysis by occlusion location: ICA/ Carotid-T final revascularization (mTICI 2b/3) was 95.3% (41/43), 90-day mRS 0-2 was achieved in 64.3% (27/42); MCA M1 final revascularization was 92.7% (115/124) and 90-day mRS 0-2 rate was 57.0% (69/121); MCA M2 final revascularization was 91.9% (34/37) with 70.3% (26/37) having 90-day mRS 0-2. Safety rates were favorable (sICH=2.9%; ENT=1.5%); 90-day morbidity (mRS 3-5) was observed in 31.5% (63/200), and 90-day all cause-mortality was observed in 7.5% (15/200). Device and procedure-related SAEs at 30-days were reported in 2.0% (4/204) and 4.4% (9/204) of subjects, respectively. There was no significant difference in safety rates by treatment location. Conclusion(s): This subset analysis of the PROMISE study demonstrates the ACE68/64 Reperfusion Catheters are able to achieve high mTICI scores, with comparable safety profile and 3-month mRS in all studied locations for patients with LVO-AIS.