# Superselective Embolization in Posttraumatic Priapism with Glubran 2 Acrylic Glue

Roberto Gandini, Alessio Spinelli, Daniel Konda, Carlo Andrea Reale, Sebastiano Fabiano, Vincenzo Pipitone, Giovanni Simonetti

Department of Diagnostic Imaging and Interventional Radiology, University of "Tor Vergata", V.le Oxford 81, 00133, Rome, Italy

## Abstract

Two patients with posttraumatic priapism underwent transcatheter embolization using microcoils, resulting in temporary penile detumescence and an apparent resolution of the artero-venous fistula. In both cases, priapism recurred 24 hours after the procedure and was successfully treated through selective transcatheter embolization of the nidus using acrylic glue (Glubran 2). The patients showed complete recovery of sexual activity within 30 days from the procedure and persistent exclusion of the artero-venous fistula after a 12-month follow-up.

**Key words:** Priapism—Posttraumatic—Fistula—Embolization— Glubran 2—Coils

Priapism is defined as a prolonged penile erection not related to sexual arousal, often causing pain [1]. There are two types of priapism: the more frequent low-flow priapism associated with a veno-occlusive pathogenesis and the less frequent high-flow posttraumatic priapism. Low-flow priapism is a medical emergency due to ischemic damage of the corpora cavernosa causing fibrosis and impotence [2]. High-flow priapism is the consequence of direct perineal or penile trauma with laceration of a cavernosal artery, causing direct blood flow into the cavernosal sinusoids with exclusion of the high- resistance helicine arterioles [1]. The late onset of posttraumatic priapism is due to the early formation of a blood clot at the site of the arterial lesion. Subsequent nocturnal penile tumescences and the action of physiological anticoagulant agents cause mobilization of the clot itself establishing the artero-venous fistula (AVF) and resulting in the prolonged penile erection. We present two patients with posttraumatic priapism who underwent embolization with microcoils, with recurrence of the AVF after 24 h, and were definitely treated with superselective embolization using acrylic glue (Glubran 2).

## Case Report

### Case 1

As a consequence of a snowboard accident, a 26-year-old man suffered perineal trauma developing a painful penile swelling. After a few days the symptoms resolved. Two months later, the patient developed painless penile erection in the absence of hematoma. The physical examination demonstrated the presence of fully tensed corpora cavernosa. A Color-Doppler US examination revealed the presence of a right bulbo-cavernosal fistula with increased flow velocity within the only feeding branch (Fig. 1A).

The right internal pudendal artery was catheterized left transfemorally with a 4 Fr introducer sheath (Radiofocus; Terumo Japan) and 4 Fr Cobra C3 diagnostic catheter (Radiofocus; Terumo Japan). The diagnostic angiography revealed the presence of an artero-venous nidus in correspondence with the right corpus cavernosus due to a right bulbo-cavernosal fistula with a single supplying vascular branch. The superselective catheterization of the feeding branch was attempted, though the microcatheter (SP; Terumo Japan) could not be advanced through the right bulbourethral artery because of an acute angulation of the vessel. The selective left internal pudendal arteriography performed by right transfemoral approach using the above materials confirmed the presence of the AVF and was accessible for catheterization (Fig. 1B). A microcatheter (SP; Terumo Japan) was introduced into the diagnostic catheter and advanced through the left bulbourethral artery into the feeding branch (Fig. 1C). The branch was embolized using a 5 mm  $\times$  0.018 inch straight microcoil (Target; Boston Scientific). The immediate postprocedural bilateral selective internal pudendal arteriography showed complete resolution of the AVF with optimal flow within the cavernous, bulbourethral and dorsal penile arteries bilaterally, in addition to the absence of any other feeding branches (Fig. 1D). A second angiography 30 minutes later with a Color-Doppler US performed 4 hours after the procedure confirmed the resolution of the AFV (Fig. 2A).

The patient developed a gradual penile detumescence, however, 24 hours after the procedure a frank priapism was observable. The Color-Doppler US revealed the presence of floating thrombus within the nidus and an intermittent blood flow from the feeding branch (Fig. 2B). The DSA, repeated by the right transfemoral approach, revealed the recurrence of the AVF with patency of the previously embolized supplying branch, despite the correct positioning of the microcoil (Fig. 2C).

The supplying branch was superselectively catheterized using a microcatheter (SP; Terumo Japan) which, however, could not be advanced distally to the microcoil. A small amount of contrast agent was thereby gently injected to confirm the presence of anterograde flow and the optimal volume of acrylic glue required for embolization of the nidus. The nidus was then embolized by injecting 0.10 ml of Glubran 2 (GEM Srl, Viareggio, Italy) acrylic glue mixed with 0.10 ml of lipiodol (1:1 ratio) to enable fluoroscopic visualization (Fig. 2D). The bilateral selective internal pudendal arteriographic control carried out immediately and 30 minutes after the procedure confirmed the complete devascularization and thrombosis of the nidus with physiological blood flow within the cavernous, bulbourethral and dorsal penile arteries bilaterally (Fig. 2E). After the procedure the patient developed a gradual penile detumescence with complete resolution of priapism within 3 days. Control Color- Doppler US examinations performed 1, 5, 30 days, and 6 and 12 months after the procedure confirmed the absence of the AVF. Sexual activity was completely recovered after 30 days.

Correspondence to: Daniel Konda; email: danielkonda@yahoo.com



Fig. 1. A) Color-Doppler US showing the presence of a right bulbo-cavernosal fistula. B) DSA of the left internal pudendal artery confirming the presence of an artero-venous nidus in correspondence with the right corpus cavernosus due to a bulbo-cavernosal fistula with a single supplying vascular branch. C) Superselective catheterization and embolization of the feeding branch using a 5 mm  $\times$  0.018 inch straight microcoil (Target; Boston Scientific). D) Postprocedural selective left internal pudendal arteriography showing the complete resolution of the AVF with optimal flow within the left cavernous, bulbourethral and dorsal penile artery.



Fig. 2. A) Color-Doppler US performed immediately after coil embolization demonstrating the complete resolution of the AVF. B) Color-Doppler US performed 24 h after the procedure revealing the presence of a floating thrombus within the nidus and an intermittent blood flow from the feeding branch. C) Selective left internal pudendal arteriography revealing the recurrence of the AVF with patency of the previously embolized supplying branch. D) Superselective catheterization of the feeding branch and embolization of the nidus with 0.10 ml of Glubran 2 acrylic glue mixed with 0.10 ml of lipiodol. E) Postprocedural selective left internal pudendal arteriography confirming the complete devascularization and thrombosis of the nidus with normal visualization of the left cavernous, bulbourethral and dorsal penile artery.



Fig. 3. A) Color-Doppler US revealing a left bulbo-cavernosal AVF with a single feeding branch. B) Selective left internal pudendal arteriography confirming the existence of a left bulbo-cavernosal AFV supplied by a single feeding branch. (C) Postprocedural selective left internal pudendal arteriography after embolization with 0.08 ml of Glubran 2 acrylic glue mixed with 0.08 ml of lipiodol, performed 24 hours after coil embolization confirming the complete devascularization and thrombosis of the nidus.

## Case 2

A 24-year-old man developed a painful penile swelling after a ski accident, causing perineal trauma. Five days after the trauma, the patient experienced a gradual increase in penile tumescence, resulting in an enduring painless penile erection in the absence of hematoma. Physical examination showed fully tensed corpora cavernosa and Color- Doppler US revealed a left bulbo-cavernosal AVF with single feeding branch (Fig. 3A). A diagnostic angiography performed by right transfemoral approach with catheterization of the left internal pudendal artery using a 4 Fr introducer sheath (Terumo; Japan) and 4 Fr Cobra C3 diagnostic catheter (Radiofocus; Terumo Japan) showed the existence of a left bulbo-cavernosal AFV (Fig. 3B). The nidus presented a single supplying branch.

The feeding branch was superselectively catheterized with a microcatheter (SP; Terumo Japan) and embolized with a  $3 \times 2$  mm spiral microcoil (Tornado; Cook Inc. Canada). The immediate postprocedural bilateral selective internal pudendal arteriography demonstrated the resolution of the AVF with physiological blood flow within the cavernous, bulbourethral and dorsal penile arteries bilaterally and no contralateral feeding branches. A second angiography performed 30 minutes later and a Color-Doppler US performed 4 hours after the procedure confirmed the resolution of the AVF.

The patient underwent a partial penile detumescence with recurrence of penile erection after 24 hours. Color-Doppler US revealed patency of the feeding branch with turbulent flow within the AFV. A selective left internal pudendal arteriography performed by right transfemoral approach confirmed the persistence of the AFV with flow within the supplying vessel. No other ipsilateral or contralateral feeding branches were detected.

The nidus was embolized with a mixture of 0.08 ml of Glubran 2 (GEM Srl, Viareggio, Italy) acrylic glue and 0.08 ml of lipiodol (1:1 ratio) injected with a microcatheter (SP; Terumo Japan) advanced through the diagnostic catheter to the microcoil in the feeding branch. The bilateral selective internal pudendal arteriography performed immediately and 30 minutes after the procedure confirmed the complete devascularization and thrombosis of the nidus with physiological blood flow within the cavernous, bulbourethral and dorsal penile arteries bilaterally (Fig. 3C). The patient underwent gradual penile detumescence with complete resolution of the

symptoms within 5 days. The Color-Doppler US examinations performed 1 day, 30 days and 6 and 12 months after the procedure confirmed the absence of the AFV. Sexual activity was completely recovered after 35 days.

## Discussion

Posttraumatic priapism is a fairly uncommon cause of priapism, usually caused by direct perineal or penile trauma. The occurrence of a bulbo-cavernosal fistula causes a high arterial flow within the lacunar spaces, bypassing the high-resistance helicine arteries [3, 4]. Diagnosis is usually suspected on the basis of the history of trauma and physical examination. When such a suspicion exists, confirmation may be obtained through arterial blood aspiration from the corpora cavernosa. On the other hand, a precise determination of the number, location and lateralization of the arterial lacunar fistulas may be obtained using Doppler-US. In addition, in case of high-flow priapism, this noninvasive exam can show the increased flow velocities in the cavernosal artery that are above the characteristic velocities of a normal erection [5].

Posttraumatic priapism is treated with conservative therapy including perineal ice-pack compression, intracavernosal  $\alpha$ -agonist agents or methylene blue injections. If conservative therapy fails an invasive alternative such as surgical internal pudendal artery ligation or selective transcatheter embolization may be employed. Surgery is, however, associated with a high risk of impotence or partial erection in the long term [6, 7].

The first successful radiological treatment of high-flow priapisms was reported in 1977 by Wear et al. [8-11] in a 26-year old man who underwent selective embolization using autologous blood clot. This technique has proven to be safe, feasible and effective in most cases, however, some authors report the occurrence of thrombolysis followed by recurrent priapism, requiring reintervention [12, 13]. An advantage of this technique is represented by the reconstitution of normal arterial flow after slow physiological dissolution of the clot, reducing the risk of impotence [14]. Nevertheless, cases of permanent arterial occlusion and impotence using autologous clot are reported in the literature [15]. When complete procedural success is achieved, erectile recovery is obtained only after several months [8, 12, 13].

Gelfoam has been shown to be effective for treating high-flow priapism, however, it enables the reconstitution of normal blood flow in a manner similar to autologous clot and shares with the latter the drawback of not being radiopaque, and thereby not allowing precise control of embolization [9, 16–18]. Gelfoam may be considered useful in the treatment of high-flow bilateral priapism where recanalization of the embolized artery is important since no collateral blood flow is available from the contralateral side [19].

Embolization using straight and spiral microcoils has been reported only in a few cases [12, 18, 20, 21]. The disadvantages of this technique are its permanent nature and the usual need of more than one microcoil for complete occlusion of the branch. The use of microcoils in small caliber vessels frequently determines spasm with an apparent occlusion of the feeding branch at the immediate post-procedural DSA control. The subsequent resolution of this spasm favors the recurrence of the AVF which can require the use of several microcoils to obtain the permanent occlusion of the feeding branch. In fact, during the physiological nocturnal penile tumescence, the vasodilation of the embolized branch, in addition to the endogenous lysis of the thrombus, may cause recurrence of priapism. Furthermore, in case of additional branches supplying the AVF, several microcoils may be required for optimal treatment. In addition, even when several feeding branches exist, the preliminary angiography frequently demonstrates the presence of a single branch due to blood flow predominance in the latter.

N-butyl-Cyanoacrylate (NBCA) is a monomer acrylic glue which rapidly polymerizes when in contact with ionic media such as blood and causes a permanent occlusion. To avoid adherence to the tissue of the thin catheters required for the superselective embolization, NBCA has to be injected through a catheter washed with a 5% dextrose solution and the catheter has to be withdrawn promptly after injection [1, 22]. The technique requires considerable expertise; there is also the risk of undesired embolizations. Moreover, NBCA polymerizes with an exothermic reaction, causing pain to the patient.

Glubran 2 (GEM Srl, Viareggio, Italy) is an acrylic glue bearing a CE mark authorized for surgical and endovascular use in neuroradiology. The comonomer of Glubran 2 comprises a monomer of NBCA and a monomer of MS (owned by GEM Srl). MS allows the monomer of NBCA to polymerize with a lower exothermic reaction (45°C) and a slightly longer polymerization time [23]. Compared to the monomer NBCA, the Glubran 2 causes less pain to the patient and is associated with a lower risk of adherence of the catheter to the tissue, hence showing a greater ease of use. Differently, acrylic glues, once deposited into the nidus, determine its permanent occlusion and prevent its replenishment through feeding branches.

In the two cases presented here, the microcoil caused an initial vascular spasm of the small feeding branch which was erroneously interpreted as the successful embolization of the latter. In both patients, the complete occlusion of the feeding vessel observed at the immediate postprocedural angiography was confirmed by a second angiography performed 30 minutes after embolization and by Color- Doppler US performed 4 hours after the procedure. The subsequent recanalization of the feeding branch, which occurred 24 hours after the procedure, might have been determined by the

physiological nocturnal penile tumescence, causing vasodilation of the embolized vessel along with the endogenous lysis of the thrombus.

During microcoil embolization, the placement of additional microcoils could have favored the durability of the occlusion, however, it would have also exposed the patient to an increased risk of nontarget artery embolization. In fact, additional microcoils could have only been placed more proximally to the first, increasing the probability of dislocation from the target site. To avoid possible further recurrences of the AFV and reduce the risk of undesired embolizations using microcoils, in both cases we preferred the direct embolization of the nidus using Glubran 2.

Although, according to the literature, the use of permanent embolic materials should be reserved for cases with recurrent priapism after an initial success with absorbable embolic agents [24], we believe that primary embolization using acrylic glues such as Glubran 2 is safe, feasible and more efficient. In fact, in case of absorbable embolic materials the recurrence of priapism is frequent due to recanalization of the AVF following reabsorbption, and a long period is required before a complete recovery of the erectile and sexual function is obtained. With microcoils, on the other hand, postprocedural penile detumescence may be associated with vascular spasm which can be erroneously interpreted as a procedural success. Furthermore, when the AVF is supplied by more than one blood vessel, the placement of several microcoils may be needed for complete resolution of the disease, with a consequent higher risk of non-target artery embolization. Differently, acrylic glues may also be deposited directly into the angiographic nidus corresponding to the AVF, thus at once permanently excluding the pathological communication regardless of the number of feeding branches.

Compared to other acrylic agents, Glubran 2 presents a greater ease of use, causes less discomfort to the patient and is the only acrylic glue bearing a CE mark. Glubran 2 enables the direct embolization of the arterovenous nidus and should be regarded as a valid alternative in the treatment of posttraumatic priapism; however, expertise is required to prevent undesired embolizations.

#### References

- Numan F, Çakirer S, Işlak C, Öğüt G, Kadioğlu A, Çayan S, Tellaloğlu S (1996) Posttraumatic high-flow priapism treated by N-butyl-cyanoacrylate embolization. Cardiovasc Intervent Radiol 19:278–280
- Spycher MA, Hauri D (1986) The ultrastructure of the erectile tissue in priapism. J Urol 135:142–147
- Witt MA, Goldstein I, De Tejada IS, Greenfield A, Krane RJ (1990) Traumatic laceration of intracavernosal arteries: the pathophysiology of nonischemic, high flow, arterial priapism. J Urol 143:129–134
- Walker TG, Crant PW, Goldstein I, Krane RJ, Greenfield AJ (1990) "High-flow" priapism: treatment with superselective transcatheter embolization. Radiology 174:1053–1056
- Schwartz AN, Wang KY, Mack LA, Lowe M, Berger RE, Cyr DR, Feldman M (1989) Evaluation of normal erectile function with color flow Doppler sonography. Am J Roentgenol 153:1155–1160
- Burt FB, Schirmer HK, Scott WW (1960) A new concept in the management of priapism. J Urol 83:60–61
- Brock G, Breza J, Lue TF, Tanagho EA (1993) High flow priapism: a spectrum of disease. J Urol 150:968–971
- Wear JB Jr., Crummy AB, Munson BO (1977) A new approach to the treatment of priapism. J Urol 117:252–254
- Miller SF, Chait PG, Burrows PE, Steckler RE, Khoury AE, McLorie GA, Connolly BL, Pereira JK (1995) Post-traumatic arterial priapism in children: management with embolization. Radiology 196:59–62
- De Stefani S, Capone M, Carmignani G (1993) Treatment of posttraumatic priapism by means of autologous clot embolization: a case report. Eur Urol 23:506–508
- 11. Kim SC, Park SH, Yang SH (1996) Treatment of posttraumatic chronic

high-flow priapisms by superselective embolization of cavernous artery with autologous clot. J Trauma 40:462-465

- Kress O, Heidenreich A, Klose KJ, Wagner H-J, Alfke H (2002) Superselective embolization with coils in high-flow priapism. Cardiovasc Intervent Radiol 25:326–329
- Bastuba MD, Saenz de Tejada I, Dinlenc CZ, Sarazen A, Krane RJ, Goldstein I (1994) Arterial priapism: diagnosis, treatment and longterm follow-up. J Urol 151:1231–1237
- Warren BA (1964) Fibrolytic activity of vascular endothelium. Br Med Bull 20:210
- Hakim LS, Kulaksizoglu H, Mulligan R, Greenfield A, Goldstein I (1996) Evolving concepts in the diagnosis and treatment of arterial high flow priapism. J Urol 155:541–548
- Ji MX, He NS, Wang P, Chen G (1994) Use of selective embolization of the bilateral cavernous arteries for post-traumatic arterial priapism. J Urol 151:1641–1642
- Cohen GS, Braunstein L, Ball DS, Roberto PJ, Reich J, Hanno P (1996) Selective arterial embolization of idiopathic priapism. Cardiovasc Intervent Radiol 19:47–49
- 18. Lanzinger M, Beckmann CF, Cossi A, Roth RA (1996) Selective

embolization of bilateral arterial cavernous fistulas for posttraumatic penile arterial priapism. Cardiovasc Intervent Radiol 19:281–284

- Ming-Xian J, Neng-Shu H, Ping W, Gui C (1994) Use of selective embolization of the bilateral cavernous arteries for posttraumatic arterial priapism. J Urol 151:1641–1642
- Kerlan RK, Gordon RL, LaBerge JM, Ring EJ (1998) Superselective microcoil embolization in the management of high-flow priapism. JVR 9:85–89
- Callewaert P, Stockx L, Bogaert G, Baert L (1998) Post-traumatic high-flow priapism in a 6-year-old boy: management by percutaneous placement of bilateral vascular coils. Urology 52:1334–1347
- Alvarez Gonzalez E, Pamplona M, Rodriguez A, Garcia-Hidalgo E, Nunez V, Leiva O (1994) High flow priapism after blunt perineal trauma: resolution with bucrylate embolization. J Urol 151:426–428
- Leonardi M, Barbara C, Simonetti L, Giardino R, Nicoli Aldini N, Fini M, Martini L, Masetti L, Joechler M, Roncaroli F (2002) Glubran 2: a new acrylic glue for neuroradiological endovascular use. Experimental study on animals. Intervent Neuroradiol 8:245–250
- Kessel D, Robertson I, McWilliams RG (1997) Letter to the editor. Re: Embolization of posttraumatic high flow priapism. Cardiovasc Intervent Radiol 20:237–238