**Introduction**

Percutaneous transluminal procedures represent an increasingly used and effective treatment of peripheral artery disease (PAD) [1, 2]. The introduction of self-expanding nitinol stents allowed treatment of long lesions/occlusions of the superficial femoral arteries (SFA) improving the long-term results of the endovascular revascularization, with primary patency in SFA occlusions up to 80% at 2 years follow-up [3-5]. Antiplatelet therapy with acetylsalicylic acid is indicated for all patients with PAD undergoing peripheral transluminal angioplasty, while dual antiplatelet therapy, associating clopidogrel with aspirin, is suggested in clinical practice as antithrombotic therapy after peripheral stent implantation. In particular, dual antiplatelet therapy is particularly recommended in high-risk patients with lesions in the femoropopliteal segments and in the smaller diameter tibial arteries [6-8]. SFA nitinol stent thrombosis after discontinuing antiplatelet therapy, although uncommon, is an urgent clinical complication of acute limb ischaemia (ALI), necessitating immediate percutaneous intervention.

Rheolytic thrombectomy (RT) with AngioJet® (Possis Medical, Inc., Minneapolis, MN, USA) has been previously approved by the Food and Drug Administration for the treatment of acute thrombosis of venous grafts and native coronary arteries. The AngioJet® system consists of a drive unit control console, a pump set and an AngioJet® spiroflex rapid exchange rheolytic thrombectomy catheter. The latter is a dual lumen, sterile, single-use catheter designed to remove thrombi from coronary conduits. High velocity saline jets directed back into the catheter create a localized low-pressure zone at the distal tip (Bernoulli principle), which results in the suction, break-up and removal of thrombus through the outflow lumen.

During percutaneous transluminal coronary angioplasty (PTCA) of venous grafts, RT has been shown to be safe and its use has significantly improved angiographic results, without increasing mortality [9]. Conversely, data regarding the use of...
this procedure in native coronary artery are controversial and while one randomized study leaded in a high volume center (Florence trial) showed the same results obtained for venous grafts [10], Ali et al. [11] demonstrated neither an improvement of angiographic results nor an improvement of the mortality by the RT.

Despite no randomized clinical trials support the utility of AngioJet® in other districts, RT could be potentially used during percutaneous transluminal angioplasty (PTA) of other vessels, in particular of femoro-popliteal arteries. Indeed, here we describe the cases of three patients presenting with acute limb ischemia due to nitinol stent or PTFE femoro-popliteal graft thrombosis, in which limb salvage was obtained by AngioJet rheolytic thrombectomy and re-stenting.

The patients were admitted to our Department due to severely reduced walking capacity, rest pedal pain with or without ulcer toe, pointing out the diagnosis of ALI. All patients were aged 72±1 years, presented hypertension and hypercholesterolemia. Two patients had rheumatoid arthritis and history of previous PTA and stent implantation for total occlusion of SFA, while one patient presented a history of right femoro-popliteal PTFE by-bass for ALI followed by PTA and stent implantation of the distal anastomosis.

After the first procedure, all patients were discharged with optimal dual antiplatelet therapy excepted for the second patient who referred history of ASA allergy. Because of the onset of other clinical conditions (see below), two patients discontinued the antiplatelet therapy during the follow-up and presented stent thrombosis which was successfully removed using the Possis AngioJet® catheter. To the best of our knowledge, these are the first three reports on the use of the AngioJet® rheolytic thrombectomy in ALI for stent or PTFE graft thrombosis.

Case Presentation

Patient 1

A 74-year-old woman with hypercholesterolemia, rheumatoid arthritis and a history of ASA allergy was admitted to our Department complaining severely reduced walking capacity (maximum walking distance < 20 meters), rest right pedal pain and ulcer toe, suggesting the diagnosis of critical lower limb ischemia. At that time the patient’s therapy consisted of atorvastatin 10 mg, ASA 175 mg, bisoprolol 2.5 mg, valsartan 160 mg and metotrexate 2.5 mg/die. Autoimmunity evaluation on blood samples showed increased IgG anti-cardiolipin levels in the absence of IgM autoantibodies. Duplex ultrasound confirmed the occlusion of the proximal right superficial femoral artery. According to symptoms and clinical signs, the patient was classified as Fontaine’s stage IV.

Peripheral angiography confirmed the proximal occlusion of SFA (TASC classification type C lesion). After heparin infusion and using crossover technique, PTA with stent implantation of the SFA and popliteal artery was performed (nitinol stents Xpert 8.0x40 mm and 2 Protégé Everflex 7.0x120 mm and 6.0x120 mm), providing an optimal angiographic result and a good runoff to distal vessels.

The patient received a loading dose of clopidogrel (300 mg) P.O. and was discharged on dual antiplatelet therapy (ASA 100 mg/die plus clopidogrel 75 mg/die) in addition to her homing therapy. Duplex ultrasound performed two days later showed normal flow of treated segments and downstream.

Seven days after the procedure, the onset of gastrointestinal bleeding (Hb 8.70 g/dl) induced her primary care physician to stop her dual antiplatelet medication. After 15 days, the patient returned to our Department for rest pain and right ALI. Right pedal pulse was absent. The peripheral angiography showed acute stent thrombosis from the proximal right SFA up to popliteal segment (Figure 1, A and B). Thus, anticoagulation with heparin was immediately started and several attempts to open the occluded stent were performed. First, a 0.035-inch J-shaped Terumo stiff guide-wire was used to cross the thrombotic occlusion and then multiple dilatations of the stents within the SFA were performed with a 3.0x40 mm balloon catheter (Amphirion), without any good result. Thus, we decided to operate a mechanical thrombectomy using the Possis AngioJet® catheter (Figure 1C). As a result, mechanical aspiration of the thrombus allowed a good angiographic runoff and then two self-expandable stents (Dyna Link 6.0x100 mm and Protégé Everflex 6.0x60.0 mm) were implanted. After the procedure, the patient reported that pain disappeared and control angiography of the treated arterial segment revealed an optimal angiographic result and a good runoff to patent vessels below the knee (Figure 1, D-E-F). At 1 year follow-up, the ulcer on her toe healed and no stenotic lesions were found at colour Doppler ultrasound.

Patient 2

A 74-year-old woman with hypercholesterolemia, rheumatoid arthritis and a history of ASA allergy was admitted to our Department with rest pedal pain and duplex ultrasound evidence of left proximal SFA occlusion (Fontaine’s stage III). Peripheral angiography confirmed the proximal occlusion of SFA (TASC Classification type C). After heparin infusion, using crossover technique, PTA with stent implantation in the left SFA and popliteal artery was performed (2 nitinol stents, Protégé Everflex 6.0x120 mm) obtaining a good angiographic result. The patient received a loading dose of clopidogrel (300 mg) P.O. and she was discharged on clopidogrel (75 mg/die) without ASA.

At 2-month follow-up, duplex ultrasound scan showed a normal flow through the stents and the downstream segments. The patient referred an improvement of walking capacity and the disappearance of rest symptoms until 4 months later when she returned to our Department for ALI. Notably, patient never stopped antiplatelet therapy with clopidogrel. Pedal pulses were absent and duplex ultrasound indicated an acute thrombosis in the distal SFA/popliteal segment. The angiography confirmed an occlusion of the left SFA 4 cm before the previous implanted stent, up to the upper border of patella (the total length of thrombotic lesion was approximately 20 cm; Figure 2, A and B).

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Using crossover technique, PTA of the left SFA was performed. Initially, a 0.035” J-shaped Terumo guide-wire was passed through the thrombotic lumen up to the end of the obstruction; then multiple PTAs were performed, using an Admiral Xtreme Balloon Catheter 5.0x60 mm, without obtaining any good result. Thus, the Possis AngioJet® catheter was passed on the 0.014” guide-wire through the thrombotic lumen (Figure 2C), and the thrombus was aspirated. Once previously stented vessel was re-opened, the result was optimized with nitinol self-expanding re-stenting using two Protégé Everflex nitinol stents (6.0x150 mm and 6.0x100 mm), leading to satisfactory angiographic outcome (Figure 2, D-E-F). A month after discharge, pain and walking limitations disappeared. Twelve months later, colour Doppler ultrasound showed patent stent segment and normal flow pattern.

**Patient 3**

A 72-year-old man, with atrial fibrillation, hypercholesterolemia and history of right femoropopliteal PTFE bypass implantation for right ALI, was admitted to our Department because of rest pedal pain and duplex ultrasound evidence of both left proximal SFA and by-pass occlusion with distal collateral flow (Fontaine’s stage III).

Peripheral angiography confirmed the proximal occlusion of both SFA and the femoro-popliteal by-pass. Moreover, a critical stenosis of the left external iliac artery was found. In order to perform a PTA, 6F long sheaths (Therumo Destination) were positioned, 5000 U.I. of heparin were administered intravenously and the PTFE graft was then passed with a 0.035” J-Shaped Terumo stiff guide-wire. The critical stenosis of external iliac artery was treated with PTA and stent implantation (Complete 7x40 mm) followed by PTA and stenting of the distal graft anastomosis (Protégé Everflex 7x60 mm). Final angiography revealed optimal result. The patient received a loading dose of clopidogrel (300 mg) P.O. and ASA (500 mg) i.v. and he was discharged with clopidogrel (75 mg/die) and ASA (100 mg/die). Two months later, duplex ultrasounds scan demonstrated a normal flow through the stents and the downstream segments.

After the procedure, the patient was asymptomatic for 4 months under dual antiplatelet therapy. However, he was recommended to stop clopidogrel and to start warfarin anticoagulation therapy in addition to ASA, in order to prevent thromboembolic events because of the persistence of atrial fibrillation. Nevertheless, clopidogrel was incorrectly interrupted before reaching therapeutic range of INR. One month later, he returned to our Department for ALI (Fontaine’s stage IV). Pedal pulses were absent and duplex ultrasound showed acute thrombosis of both right PTFE graft and previous implanted stent at the distal anastomosis. Angiography confirmed the complete occlusion of both by-pass and the previously deployed stent at the distal anastomosis (Figure 3A). PTA of right PTFE by-pass was performed by using the crossover technique. A 0.035” J-shaped Terumo stiff guide-wire was crossed

![Image](https://via.placeholder.com/150)

**Figure 1.** - Proximal thrombotic occlusion of the left superficial femoral artery (A) and slow flow to the popliteal artery at angiography (B). C: AngioJet catheter positioned in the left superficial femoral artery to remove thrombus. D, E, F: thrombus aspiration and vessel patency.
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through the lumen up to the distal end of the tibio-peroneal trunks. Using an Admiral Xtreme Balloon Catheter 6.0x40 mm multiple PTAs were performed without reaching any good result. Furthermore, the angiography showed multiple minus images, revealing thrombus persistence (Figure 3B). Thus, a mechanical thrombectomy using Possis AngioJet® was performed using the Spiroflex catheter on a 0.014” guide-wire obtaining an optimal result up to the distal segment of the by-pass. Finally, a new stent (absolute Pro 7.0x60 mm) was implanted in the distal femoral popliteal artery in order to reach a better clinical outcome (Figure 3C). One month after discharge, pain and walking limitations disappeared. Six months later, colour Doppler ultrasound demonstrated patent stent segment and a normal flow pattern.

Discussion

The primary goals of ALI treatment are to relieve ischemic pain and to prevent limb loss. Primary percutaneous revascularization with new nitinol stents in association to dual antiplatelet therapy is safe and improves late patency rates [1, 3, 5]. Here we describe three cases of SFA stent thrombosis as a consequence of inadequate or prematurely discontinued dual antiplatelet therapy after SPA stenting. It is important to emphasize that much of the supporting evidence for antiplatelet therapy after SPA stenting is extrapolated from what is related to the coronary circulation [6]. Dual antiplatelet therapy for three to six months after PTA and nitinol stent placement is usually recommended to prevent early stent failure because of thrombosis at the intervention site. Our patients were characterized by high-risk metabolic-profile, with long peripheral lesions (type C TASC classification) and, strikingly, two of them were affected by rheumatoid arthritis, a chronic inflammatory disease associated with pro-atherothrombotic state. Notably, in this setting the discharge of dual antiplatelet therapy for major bleeding (case report 1) or a suboptimal antiplatelet therapy for ASA allergy (case report 2) may have induced the thrombotic event development following the primary SFA stent placement. In addition, the clinical case 3 reflects the different roles of anticoagulation and antiplatelets drugs, and the key role of reaching the therapeutic INR target.

Although it is well known that initial therapeutic strategy for ALI aims at preventing thrombus propagation and worsening of ischemia, which technique should be considered the gold standard during ALI caused by stent thrombosis is still uncertain. Data from randomized studies suggest that catheter directed-thrombolysis is associated to reduced mortality rate and to a less complex surgical procedure when compared to surgical revascularization, despite a higher rate of recurrent ischemia and risk of amputation [12-15]. Several non-randomized studies validated the role of combined percutaneous endovascular approach including mechanical thrombus aspiration, catheter thrombolysis, and percuta-
neous transluminal angioplasty to treat acute and sub-acute occlusions of native leg arteries [16]. Thus, percutaneous thrombectomy devices represent a non-surgical alternative for the treatment of ALI and they may be used together with fibrinolysis to reduce time and dose of the fibrinolytic agent or as a stand-alone procedure without using pharmacologic thrombolytic agents [17, 18]. However, no data regarding their use for late thrombus removal after peripheral stent implantation are available.

Here we describe, for the first time, the efficacy of percutaneous thrombectomy for treating acute peripheral nitinol-stent thrombosis, suggesting its use not only in native arteries or aortic-coronary grafts, but also in long-stented segment of SFA. The (AngioJet®) rheolytic thrombectomy (AngioJet®) system has shown promising results in terms of safety and efficacy to open thrombotic coronary arteries in acute myocardial infarction, despite it has never been tested in randomized trials. During ALI, the rheolytic thrombectomy system has been shown to be useful together with thrombolytic therapy in order to re-establish blood flow into the native lower extremity arteries [18]. Currently, it is not clear to what extent this technique is applicable for peripheral stent thrombosis. Our reports indicate the efficacy of this mechanical thrombus aspiration system in SFA stent or PTFE prosthesis thrombosis. Indeed, AngioJet® rheolytic thrombectomy has been effective in restoring immediate blood flow in stented segment, supporting an efficacy and safety profile of the Possis AngioJet catheter and its use in patients with peripheral acute stent-thrombosis. Thus, these cases suggest that AngioJet® rheolytic thrombectomy might represent a novel effective strategy in the percutaneous treatment of stent or graft thrombosis determining ALI.

**Riassunto**

L’interruzione della terapia antiaggregante, determinante trombosi degli stent in nitinol o dei by-pass femoro-poplitici in politetrafluoroetilene (PTFE) della femorale superficiale, costituisce una problematica emergente nella gestione dei pazienti con ischemia acuta dell’arto (ALI), che necessitano un intervento immediato. Ad oggi, infatti, non esiste alcuna strategia codificata per la gestione di queste complicanze. Descriviamo di seguito due casi di ALI derivanti dalla interruzione della terapia antiaggregante con la conseguente trombosi dello stent in nitinol, ed un caso di ALI conseguente alla trombosi di un by-pass in PTFE; in tutte le circostanze il salvataggio d’arto è stato ottenuto rimuovendo il trombo attraverso trombectomia percutanea di tipo reolitico (AngioJet® rheolytic thrombectomy) e attuando una successiva angioplastica transluminale percutanea (PTA). In tutti i casi, è stato ottenuto un ottimo risultato angiografico. Ad oggi, i casi da noi descritti sono i primi in cui è stato usato l’AngioJet® nel trattamento della ALI.
dovuta a trombosi dello stent o del graft. Nell’insieme, questi casi suggeriscono che la thrombectomia percutanea di tipo reolitico con AngioJet® costituisce una valida alternativa nel trattamento della trombosi di stent o graft, determinanti ALI.

ABBREVIATIONS LIST

ASA: Acetylsalicylic Acid
PAD: peripheral artery disease
SFA: superficial femoral arteries
ALI: acute limb ischaemia
RT: Rheolytic thrombectomy
PTCA: percutaneous transluminal coronary angioplasty
PTA: percutaneous transluminal angioplasty
PTFE, polytetrafluoroethylene

References