

**TECHNOLOGY NOTE** 

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## A novel hybrid catheter-directed technique to treat intermediate-high risk pulmonary embolism

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Despite advances in the management of acute pulmonary embolism (PE), the mortality rate in intermediate-high-risk and high-risk PE remains significant [1, 2]. Recently, percutaneous catheter-directed therapies have progressed with contemporary PE management [3–6].

A 44-year-old woman with a history of disseminated ovarian cancer was admitted with intermediate-high risk PE. Several days of unfractionated heparin (UFH) infusion was ineffective despite therapeutic range of activated partial thromboplastin time (APTT). Physical examination revealed heart rate (HR) 120 bpm, respiratory rate (RR) 30/ /min and blood pressure (BP) 95/60 mmHg. Blood test showed elevated troponin concentration 0.6 ng/L (normal range < 0.01 ng/L) and low arterial saturation (SaO<sub>2</sub>) 88% despite oxygen supplementation (flow rate 15 L/min). Echocardiography demonstrated right ventricle dysfunction (right ventricle/ /left ventricle [RV/LV] diameter ratio 1.5 and tricuspid annular plane systolic excursion [TAPSE] 13 mm). A bedside lower-limb compression ultrasonography showed a deep vein thrombosis of the left popliteal vein. The Pulmonary Embolism Severity Index indicated very high mortality risk (144 points — class V). Our institutional PE response team qualified the patient to catheter-directed treatment.

The right common femoral venous access was obtained with a 6-F vascular sheath. A pigtail catheter (6-F; Cordis, Hileah) was inserted into the right and left pulmonary arteries and a selective angiography was performed. Then, pulmonary arterial pressures (PAPs) were recorded under fluoroscopy according to current guidelines [7]. The systolic/diastolic/mean PAPs were 63/17/35

mmHg, respectively. Pulmonary angiography revealed large thrombus burden bilaterally mainly in lobar pulmonary arteries.

Due to well thrombi organization it was decided to give alteplase directly to the thrombus in preparation for aspiration.

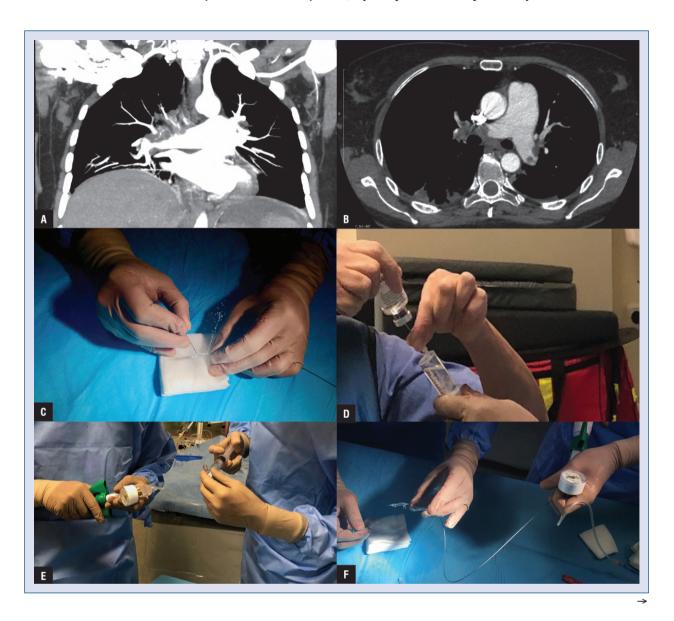
For this purpose, an angioplasty balloon catheter Maverick 2  $^{\text{TM}}$  4.0  $\times$  20 mm (Boston Scientific, Marlborough) was manually perforated with a needle at different sites and subsequently inserted into the target segmental artery of the right upper lobe tracked along Sion blue guidewire (Ashahi, Aichi). The balloon catheter was inflated with an alteplase solution (concentration 1 mg/1 mL) using a deployment pressure of 6 atmospheres (Fig. 1A–F). The balloon catheter was subsequently placed and inflated in the two other segmental arteries of the right upper lobe. The total amount of 3 mg of alteplase was injected via the balloon catheter within 3 min — each inflation lasted 1 min.

Subsequently, a 115 cm, 8-F Indigo CAT8 XTORQ catheter (Penumbra, Alameda, California) was inserted through a long 90 cm, 8 F Flexor sheath (Cook; Bloomington) to perform continuous aspiration thrombectomy (CDT) procedure in the right and left lobar pulmonary arteries. A direct-aspiration first-pass technique was performed in order to attach a large part of the thrombus to the catheter tip by suction and then pull it out through the sheath. Subsequently, separator wire was repeatedly passed through the thrombus to fragment and facilitate aspiration (Fig. 1G–I). The procedure was terminated with the improvement of hemodynamic parameters (on table HR normalization < 100 bpm and PAPs reduction to 46/15/26 mmHg).

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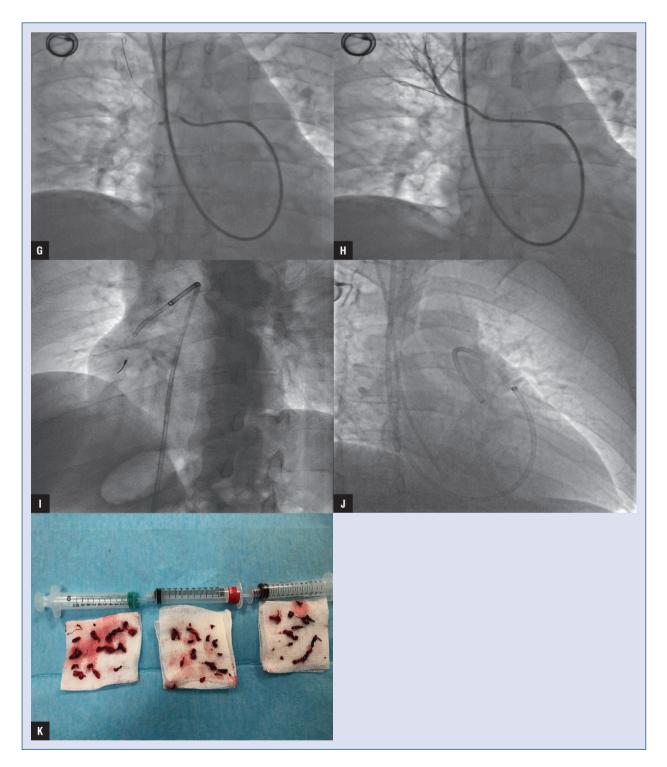


Eventually, blood flow significantly improved bilaterally in proximal pulmonary arteries as well in the lung periphery. During the procedure, intravenous UFH was continued at normal intensity with a goal APTT of 1.5–2.5 upper normal range.

During the subsequent monitoring in the cardiac intensive unit the patient's clinical status significantly improved. There were no complications related to the procedure. 24 hours after CDT patient's HR was 80 bpm, RR was 20/min and  $\rm SaO_2$  was 94% on room air, respectively. Echocardiography showed significant RV function improvement (TAPSE 20 mm, RV/LV diameter ratio 1.0). Anticoagulation with UFH was continued for 48 hours after the procedure and then a weight-adjusted low molecular weight heparin was introduced. The clinical 6-month follow up was uneventful and the patient continued with chemotherapy.

Interventional techniques are focused to quickly relieve blockage and restore pulmonary blood flow with subsequent improvement in hemodynamic status, oxygenation and RV function [8, 9]. The use of two catheter-directed techniques, balloon-delivered local thrombolysis and continuous aspiration thrombectomy, during one procedure was unique. In the present case, efficient clot removal immediately improved the patient's condition and appeared important to ameliorate the outcome with low bleeding risk. An alteplase given directly into the pulmonary arterial circulation reaches a steady-state and is capable of dissolving thrombi [10]. It was decided to administer a thrombolytic drug directly into the segmental arteries to provide the highest drug concentration in most obstructed vessels. These results suggest a possible synergistic effect of low dose

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**Figure 1.** Step by step a hybrid catheter-directed treatment of pulmonary embolism; **A, B.** The initial computed tomography angiograms showing extensive thromboembolic material burden in right pulmonary arteries; **C.** The angioplasty balloon perforation with a needle at different sites; **D.** The alteplase solution preparation (concentration 1 mg/1 mL); **E, F.** Filling of the handheld pump and the angioplasty balloon with the alteplase solution; **G.** The angioplasty balloon placement and inflation with the alteplase solution into the target segmental artery of the right upper lobe; **H.** The selective angiogram performed after transballoon alteplase delivery; **I, J.** The catheter-directed mechanical aspiration thrombectomy procedure with the Indigo device; **K.** An image of the removed clots.

thrombolytic therapy and aspiration thrombectomy, even in patients with a relatively long duration of PE symptoms.

## Conflict of interest: None declared

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