Role of cyanoacrylate glue and its efficacy in the treatment of varicose vein

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Abstract

To evaluate the role of cyanoacrylate glue therapy and its efficacy and technical feasibility in the management of primary varicose vein due to great saphenous vein incompetency with or without incompetent perforators. This is a prospective study done in 30 patients who presented with varicose veins due to great saphenous vein reflux with or without incompetent perforators and with or without venous ulcer. All patients were subjected to cyanoacrylate glue therapy in great saphenous vein followed by foam sclerotherapy for any incompetent were done at months 1, 3, 6 and 12. The study’s Primary end point was closure of the target vein at 12 month as assessed by duplex ultrasound. Technical success rate of procedure was 100%. Of all the 30 patients Saphenous vein closure rate was 96.6% at one year. There was no femoral venous extension of cyanoacrylate in any of the patients. Significant improvement was found in reduction in pain and in diameter and tortousity of saphenous vein. Ulcer healing rate was 100%. Symptoms and quality of life improved in all the patients. There was no significant adverse effects noted during the procedure and in follow up. In patients with incompetent GSV, treatment with cyanoacrylate glue therapy results in high occlusion rates. Cyanoacrylate adhesive embolization and sclerotherapy is safe, easy, cost effective, minimally invasive, non surgical outpatient procedure for the treatment of primary varicose veins with 100% patient acceptance and very less morbidity compared to surgery.

Keywords: Ultrasonographic, fatty liver, cyanoacrylate glue therapy, technical feasibility

Introduction

Varicose veins are veins that have become enlarged and twisted. When veins become varicose, the leaflets of the valves no longer meet properly, and the valves do not work (Valvular incompetence). This allows blood to flow backwards and they enlarge even more. Varicose veins are most common in the superficial veins of the legs, which are subject to high pressure when standing. Besides being a cosmetic problem, varicose veins can be painful, especially when standing. Severe long-standing varicose veins can lead to leg swelling, venous eczema, skin thickening (lipodermatosclerosis) and ulceration. Although life-threatening complications are uncommon, varicose veins may be confused with deep vein thrombosis, which may be life-threatening [1].

The traditional surgical treatment has been vein stripping to remove the affected veins. Newer, less invasive treatments which seal the main leaking vein are available. Alternative techniques, such as ultrasound-guided foam sclerotherapy, radiofrequency ablation and endovenous laser treatment, are available as well. Because most of the blood in the legs is returned by the deep veins, the superficial veins, which return only about 10% of the total blood of the legs, can usually be removed or ablated without serious harm [2, 3].

In the present Cyanoacrylate Adhesive Embolization and Sclerotherapy (CAES) study, we explore the feasibility and efficacy of an inexpensive treatment combining adhesive embolization of GSV and perforators with the commonly available n-butyl cyanoacrylate glue (which is used as topical skin adhesive or for endovascular embolization Of arteriovenous malformations and vascular tumors 28, 29) and sclerotherapy for the treatment of primary varicose veins.
Aims and Objective
To evaluate the role of cyanoacrylate glue therapy and its efficacy in the management of primary varicose vein due to great saphenous vein incompetency with or without incompetent perforators and with or without venous ulcer.

Material and Methods
This prospective study was conducted as collaboration between the department of Radio diagnosis and department of Surgery. The study period was from July 2018 till March 2018. Consecutive patients who had presented to the Surgery department with complaints of dilated and tortuous saphenous veins with or without venous ulcer.

Inclusion criteria
1. Males and non-pregnant females of any age over 18 years.
2. Primary varicose veins due to great saphenous vein reflux with or without perforator reflux.
3. Good mental health to understand and consent for the procedure with investigative nature.
4. Patients willing for follow-up.

Exclusion criteria
1. Great saphenous vein diameter more than 8 mm.
2. Recurrent varicose veins.
3. Present or previous DVT.
4. Lower limb arterial disease.
5. Hypercoagulable status.

All patients who were referred to our Radiology department with symptoms of lower limb primary superficial venous insufficiency were subjected to physical examination and a thorough Doppler study to map the sources of reflux in pathological veins. Patients satisfying inclusion and exclusion criteria were considered for the study.

Procedure
These patients underwent routine colour duplex ultrasound using ALOKA Pro sound ALPHA-6, linear array probe with frequency 10 MHz to confirm the venous insufficiency and to know the extent of the venous disease. The procedure was explained to the patient in their own languages and their consent for participating in the study was taken. Procedures was performend in supine position with maintaining proper aseptic precautions. Cyanoacrylate adhesive embolization of GSV in the thigh and leg was performed followed by foam sclerotherapy of any incompetent perforators in the legs whenever required. The principle of adhesive embolization was to inject cyanoacrylate glue into dilated saphenous vein or perforators followed immediately by external compression to cause collapse and obliteration of the vessel lumen, with special attention to prevent any spread of glue into the deep veins. (BERRY). The glue used in our study was a very low-viscosity, rapidly polymerizing fluid N butyl cyanoacrylate glue (RECKSEAL TM). Under USG guidance a 20G, venflow needle was percutaneously placed within the dilated GSV in the lower thigh and leg having the straight course for ease of puncture after applying the tourniquet at saphenofemoral junction, the intravascular position of the needle was confirmed by movement of the needle within the vein. The dextrose-containing syringe was exchanged quickly for a 1-ml syringe loaded with 0.5 ml of cyanoacrylate glue (Reckseal or Endocryl, Samarth Pharma Pvt. Ltd) and 0.15 ml of air (0.10 ml of air is taken to fill the intraluminal dead space of the venflow needle and push the full dose of cyanoacrylate glue into the cannulated vein when injected). The glue was injected using a fairly quick and continuous push of the plunger as the needle was withdrawn and firm immediate external compression was applied for 30 s. This resulted in intravenous spread of the glue, and collapse – adhesive occlusion of the vein. Deposited glue in the vein appeared echogenic and if dense, cast a post-acoustic shadow. Short unoccluded segments between treated segments were not treated unless there was an associated visible perforator in ultrasound entering the segment. After occlusive embolization of GSV in the thigh, perforators were treated if they were found to be incompetent – defined as reflux time of more than 0.35 s and diameter of more than 3 mm at the deep fascial level. Perforators smaller than 3 mm showing long reflux times were also treated, if they were located close to a venous ulcer, or communicate directly with symptomatic dilated superficial veins. For dilated and tortuous vein and incompetent perforators in legs, foam sclerotherapy was done using Fagan’s method of foam sclerotherapy. The Tessari micro bubble method of UGFS sclerotherapy was used. Polidocanol was the detergent sclerosant used in the procedure (pharmaceuticals – injection asklero 3% / 2ml, manufacturer Samarth pharma private limited. Price-Rs 85/-). The detergent sclerosant (Polidocanol) was mixed with room air to form the foam. This was done using the three way tap switch and two 10 ml disposable syringes. A 3:1 ml room air to sclerosant ratio was mixed with approximately 18-20 rapid exchanges which causes the turbulent flow via three way tap switch to produce micro bubble foam (Tessari micro bubble method). Typical ratios described of air to sclerosant are 3:1 to 5:1. The foam was stable for 1-2 minutes and remixing was done if reduced to froth. The maximum dose of detergent sclerosant, polidocanol that can be used per session is 12 mL of 1% (4ml of 3%) . After the procedure, roller bandage was applied firmly from groin to foot and the patients were mobilized. Patients were instructed to apply the bandage for three -four weeks. If the bandage get loosened then reapplied it firmly again. Patients had given ecosprine and clopidogrel prophylactically. Any pain in the postoperative period was managed with oral analgesics and cold fomentation.

Follow-up
The patients were asked to revisit us at 1, 3, 6, 9 and 12 months. Clinical assessment, and ultrasound were performed on patients during follow-up.

Fig 1: USG image showing thrombosed GSV with echogenic glue within its lumen.
Incompetent SFJ with incompetent perforators. Successful outcomers was found in 29 patients with presence of echogenic glue within the vessels at 12 month follow up. Failure was found in 1 patient probably patient was on antiplatelet therapy for cardiac pathology. No serious complication was found in successive follow up apart from minor pain at the injection site which was managed with nonsteroidal anti-inflammatory medicines.

Discussion

Various surgical ie vein stripping to remove the affected veins and non surgical treatment options such as ultrasound-guided foam sclerotherapy, radiofrequency ablation and endovenous laser treatment. The intravascular use of NBCA also known as superglue has a history of >20 years in arteriovenous malformations and venous applications in other parts of the body. Various Study results have revealed that on contact with intravascular tissue, NBCA rapidly undergoes a polymerization reaction and begins to solidify on coming in contact with the anions of the blood [8, 9]. This polymerization creates an acute inflammatory effect over the vein wall which progresses to granulomatous inflammation; simultaneous gradual resorption of the polymer takes place, resulting in the transformation of a patent vein to an occluded fibrotic cord [9, 10].

The glue used in this study is low viscosity liquid with quick polymerization quality hence the injected venflow withdrawl- external compression had to be fairly quick. Since the glue polymerize very quickly in coming in contact with the blood, the deliver of the glue could not be done with the catheter as the risk of catheter lumen blockage. For the same reason cyanoacrylate glue was injected segmentally at multiple sites directly using the disposable venflow (18G) within the vein percutaneously. Because of the low viscosity and unpredictability of extent of spread of glue, to prevent the catastrophic complication on DVT, tight tourniquet was used at SFJ and site of injecting the glue should be atleast 8cm from the SFJ. The clinical and ultrasonographic findings during follow-up in our patients correlate with the pathology of cyanoacrylate embolization where veins with deposited palpably hard cyanoacrylate within them cause a transient thrombophlebitic reaction (corresponding to acute inflammatory response) and gradually reduce to a slightly firm fibrotic cord (corresponding to gradual resorption of cyanoacrylate and scarring).

Results from this study confirm that CAE is safe and highly effective for the treatment of CVD. The study showed that occlusion of the target vein at 3 months was high and 12 month occlusion rate in our study found to be 96.5% and similar to that observed in a prior single-arm CAE study (95% in a small Europaen study [96%]) feasibility study [11] and in a prospective CAE multicenter [12]. No serious procedural and post procedure adverse effects were noted. Although some cases of phlebitis of the treated GSV had occurred, were mild, transient, and successfully treated with the over counter nonsteroidal anti-inflammatory medications. Prophylactically antibiotics were given to prevent any infection post procedural.

Primary advantage of this treatment was low cost and easily availability of cyanoacrylate glue required for CAES. Other advantages of CAES include outpatient nature of the procedure, no requirement of tumescent anaesthesia or immobilization, efficiency comparable to multiple published
studies using proprietary cyanoacrylate, clinical improvement in all patients, very good ulcer healing rate and low complication rate. The primary disadvantage is the unpredictability of the length of glue extension and a possibility of catastrophic occlusion of common femoral vein (although not seen in any patient in our study), because of which it should be performed by interventionists with good prior experience with usage of cyanoacrylate glue. A minor disadvantage of this procedure is that it requires multiple skin punctures compared to single puncture required to treat the whole length of saphenous vein when proprietary cyanoacrylate glue (Sapheon venaseal or Variclose) is used.

Conclusion
CAES is a minimally invasive alternative treatment option for the treatment of primary varicose veins. In current study, CAES was found to be efficacious, feasible easily affordable, cost effective treatment option especially for poor patients which can be performed in an outpatient basis. Result of study was found to be comparable with the various other treatment options.

References
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