

Journal of Clinical Imaging Science



Original Research Vascular and Interventional Radiology

Percutaneous n-butyl cyanoacrylate glue embolization combined with arterial embolization for extracranial arteriovenous malformations

Yang-Chao Tong^{1#}, Yang Zhao^{2#}, Hai-Peng He², Ren Lin², Jun-Bing Lv², Yi-Bo Zhang², Hui-Ning Chen², Heng-Hui Yin², Jie-Sheng Qian^{2*}

¹Department of Interventional Vascular, The Third Affiliated Hospital of Sun Yat-sen University, Guangzhou, ²Department of Vascular Surgery, The Third Affiliated Hospital of Sun Yat-sen University, Guangzhou, China. [#]These author contributed equally to this work.

*Corresponding author: Jie-Sheng Qian, Department of Vascular Surgery, The Third Affiliated Hospital of Sun Yat-Sen University, Guangzhou, China.

qjsyf@163.com

Received: 02 March 2025 Accepted: 24 April 2025 Published: 29 May 2025

DOI 10.25259/JCIS_33_2025

Quick Response Code:



ABSTRACT

Objectives: Congenital arteriovenous malformations (AVMs) are characterized by abnormal connections between arteries and veins, often presenting challenges in treatment due to their complex vascular structure. Endovascular therapies, including embolization techniques, have become integral in managing AVMs, yet optimal treatment strategies remain under investigation. This retrospective study aimed to evaluate the safety and efficacy of percutaneous glue embolization of n-butyl cyanoacrylate (nBCA) combined with arterial embolization in treating extracranial AVMs.

Material and Methods: This retrospective study included 11 patients with extracranial AVMs who underwent percutaneous injection of nBCA glue embolization combined with arterial embolization at our institution between May 2015 and October 2023. Angiographic classification was performed using the Cho-Do vascular imaging system and the Schobinger classification system to categorize the AVMs. Treatment efficacy was assessed based on the percentage of vessel occlusion observed in angiography or imaging studies. The occurrence of major and minor complications was also evaluated.

Results: Eleven patients received 16 treatments, involving a combination of arterial and percutaneous embolization techniques. Five cases achieved complete recovery, while four cases showed significant improvement, resulting in an overall treatment success rate of 81.8%. Favorable outcomes were observed in terms of symptom alleviation and lesion occlusion. Adverse events were minimal, with transient pain and edema being the most common post-operative complication occurred.

Conclusion: Percutaneous glue embolization combined with arterial embolization proves to be a safe and effective treatment modality for extracranial AVMs, with acceptable rates of complications and favorable treatment outcomes.

Keywords: Arteriovenous malformations, Embolization, n-butyl cyanoacrylate, Percutaneous

INTRODUCTION

Congenital arteriovenous malformations (AVMs) are a form of high-flow vascular malformations characterized by underdeveloped arteries and veins directly interconnected through a convoluted vascular mass termed the "nidus."^[1] AVMs can be classified into central and peripheral types, with the head and neck representing the most common sites for peripheral AVMs,^[2] followed

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by the limbs. Symptoms vary depending on the location and size of the lesion, including pain, swelling, skin discoloration, pulsatile masses, ulceration, bleeding, and necrosis.^[3] In massive AVMs, direct arterial blood flow into the venous system can lead to increased cardiac pre-load and subsequent heart failure.^[4] According to the Schobinger classification, AVMs can be divided into four stages based on clinical symptoms: Quiescence (Stage I), expansion (Stage II), destruction (Stage III), and decompensation (Stage IV).^[5] Advancement to higher stages indicates more severe clinical symptoms and typically necessitates more intricate clinical management.

Resecting AVMs surgically presents challenges, including achieving complete excision for larger lesions, the risk of recurrence, intraoperative hemorrhage, and potential damage to surrounding structures.^[6,7] Endovascular therapy employing diverse embolic agents and sclerosants has emerged as a principal treatment modality for AVMs.^[8,9] Among these, an extensively used embolic agent is absolute ethanol, which has demonstrated superior efficacy compared to other embolic materials in numerous studies.^[10,11] The complete cure rate of AVMs with ethanol embolization ranges from 40% to 88%.^[4] However, the use of absolute ethanol is associated with a high incidence of complications, with an overall complication rate ranging from 10% to 52%.[12] In addition to absolute ethanol, other commonly used embolic materials include liquid embolic agents such as n-butyl cyanoacrylate (nBCA),^[13] Onyx,^[14] particulate embolic agents like polyvinyl alcohol (PVA) embolic particles,^[15] and detachable coils.^[16]

Arterial embolization using nBCA, also known as tissue glue embolization, is frequently employed in the clinical management of AVMs.^[17] However, relying solely on nBCA may not always achieve complete occlusion of AVMs, particularly in cases involving multiple feeding arteries and draining veins,^[18] often leading to incomplete occlusion and subsequent recurrence.^[19]

When arterial access poses challenges or complete occlusion cannot be achieved via arterial approaches, percutaneous puncture approaches serve as valuable alternatives.^[20] Percutaneous injection of ethanol via catheter or needle puncture has shown promising outcomes but is associated with an increased risk of complications.^[5,21] Combining arterial embolization with nBCA and percutaneous sclerotherapy using ethyl oleate has been reported as a safe and effective treatment method with an acceptable risk profile.^[22] Similarly, PVA embolization combined with nBCA arterial embolization has shown promising results for extracranial AVMs with an acceptable rate of complications.^[17]

Percutaneous injection of nBCA has demonstrated pathological success in embolizing AVMs,^[23,24] venous

malformations,^[25] and pseudoaneurysms^[26] in various patient populations. However, clinical studies specifically focused on percutaneous nBCA glue embolization therapy remain relatively limited, particularly in China. Therefore, this study aimed to evaluate the safety and efficacy of percutaneous glue embolization of nBCA combined with arterial embolization in the treatment of extracranial AVMs.

MATERIAL AND METHODS

Study population

This was a retrospective study conducted on all patients with AVMs who received treatment involving percutaneous injection of tissue glue embolization, either with or without adjunctive arterial embolization, at our institution between May 2015 and October 2023.

All enrolled patients underwent at least one embolization treatment. Exclusion criteria included other types of vascular malformations such as vascular tumors, venous malformations, and patients who did not undergo percutaneous injection of tissue glue embolization treatment.

The diagnosis of AVMs was established through a combination of clinical history, physical examination, and imaging studies including magnetic resonance imaging, computed tomography, and duplex ultrasonography.

The data collected included patients' medical records such as demographic data, pre-operative imaging studies, details of all embolic agents and embolization approaches used during surgery, post-operative imaging studies, and followup information. This study was approved by the institutional review board of the Third Affiliated Hospital of Sun Yat-sen University (No. II2023-250-01). Written informed consent was obtained from each patient.

Angiographic classification

All patients, except one, underwent selective angiography through arterial access to acquire detailed anatomical and hemodynamic data. Prior to embolization, two interventional radiologists independently reviewed angiographic images and categorized AVMs based on Cho *et al*.'s classification,^[27] reaching a consensus. A highly experienced vascular surgeon made the final decision. The AVMs were classified into four types based on their angiographic morphology.

Procedures

All embolization procedures were conducted under local anesthesia and guided by digital subtraction angiography (DSA). Initially, diagnostic angiography was performed to confirm the diagnosis and comprehend the vascular architecture of AVMs. When the draining veins of the lesion were identifiable and relatively large, methods such as compression, tourniquet application, or ligation were employed to decelerate blood flow. Subsequently, suitable microcatheters were chosen based on angiographic images to attempt superselection of each feeding artery of the AVMs. After verifying the position with repeat angiography, appropriate embolic agents were meticulously prepared and slowly injected through the microcatheter. Injection was halted upon stagnation or retrograde flow of blood toward the microcatheter tip, followed by angiography to confirm the extent of embolization. Comprehensive angiography was performed upon achieving satisfactory embolization of all superselected feeding arteries to evaluate the efficacy of embolization.

In instances where small, tortuous feeding arteries of the AVMs were challenging to superselect via arterial access alone, a combined approach was implemented to augment treatment efficacy, involving arterial embolization followed by percutaneous embolization. Utilizing techniques like compression of superficial feeding arteries and draining veins to attenuate blood flow, a 7-French scalp needle was initially employed for percutaneous puncture of the lesion until blood was visualized. Subsequent angiography through the puncture needle confirmed its position, assessed vascular structure and size, and facilitated the preparation of tissue glue embolic agents in varied proportions based on the flow rate. Under DSA guidance, tissue glue was slowly injected until favorable outcomes were attained, followed by subsequent arterial angiography to assess treatment outcomes. Multiple sessions of arterial/percutaneous embolization were conducted for larger lesions to achieve complete nidus occlusion.

Follow-up

During hospitalization and outpatient or telephone followup, all patients were closely monitored, and occurrences of complications or changes in symptoms were recorded. Assessment of treatment efficacy was conducted 3–6 months after the final treatment session. Additional embolization or sclerotherapy was recommended if symptoms recurred or worsened post-treatment, affecting quality of life, or if residual AVMs were detected on follow-up imaging examinations.

Statistical analysis

Statistical analysis was performed using Fisher's exact test to determine the relationship between AVM staging/types and treatment efficacy. IBM Statistical Package for the Social Sciences Statistics Version 22.0 (IBM, New York, USA) was utilized for statistical analysis, with P < 0.05 being considered statistically significant.

RESULTS

Patients and demographics

Demographic and clinical characteristics of patients with AVM are presented in Table 1. A total of 11 patients meeting the inclusion criteria were included. Among them, there were 5 male and 6 female patients, with a mean age of 36.1 \pm 7.1 years (range: 12–66 years). Pain and swelling were the most common symptoms, with 5 cases classified as Schobinger II and 6 cases as Schobinger III [Table 1]. All patients had type IIIb AVMs, except for Patient 8, who had a combination of type IIIa and IIIb. AVMs were predominantly located in the limbs and head, including the fingers, forearms, lower legs, thighs, and head. Notably, the highest prevalence of AVMs was observed in the hands (including fingers and forearms).

Treatment details

The detailed treatment procedures are outlined in Table 2. Among the 11 patients, 5 achieved complete devascularization of AVM lesions. Notably, in one case [Patient 3], due to tortuous and small feeding arteries, a direct percutaneous injection of glue was performed after ligation and dilation of the draining vein. A patient (Patient 7) diagnosed with a left frontal AVM (Figures 1a-c) initially underwent the release of spring coils followed by n-butyl cyanoacrylate (NBCA) embolization via arterial access under manual compression of superficial drainage veins (Figure 1d). Subsequently, percutaneous glue injection was performed (Figure 1e). Post-embolization angiogram demonstrated complete occlusion of the AVM nidus (Figure 1f). Another patient (patient 11) diagnosed with a right forearm AVM (Figures 2a-b) underwent spring coils release via arterial access under manual compression of superficial drainage veins (Figure 2c). Subsequently, percutaneous glue injection was performed (Figure 2d). Post-embolization angiogram demonstrating complete occlusion of the AVM nidus (Figure 2e). One-month post-embolization MRI showed a reduction in the lesion size compared to previous imaging, with no visible flow voids (Figure 2f). Patient 8, diagnosed with a left thigh and left knee AVM (Figures 3a-b), underwent polyvinyl alcohol (PVA) and glue embolization via arterial access under tourniquet compression (Figure 3c). Post-embolization angiography demonstrated near-complete occlusion of the AVM nidus (Figure 3d). Subsequently, percutaneous glue injection was performed for the residual left knee AVM (Figure 3e). Final angiogram confirmed complete occlusion of the entire AVM nidus (Figure 3f).

Four patients achieved over 90% occlusion of AVM vessels. In one case (Patient 6), direct percutaneous glue injection was performed due to tortuous and small feeding arteries. In two cases (Patient 1 and 10), arterial embolization was

Table 1: Characteristics of patients with arteriovenous malformations.								
Patient ID	Gender	Age	Location	Туре	Stage	Past treatments	Symptoms	
1	Female	38	Left lower jaw	IIIb	II	None	Swelling	
2	Female	22	Right calf	IIIb	III	Embolization	Swelling, Pain	
3	Male	66	Right forearm	IIIb	II	Excision	Swelling	
4	Female	23	Left finger	IIIb	III	None	Swelling, Pain, Vascular tufts	
5	Female	26	Left hand, Left forearm	IIIb	II	2 times Embolization	Vascular tufts	
6	Male	49	Left finger	IIIb	III	Excision	Swelling, Pain	
7	Female	27	Left forehead	IIIb	III	Excision	Swelling, Pain	
8	Female	12	Left thigh, Left knee	IIIa, IIIb	II	Excision	Swelling	
9	Male	36	Forehead	IIIb	II	None	Swelling	
10	Male	42	Right forearm, Right hand	IIIb	III	2 times Excision	Swelling, Pain, Vascular tufts	
11	Male	28	Right forearm	IIIb	III	None	Swelling, Pain	

Patient ID	Number of Treatments	Treatment Pathway	Transarterial	Percutaneous	Transvenous
1	1	TA, TP	PVA, gelatin sponge, coil, glue	Glue	None
2	3	TA, TV, TP+TP+TA, TV, TP	Glue+0+PVA	Glue+polyvinyl alcohol foam+glue	Ligation+ None+Ligation
3	1	TV, P	None	Glue	Ligation
4	2	TA, TV, TP+TA	PVA+PVA	Glue+None Ligation-	
5	2	P+P	Compression +Compression	Glue+Glue	Compression +Compression
6	1	Р	Compression	Glue	Compression
7	1	A, P	Coil, Glue	Glue	Compression
8	1	A, P	PVA, Bleomycin	Glue	Tourniquet
9	2	A, P+A	Glue+PVA	Glue+None	Compression +Compression
10	1	A, P	PVA, Bleomycin	Glue	Compression
11	1	A, P	Coil	Glue	Compression
Patient ID	Occlusion rate (%)	Outcome	Complications	Follow-up ti	me (months)
1	>90	Significant improvement	None	7	'5
2	50-90	Partial improvement	None	67	
3	100	Complete recovery	None	53	
4	>90	Significant improvement	None	55	
5	<50	No change	None	3	3
6	>90	Significant improvement	None	41	
7	100	Complete recovery	None	25	
8	100	Complete recovery	None	(6
9	100	Complete recovery	None	1	5
10	>90	Significant improvement	Mild	4	
11	100	Complete recovery	None	4	4

performed initially followed by percutaneous glue injection. Patient 4 underwent staged embolization due to the large

lesion, with residual nidus embolization during the second hospitalization.

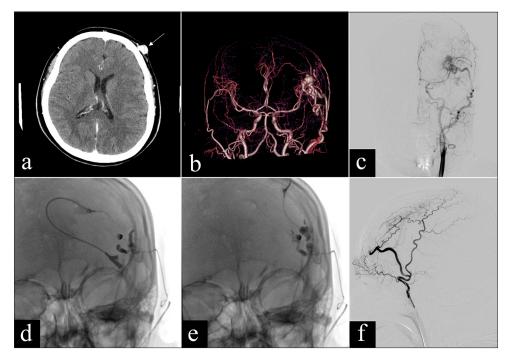


Figure 1: A 27-year-old female (Patient 7) diagnosed with left frontal arteriovenous malformation (AVM), characterized by swelling, locally elevated skin temperature, occasional throbbing pain, and palpable tremor, classified as type IIIb. (a) Axial enhanced computed tomography (CT) revealing subcutaneous AVMs in the left frontal region, with visible abnormal vascular clusters (yellow arrow). (b) CT angiography displaying the vascular architecture of the AVM, highlighting multiple dilated feeding arteries and draining veins. (c) Angiography revealing the vascular structure and dilated feeding arteries and draining veins. (d) Fluoroscopic image after the release of spring coils (3 mm and 4 mm each) via arterial access and subsequent use of tissue glue embolization, depicting the coils in place and n-butyl cyanoacrylate (nBCA)/iodized oil cast. (e) Fluoroscopic image after percutaneous injection of tissue glue, displaying nBCA/iodized oil cast. (f) Post-embolization angiogram demonstrating complete occlusion of the AVM nidus. Following the procedure, the patient experienced complete resolution of symptoms, with no recurrence observed during follow-up. The treatment outcome was categorized as "complete recovery."

One patient had an occlusion rate between 50 and 90%. This patient (Patient 2) underwent three embolization treatments. Initially, after ligation of the draining vein, arterial embolization with glue was performed, followed by percutaneous glue injection. During the second hospitalization, the patient underwent percutaneous polidocanol sclerotherapy. During the third hospitalization, after ligation of the draining vein, arterial embolization with PVA was performed, followed by percutaneous glue injection.

One patient had an occlusion rate below 50%. In this case (Patient 5), direct percutaneous glue injection was performed under manual compression of veins due to tortuous and small feeding arteries.

Treatment outcomes

The comprehensive treatment outcomes of AVM patients are summarized in Table 2. Clinical symptoms completely disappeared in three patients, with two experiencing pain cessation and three exhibiting complete regression of swelling. In addition, clinical symptoms significantly improved in four patients, with three reporting notable alleviation of pain, four showing considerable reduction in swelling, and one displaying marked reduction in varicose vein clusters. Moreover, partial relief of symptoms was observed in four patients, with three demonstrating reduced swelling and two showing decreased vascular clusters, while one experienced partial relief of pain.

Thus, 5 (45.4%) cases achieved complete recovery, 4 (36.4%) cases showed significant improvement, 1 (9.1%) case was categorized as partial improvement, and 1 (9.1%) case showed no change. The overall treatment success rate was 81.8% (9/11). During the follow-up period, none of the patients experienced symptom exacerbation or recurrence.

There were no significant differences in treatment efficacy among patients with different Schobinger stages and Cho-Do subtypes [Table 3].

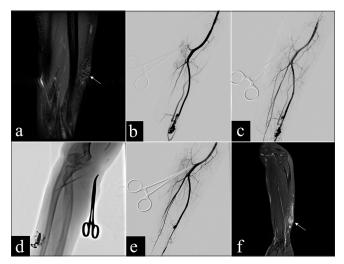


Figure 2: A 28-year-old male (Patient 11) diagnosed with arteriovenous malformations (AVMs) on the ulnar side of the right forearm, presenting symptoms including swelling, pain, locally elevated skin temperature, and palpable pulsation, classified as type IIIb. (a) Coronal T2-weighted magnetic resonance imaging (MRI) sequence revealing a lesion on the ulnar side of the right forearm, with evident flow voids indicating blood vessels (yellow arrow). (b) Angiography displaying the AVM nidus (yellow arrow). (c) Postrelease of spring coils via arterial access (two coils of 3 mm and one coil of 4 mm), angiography showing partial occlusion of the nidus and reduced blood flow in the AVMs. (d) Fluoroscopic image during percutaneous injection of tissue glue. (e) Post-embolization angiogram showing complete occlusion of the AVMs nidus. (f) One-month post-embolization, coronal T2-weighted MRI sequence indicating a reduction in the lesion size compared to previous images, with no visible flow voids (yellow arrow). Following the procedure, the patient experienced significant improvement in swelling, normalization of skin temperature, and basic disappearance of pain. No symptom exacerbation was observed during follow-up. The treatment outcome was categorized as "complete recovery."

Adverse events and complications

Most patients experienced transient local pain and accompanying edema after surgery, which was attributed to inflammatory reactions. Among a total of 16 procedures, one case of mild complication occurred. The patient presented with multiple serpentine and tortuous vascular clusters on the surface of the right forearm and hand before treatment, with involvement of the index and middle fingers. A single embolization treatment was performed, preceded by arterial angiography for diagnostic confirmation. Subsequently, the lesion nests and draining veins were embolized percutaneously under compression of superficial drainage veins using tissue glue, while PVA and bleomycin were administered transarterially to embolize the feeding arteries and residual nests. The patient experienced persistent pain in the surgical area for over 2 months post-embolization, with partial relief achieved through analgesic medication.

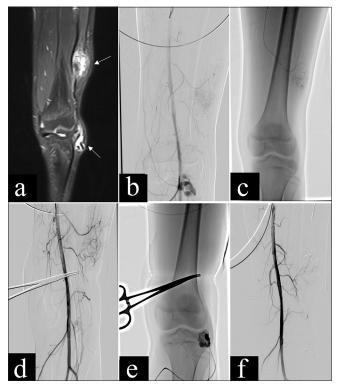


Figure 3: A 12-year-old girl (Patient 8) with arteriovenous malformations (AVMs) in the mid-lower segment of the left thigh and left knee joint, presenting with pulsatile masses. The AVM in the left thigh was classified as type IIIa, while the AVM in the left knee was classified as type IIIb. (a) Coronal T2-weighted magnetic resonance imaging sequence showed two lesion sites in the midlower segment of the left thigh and left knee joint, with visible abnormal vascular clusters (yellow arrows). (b) Angiography displayed the vascular architecture of the AVM lesions at both sites and the dilated feeding arteries and draining veins (yellow arrow). (c) Fluoroscopic image after arterial embolization using PVA and Onyx, showed cast formation within the nidus. (d) Postarterial embolization angiogram demonstrated partial occlusion of the AVM nidus in the mid-lower segment of the left thigh. (e) Fluoroscopic image after percutaneous injection of tissue glue into the AVM nidus in the left knee joint showed n-butyl cyanoacrylate /iodized oil cast. (f) Post-embolization angiogram demonstrated complete occlusion of the AVM nidus. Following the procedure, the pulsatile masses in the left thigh and left knee decreased in size compared to before. No enlargement of the masses or other symptoms were observed during follow-up. The treatment outcome was categorized as "complete recovery."

Visible pigmentation deposits were observed on the skin of the right index and middle fingers. Over the course of more than 3 months post-surgery, the pain gradually subsided and eventually disappeared, with the pigmentation on the skin of the right index and middle fingers remaining largely unchanged.

Table 3: Comparison of therapeutic efficacy in patients withdifferent Schobinger stages and Cho-Do subtypes.							
	Treatment successful	Treatment unsuccessful	P-value				
Schöbinger stages	successful	unsuccessful	P-value				
II	4	1	<i>P</i> =1.00				
III	5	1					
Cho-Do subtypes	successful	unsuccessful	P-value				
II	1	0	<i>P</i> =1.00				
IIIa; IIIb	1	0					
IIIb	7	2					

DISCUSSION

AVMs typically comprise feeding arteries, draining veins, and a nidus, with the latter being the primary focus of therapeutic intervention due to its role in sustaining abnormal circulation. Effective treatment strategies aim to disrupt the abnormal flow within the nidus, reducing hemorrhage risk and alleviating associated symptoms.

nBCA, a widely used liquid embolic agent, polymerizes rapidly upon contact with ionic substances like blood or water,^[28] offering advantages such as affordability and accessibility. Its liquid form enables distal embolization via a flow-directed strategy, facilitating precise vessel occlusion within the AVM nidus.^[29] Upon injection, nBCA initiates a polymerization reaction, leading to acute inflammatory response and eventual vascular occlusion.^[30,31] In our center, nBCA is typically mixed with iodized oil to enhance visibility and control the rate of polymerization. nBCA exhibits a high thrombogenic potential, allowing for progressive and permanent occlusion.^[32] Despite its validated efficacy in AVM embolization, studies have indicated a possibility of recurrence, particularly in cases of partial embolization without complete nidus occlusion.^[18,31]

Although ethylene vinyl alcohol copolymer (EVOH)-based agents such as Onyx and Squid have emerged as effective embolic materials for the treatment of both intracranial and extracranial AVMs over the past decade,^[33,34] their use was not incorporated in this study. This was primarily due to cost limitations and regional accessibility issues. In addition, our team possessed greater familiarity with nBCA, allowing for safer and more controlled embolization in our local context. Nonetheless, we recognize the growing body of evidence supporting EVOH as a valuable tool in AVM management and plan to incorporate it in future investigations as access and clinical experience expand.

The intricate anatomical structures of AVMs, including tortuous feeding arteries, pose challenges in positioning

microcatheter tips near the nidus for complete occlusion. In clinical practice, percutaneous access appears to facilitate better tissue adhesive infiltration into the nidus, achieving more thorough AVM embolization. Although the use of nBCA as a liquid embolic agent is well established for AVM management, our study specifically investigates the synergistic effect of combining percutaneous glue embolization with arterial embolization. This dual technique allows for more comprehensive occlusion of the AVM nidus, particularly in cases where arterial access is limited or insufficient due to small, tortuous feeding vessels. By integrating both access routes, this approach enhances the likelihood of complete nidus embolization, thereby reducing recurrence risk. Consequently, our study initially focuses on arterial occlusion of the nidus and subsequently employs percutaneous embolization for residual nidus occlusion. In Israrahmed et al's study on scalp AVMs, tissue adhesive was exclusively used as an embolic agent for complete occlusion of small lesions or as adjunctive therapy prior to resection of large lesions.^[23] With accumulated experience, the researchers found that direct percutaneous injection of tissue adhesive could penetrate lesions more effectively, achieving more complete embolization. Thus, in the later stages of their study, the investigators primarily adopted the percutaneous approach for embolization therapy, which yielded favorable outcomes. The findings of Israrahmed et al.'s^[23] study align closely with our observations.

Le Fourn et al. utilized nBCA, PVA particles, and absolute ethanol for the treatment of limb and head-and-neck AVMs, achieving a vessel occlusion rate exceeding 75% in 56% of patients (18/32). However, the complication rate was 55% (23/42).^[35] Bissacco et al. employed nBCA for AVM embolization in limbs, with 18 out of 21 (86%) patients achieving optimal or suboptimal results. Specifically, 10 patients exhibited optimal response, indicating complete vessel occlusion, while 8 patients showed suboptimal response, with residual flow within the treated vessels but decreased flow compared to before treatment. It is noteworthy that only 2 (9.4%) complications occurred in this study.^[8] Kitagawa et al.'s study confirmed the safety and efficacy of percutaneous ethanol sclerotherapy combined with arterial embolization using tissue glue for extracranial AVMs. In the study, 87% (20/23) of patients achieved a vessel occlusion rate exceeding 75%, with 2 cases of minor complications.^[17]

Different anatomical locations and angioarchitectural features of AVMs can influence both treatment strategy and risk of complications. For example, digital AVMs may involve end-arterial circulation and pose a higher risk of ischemia and possible amputation. Cirsoid AVMs, particularly those involving the scalp or face, may have complex vascular networks, including potential extracranial-intracranial

arterial anastomoses, increasing the risk of cerebral embolism or pulmonary embolism due to high-flow shunts. In our cohort, no major complications such as amputation or embolic events were observed, likely due to careful case selection, thorough pre-procedural angiographic evaluation, and real-time DSA-guided embolization. Nonetheless, further studies with larger sample sizes are warranted to better understand complication risks across different AVM subtypes.

In this study, 5 (45.4%) cases attained complete recovery, while 4 (36.4%) cases exhibited significant improvement, yielding an overall treatment success rate (vessel occlusion rate exceeding 90%) of 81.8% (9/11). Only one complication was encountered and resolved conservatively. Considering the tendency for AVM recurrence, the efficacy evaluation criteria adopted in this study were stringent. However, compared to previous relevant studies,^[8,17,35] this study demonstrated a higher treatment success rate and a lower complication rate. This outcome may be attributed to the comparatively gentle nature of tissue glue embolization. This suggests that percutaneous injection of tissue glue combined with arterial embolization is safe and effective, with an acceptable complication rate, for superficial AVMs. In this study, there was no statistically significant difference in efficacy among patients with different Schobinger stages and Cho-Do subtypes, possibly due to the small sample size.

This study has several limitations. First, the retrospective nature introduces inherent biases and limits the establishment of causal relationships. The small sample size, only 11 patients over an 8-year period, was primarily due to the rarity of extracranial AVMs and the retrospective design, which limits the ability to accumulate a large number of cases in a short timeframe. Second, the study cohort was heterogeneous, involving various anatomical sites such as the hand, limbs, and scalp (including one case of cirsoid aneurysm). This heterogeneity may introduce variability in clinical presentation and treatment response. However, it also mirrors real-world clinical scenarios and broadens the applicability of our findings. Third, the lack of continuous and standardized follow-up data hampers the comprehensive evaluation of treatment efficacy, and the risk of AVM recurrence over time remains a concern. Furthermore, treatment outcomes were primarily assessed based on imaging, while clinical symptom relief was not thoroughly evaluated. Future prospective, multicenter studies with larger and more homogeneous populations are warranted to validate our conclusions and to compare the long-term efficacy of different embolization techniques.

CONCLUSION

The combined approach of percutaneous glue embolization with arterial embolization proves to be safe and effective

in treating extracranial AVMs. However, further research validation and broader data analysis are needed to strengthen the evidence base and optimize treatment strategies.

Ethical approval: The research/study was approved by the Institutional Review Board of the Third Affiliated Hospital of Sun Yat-sen University, number No. II2023-250-01, dated September 2023.

Declaration of patient consent: The authors certify that they have obtained all appropriate patient consent.

Financial support and sponsorship: Nil.

Conflict of interest: There are no conflict of interest.

Use of artificial intelligence (AI)-assisted technology for manuscript preparation: The authors confirms that there was no use of artificial intelligence (AI)-assisted technology for assisting in the writing or editing of the manuscript and no images were manipulated using AI.

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How to cite this article: Tong YC, Zhao Y, He HP, Lin R, Lv JB, Zhang YB, *et al.* Percutaneous n-butyl cyanoacrylate glue embolization combined with arterial embolization for extracranial arteriovenous malformations. J Clin Imaging Sci. 2025;15:19. doi: 10.25259/JCIS_33_2025