ORIGINAL ARTICLE



Clinical outcomes of adjunct sinus stenting in dural arteriovenous fistulas

Role of flow restoration in steno-occlusion and cortical venous reflux

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Abstract

Purpose This study investigated the clinical outcomes and follow-up results of venous sinus stenting (VSS) for constrictive dural sinus restoration in patients with intracranial dural arteriovenous fistulas (DAVFs). We hypothesized that this treatment would have durable benefits in such patients.

Methods Patients who underwent VSS for DAVFs with cortical venous reflux between January 2008 and June 2020 were identified after a retrospective review (n = 18). Clinical and endovascular treatment data and follow-up information were reviewed.

Results The mean age of the 18 patients was 59.9 years. Stents were implanted in 10 previously occluded sinuses and 9 stenotic sinuses in addition to endovascular embolization. One patient received bilateral VSS. Subdural hemorrhage occurred in one patient after recanalization for embolization, followed by uneventful stenting. In 17 patients with clinical follow-up, the median follow-up time was 59.5 months (interquartile range 18 to 84 months). Of these, sixteen patients had follow-up vascular imaging, revealing AVF obliteration in 6 patients (38%) and stent patency in 11 (69%). Retreatment was performed for 8 (50%) patients. The mean follow-up modified Rankin scale score was 1.28. All patients had longstanding symptomatic improvement.

Conclusion Restoration of sinus flow in DAVFs with cortical venous reflux through VSS has an acceptable complication rate and long-term symptomatic control; however, retreatment is often required, and stent occlusion is not uncommon.

Availability of Data and Material Data can be made available by contacting the corresponding author via email under reasonable request.

Code Availability The SAS code can be made available by contacting the corresponding author via email under reasonable request.

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Keywords Dural arteriovenous fistula · Cerebral venous reflux · Venous sinus stenting · Endovascular · Follow-up

Introduction

Intracranial dural arteriovenous fistulas (DAVFs) are cerebrovascular malformations characterized by an abnormal connection between the dural arteries and venous sinus or cortical veins [1]. Progression of DAVF results in excess cerebral venous flow, reactive dural sinus constriction, and impaired cerebral venous drainage [2]. Therefore, outflow restriction and venous hypertension are major factors for hemorrhagic complications [3]. Endovascular treatment is effective against intracranial DAVFs because it immediately curtails the shunting flow into the cerebral venous system. Nevertheless, a sinus occlusive approach, particularly that administered via the transvenous route, is crucial for obtaining angiographic cure and achieving optimal embolization [4]; however, normal cerebral drainage function can be affected after treatment. Therefore, a sinus-preserving approach is a more attractive choice than the preceding methods because it enables the physiological venous drainage of brain tissue.

Dural sinus angioplasty and stenting have been proposed as a method for restoring venous outflow after sinuspreserving embolization for high-grade DAVFs with sinus stenosis or occlusion [5]. Moreover, in some cases stenting can help reduce the shunting flow [6]; however, dural sinus preservation occasionally reduces the likelihood of achieving complete DAVF obliteration. Therefore, despite the technical feasibility of sinus stenting for DAVF with sinus constriction, its safety, efficacy, and long-term outcomes are largely unknown. We hypothesized that a sinuspreserving approach involving venous sinus stenting (VSS) would be effective in treating DAVFs with sinus restriction.

Accordingly, this paper presents our single-center experience of a sinus-preserving approach involving dural sinus stenting in addition to embolization for the treatment of DAVFs.

Material and Methods

Study Population

The study was approved by our institutional review board, and individual patient informed consent was waived. A total of 322 patients with intracranial DAVFs were treated through endovascular therapy between January 2008 and June 2020. After a retrospective review of patient records, those who received dural sinus stents for downstream sinus stenosis or occlusion were identified and their DAVFs were classified according to the Cognard classification system [7]. Because restoration of normal venous outflow function may be potentially beneficial for brain circulation, adjunct VSS was considered to treat those with downstream sinus constriction. At our institute, stenting is applied to restore a diseased dural sinus if the following criteria are met: 1) a lateral sinus DAVF involving the dominant sinus and having a size >5 mm in normal segment and 2) a superior sagittal sinus or straight sinus DAVF with at least one normal downstream lateral sinus; however, stent implantation would not be considered if the following clinical conditions are observed: 1) massive venous infarct and large intraparenchymal hematoma and 2) failed guiding catheter placement into the dural sinus. On the basis of these criteria, our final analysis included 18 patients.

Endovascular Procedure

The endovascular procedure was performed by a board certified neurointerventionalist (C-W L) and at least one of two other neurointerventionalists (H-M L and Y-H L) in the neuroangiography suite by using a biplane angiography machine with the patient under general anesthesia. Loading doses of dual antiplatelet agents, including 300 mg aspirin and 300 mg clopidogrel, were administered before or at the start of the procedure. Maintenance doses of 100 mg aspirin and 75 mg clopidogrel were continued for 6 months, followed by single agent indefinitely. Heparinization was not applied. Angioplasty was performed first with a monorail angioplasty balloon after successful wire crossing of the involved dural sinus to perform sinus stenting. For an occluded sinus, a 0.035-inch guidewire or 0.027-inch microcatheter-microwire kit (Progreat, Terumo, Tokyo, Japan) was used along with diagnostic or guiding catheters applied in a coaxial or triaxial manner. After angioplasty, stents of variable sizes and lengths were used; such stents were primarily self-expandable nitinol stents (Precise, Cordis, Santa Clara, CA, USA and Zilver Flex, Cook Medical, Bloomington, IN, USA) and occasionally balloon-expandable stent grafts (GraftMaster, Abbott, Chicago, IL, USA). Moreover, a guiding sheath was placed into the sinus if feasible to facilitate stent placement. In addition to stent placement, endovascular embolization was performed either before or after stent placement. The goal was to sufficiently reduce the flow of AVF. Notably, the embolization process was performed through either a transarterial or a transvenous route using various embolic materials, including coils, n-butyl-2-cyanoacrylate (Histoacryl®, B. Braun, Melsungen, Germany), and ethy-

	mRS	0	-	0	ε	-	0	0	-
	Clinical results	Asympto- matic	Still dizziness, other- wise no change	Asympto- matic	Unknown	Occasional seizure	Asympto- matic	Improved neuro- logical deficit	Occasional seizure, headache, no tinnitus
	tt/ Clinical follow- ic up (month)	S	14	6	0	20	57	58	61
	Retreatmer method/ angiograph result	Yes/TAE/ trace residual	No	Yes/TAE/ trace residual	No	No	Yes/TAE/ residual type 2A	No	No
	l Radiological result/ stent patency	Recurrence and CVR/ occlusion	No re- currence/ ISRS	Recurrence and CVR/ ISRS	NA	No re- currence/ patent	Progression to type 2A/ patent	Regressed/ patent	Stable type 2A/ patent
	Radiologica follow- up (month)	2	×	7	NA	20	47	20	55
	Compli- cation	Acute stent throm- bosis	1	I	I	1	Stent acute occlu- sion	I	1
-up results of individual patients	Initial angio- graphic result	Total oblit- era- tion	Total oblit- era- tion	Trace resid- ual	Trace resid- ual	Trace resid- ual	Residual type 1	Residual type 2A	Residual type 2A
	Timing	Simul- taneous	Simul- taneous	Simul- taneous	Simul- taneous	Stenting only	Simul- taneous	Simul- taneous	Simul- taneous
	Stent location/ type	Straight sinus/ Precise	Left SS/ Graft master	Left SS/ Precise	Left SS to jugular Zilver flex	Right SS/ Zilver flex	Right TS/ Precise	Left TS/ Precise	Right TS to SS/ Precise
and follow	Emboli- zation mate- rial	Onyx, coil	Onyx	Onyx	Onyx	NA	Coil	Coil, onyx	Coil, onyx
atment details,	Treatment strategy	VSS fol- lowed by combined TAE and TVE	TAE fol- lowed by VSS	TAE fol- lowed by VSS	VSS fol- lowed by TAE	VSS fol- lowed by TAE	TVE followed by VSS	VSS fol- lowed by combined TAE and TVE	VSS fol- lowed by combined TVE
stics, endovascular tr	DAVF location/ Cognard grade/ diseased sinus	Falcotentorial region/type 2B/ straight sinus severe stenosis	Left SS/type 4/ left SS and TS occlusion	Left TS to torcula/ type 2A+B/ left SS occlusion	Left SS/type 4/ left SS occlusion	Right SS and SSS/ type 2A+B/ right SS and TS occlusion	Right TS and SSS/ type 2A + B/right TS occlusion	Bilateral TS and torcula/ type 2A+B/ bright TS oc- clusion, left TS severe stenosis	Left TS/ type 2A+B/ bilateral TS stenosis
cal characterist	Clinical presenta- tion	Chemosis, headache	ICH, sensory aphasia	Progressive memory loss	Seizure, conscious distur- bance	Hemianopia	Headache and seizure	Sudden weakness, aphasia, headache	Tinnitus
e 1 Clinic	Age (years)/ sex	47/ Male	55/ Male	66/ Female	53/ Female	55/ Male	68/ Male	68/ Female	33/ Female
Tab	No	1	7	ς	4	Ś	9	2	×

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1 (Continued)	inued) Clinical DAVE location/	DAVF location/		Treatment	Emboli_	Stent	Timina	Initial	Comoli-	Radiological	Radiolonical	Retreatment/	Clinical	Clinical	s dur
Age CLINICAL DAYFLOCATION ITEAUTIENT ENTROIT NEMT years)/ presenta- Cognard grade/ strategy zation location/ ex tion diseased sinus mate- type rial	CHILLER DAYF LOCAUCIUM ALEAUTION - SIGHT presenta- Cognard grade/ strategy zation location/ tion diseased sinus mate- type rial	Cognard grade/ strategy zation location/ Cognard grade/ strategy zation location/ diseased sinus mate- type rial	rreauticut Eurobui- Stent strategy zation location/ mate- type rial	zation location/ mate- type rial	location/ type		guilling	angio- graphic result	cation	Raunouogua follow- up (month)	result/ stent patency	Retreatment method/ angiographic result	follow- up (month)	results	
 Blurred Right SS-TS/ VSS fol- Coil, Right SS Male vision, type 2A+B/right lowed by onyx, to IJV/ throbbing SS occlusion combined parti- Zilver headache TAE and cle flex TVE 	Blurred Right SS-TS/ VSS fol- Coil, Right SS vision, type 2A+B/right lowed by onyx, to IJV/ throbbing SS occlusion combined parti- Zilver headache TAE and cle flex TVE	Right SS-TS/ VSS fol- Coil, Right SS type 2A+B/right lowed by onyx, to IJV/ SS occlusion combined parti- Zilver TAE and cle flex TVE	VSS fol- Coil, Right SS lowed by onyx, to IJV/ combined parti- Zilver TAE and cle flex TVE	Coil, Right SS onyx, to IJV/ parti- Zilver cle flex	Right SS to IJV/ Zilver flex		Simul- taneous	Residual type 2A	1	89	Stable type 2A/ patent	No	74	Mild pulsatile sensation, other improved	5
60/LeftTorcula to rightVSS fol-Coil,RightFemaleweakness,TS/type 2A+B/lowed byonyxTS todrowsyright TS-SScombinedSS andjunction stenosisTAE andleft TS/and left proximalTVEPrecise*2TS occlusionTS occlusion	LeftTorcula to rightVSS fol-Coil,Rightweakness,TS/type 2A+B/lowed byonyxTS todrowsyright TS-SScombinedSS andjunction stenosisTAE andleft TS/and left proximalTVEPrecise*2TS occlusion	Torcula to rightVSS fol-Coil,RightTS(type 2A+ B/lowed byonyxTS toright TS-SScombinedSS andjunction stenosisTAE andleft TS/and left proximalTVEPrecise*2TS occlusion	VSS fol- Coil, Right lowed by onyx TS to combined SS and TAE and left TS/ TVE Precise*2	Coil, Right onyx TS to SS and left TS/ Precise*2	Right TS to SS and left TS/ Precise*2		Simul- taneous	Residual type 2A	I	78	Progression to 2B/ occlusion	Yes/TVE/ total oblitera- tion	80	No seizure, occa- sional dizziness	-
 2.2' Right Bilateral SS- Combined Coil, Right Male tinnitus, TS/type 2A+B/ TAE and onyx TS/ headache right TS focal TVE Precise*2 stenosis, left TS followed long segment by VSS occlusion 	RightBilateral SS-CombinedCoil,Righttinnitus,TS/type 2A+B/TAE andonyxTS/headacheright TS focalTVEPrecise*2stenosis, left TSfollowedlong segmentby VSSocclusionocclusionocclusion	Bilateral SS-CombinedCoil,RightTS/type 2A+B/TAE andonyxTS/TS/type 2A+B/TVEonyxTS/right TS focalTVEPrecise*2stenosis, left TSfollowedlong segmentby VSSocclusionocclusion	Combined Coil, Right TAE and onyx TS/ TVE Precise*2 by VSS	Coil, Right onyx TS/ Precise*2	Right TS/ Precise*2		Delayed after em- boliza- tion	Residual type 1	I	79	Progression to type 2A/ patent	Yes/TAE and TVE/ trace residual	86	Asympto- matic	0
 Functional Left SS-TS Combined Coil, SSS to leaded decline, to torcula/ TAE and onyx right TS/ a poor type 2A+B/ TVE Zilver Cliver nemory left TS stenosis followed flex leaded sion, right TS stenosis 	FunctionalLeft SS-TSCombinedCoil,SSS todecline,to torcula/TAE andonyxright TS/apoortype 2A+B/TVEZilveromemoryleft TS stenosisfollowedflex1and SS occlu-by VSSstenosisstenosistenosisstenosisstenosisstenosisfollowedflex1	Left SS-TSCombinedCoil,SSS toto toreula/TAE andonyxright TS/istype 2A+B/TVEZilveroleft TS stenosisfollowedflex1and SS occlu-by VSSsion, right TSstenosisstenosisfollowedflex1	Combined Coil, SSS to TAE and onyx right TS/ is TVE Zilver of followed flex 1 by VSS 1	Coil, SSS to onyx right TS/ Zilver flex 1	SSS to right TS/ Zilver flex		Delayed after em- boliza- tion	Residual type 1	1	30	Progression to type 2A/ patent	Yes/TVE/ total oblitera- tion	86	Asympto- matic	0
19/ Seizure, SS/ TAE fol- Onyx SSS/ I Male venous type 2A+B/ lowed by Precise a infarction posterior SSS VSS vSS focal stenosis t t	Seizure, SSS/ TAE fol- Onyx SSS/ I venous type 2A+B/ lowed by Precise a infarction posterior SSS VSS e focal stenosis tt	SSS/ TAE fol- Onyx SSS/ I type 2A+B/ lowed by Precise a posterior SSS VSS vSS t focal stenosis t	TAE fol- Onyx SSS/ I lowed by Precise a VSS 6 b	Onyx SSS/ I Precise a b b t t	SSS/ I Precise a b b t	цеорт	Delayed fiter m- ooliza- ion	Trace resid- ual	I	12	Progression to type 2B/ ISRS	Yes/ direct puncture/ trace residual	84	Impaired cognitive function, mild dementia	
66/ Aphasia Left SS-TS/ VSS fol- Onyx Left TS S Male type 2A+B/ lowed by to SS/ b left TS and SS TAE Precise*2 e severe stenosis thrombosis	Aphasia Left SS-TS/ VSS fol- Onyx Left TS S type 2A+B/ lowed by to SS/ b left TS and SS TAE Precise*2 e severe stenosis thrombosis thrombosis	Left SS-TS/ VSS fol- Onyx Left TS S type 2A+B/ lowed by to SS/ b left TS and SS TAE Precise*2 e severe stenosis with partial t thrombosis	VSS fol- Onyx Left TS S lowed by to SS/ b TAE Precise*2 e t	Onyx Left TS S to SS/ b Precise*2 e t	Left TS S to SS/ b Precise*2 e b t	N D O D D	taged efore m- oliza- ion	Trace resid- ual	1	NA	NA	No	18	Stable	0
60/ Tinnitus, Left SS-TS/ VSS fol- Coil, Left S emale conscious type 4+5/left TS lowed by onyx, TS to tt distur- an SS occlusion TAE parti- jugular bance cle bulb/ Precise*2	Tinnitus, Left SS-TS/ VSS fol- Coil, Left S conscious type 4+5/left TS lowed by onyx, TS to t distur- an SS occlusion TAE parti- jugular bance cle bulb/ Precise*2	Left SS-TS/ VSS fol- Coil, Left S type 4+5/left TS lowed by onyx, TS to tt an SS occlusion TAE parti- jugular cle bulb/ Precise*2	VSS fol- Coil, Left S lowed by onyx, TS to t TAE parti- jugular cle bulb/ Precise*2	Coil, Left S onyx, TS to ti parti- jugular cle bulb/ Precise*2	Left S TS to ti jugular bulb/ Precise*2	2 H	imul- aneous	Trace resid- ual	I	6	No re- currence/ occlusion	No	19	Asympto- matic	0

mRS	2v	ŝ	ω
Clinical results	Bed rid- den and aphasia due to left MCA infarct 5 years later	Occasional dizziness, still unsteady gait; otherwise improved	Residual deficit
/ Clinical follow- c up (month)	103	107	61
Retreatment method/ angiographic result	No	Yes/TAE and TVE/ total oblitera- tion	No
1 Radiological result/ stent patency	no recur- rence/ patent	Progression to type 2A + B/ ISRS	Stable type 1/ patent
Radiologica follow- up (month)	36	Ξ	37
Compli- cation	1	1	SDH and craniec- tomy
Initial angio- graphic result	Total oblit- era- tion	Residual type 1	Residual type 1
Timing	Simul- taneous	Staged before em- boliza- tion	Delayed after em- boliza- tion
Stent location/ type	Left TS/ Precise	Left SS/ Balloon mounted stent	Right SS/ Zilver flex
Emboli- zation mate- rial	Coil	Coil, onyx	Coil, NBCA
Treatment strategy	VSS fol- lowed by TVE	VSS fol- lowed by combined TAE and TVE	Combined TAE and TVE followed by VSS
DAVF location/ Cognard grade/ diseased sinus	Left TS/type 4/ left TS and SS occlusion	Left SS-TS/ type 2A+B/left SS stenosis	Right SS/ type 2A+B/ right SS and left TS occlusion
Clinical presenta- tion	Progressive dementia, venous infarction and hem- orrhage	Cognitive decline, dizziness, tinnitus, unsteady gait	Headache
Age (years)/ sex	81/ Female	47/ Female	41/ Female
No	16	17	18

GraftMaster (Abbott, Chicago, IL, USA); Onyx (Medtronic, Dublin, Ireland); Precise (Cordis, Santa Clara, CA, USA); Zilver Flex (Cook, Bloomington, IN, USA) *ISRS* in-stent restenosis, *IJV* internal jugular vein, *MCA* middle cerebral artery, *NA* not applicable, *NBCA* N-butyl cyanoacrylate, *SDH* subdural hemorrhage, *SS* sigmoid sinus, *SSS* superior sagittal sinus, *TAE* transverse sinus, *TVE* transverse sinus, *TSE* venous sinus stenting

lene-vinyl alcohol (OnyxTM, Medtronic, Dublin, Ireland). Balloon-assisted techniques were not used during Onyx injection. The procedures were occasionally performed in stages, with the stenting and embolization performed at different sessions.

Clinical Assessment

Clinical data were retrieved from the electronic medical records of each patient for analysis. Presenting symptoms were classified as aggressive or nonaggressive symptoms [8]. Aggressive symptoms included focal neurological deficits, seizures, rapidly progressive dementia and hemorrhagic presentation; nonaggressive symptoms included ocular symptoms, pulsatile tinnitus, headache, dizziness, and vertigo. Radiological investigations, including catheter angiography and magnetic resonance imaging (MRI), were conducted routinely before the procedure. After the procedure, patients were instructed to undergo follow-up examination in clinics. Noninvasive radiological follow-up was ordered according to physicians' requests. In principle, MRI was arranged 6 months after the procedure and annually thereafter. If there was evidence of disease progression on MRI, catheter angiography would be considered. All imaging studies were re-evaluated by two experienced neuroradiologists (C.W.L and Y.H.L with 19 and 11 years of experience, respectively). The conditions of presenting symptoms were investigated. In addition, any retreatment during the follow-up period was recorded. The clinical functional status was assessed using the modified Rankin scale (mRS).

Statistical Analysis

Herein, descriptive statistics are presented for demographic, angiographic, and clinical treatment data, and individual treatment details are provided. In addition, the follow-up results are presented. Because of the limited sample size, other inference statistics were not generated. All statistical analyses were performed using SAS software version 9.4 (SAS Institute, Cary, NC, USA).

Results

Study Patients

The mean age of the 18 study patients was 59.9 years (standard deviation 14.2 years) and 8 patients (44%) were male.



Fig. 1 Illustrative case of stent placement in an isolated occluded right lateral sinus (patient 5). This 55-year-old man presented with hemianopia. **a,b** Frontal and lateral view arterial phase of right external carotid artery reveals a right sigmoid sinus Cognard type 4 dural arteriovenous fistula with an isolated right lateral sinus with cortical reflux into the vein of Labbé (*asterisk*). *White arrows* indicate locations of stenosis and occlusion at right transverse and sigmoid sinus. **c** A self-expandable stent (Zilver flex, Cook) was placed to right sigmoid sinus after angioplasty. The locations of stent edges are indicated by the *black arrows*. Transarterial embolization by Onyx was then performed after stent placement. **d,e** Lateral view arterial phase image after stent placement and embolization reveals nearly complete obliteration of fistula with residual trace amount of shunting flow at jugular bulb (*dashed circle*). Venous phase image shows antegrade flow of vein of Labbé draining into sigmoid sinus (*asterisk*). **e** Contrastenhanced MR venogram 20 months after endovascular treatment reveals enhancement within stent, suggestive of patency. No evidence of fistula progression was found on MR angiography (not shown). The neurological symptom improved clinically



Fig. 2 Patient 11 with bilateral sinus placements. This 80-year-old woman had left side weakness and consciousness disturbance at time of diagnosis. **a,b** Frontal view arterial phase of bilateral common carotid artery injection reveal dural arteriovenous fistula at bilateral lateral sinus and torcular area. There are right transverse sinus focal stenosis and left transverse sinus occlusion, as well as extensive venous reflux into the superior sagittal sinus and cortical veins (*asterisk*), suggestive of Cognard type 2A+B disease. **c,d** After the guiding catheter was navigated across the right jugular bulb, angioplasty was performed on the occluded segment of bilateral lateral sinuses. The stents were deployed thereafter sequentially in the same session (*arrows*). Combined transarterial and transvenous embolization was performed after stent placement. **e, f** Frontal view of bilateral common carotid artery injection after embolization reveals substantially decreased shunting flow but residual type 2A disease at torcular region with retrograde flow into superior sagittal sinus. **g** After 45 months later, follow-up MRI shows engorged cortical and medullary vein in the bilateral cerebral hemispheres, suggestive of disease progression and venous congestion. **h** Left common carotid angiography reveals recurrent fistula with superior sagittal sinus and cortical venous reflux (*asterisks*) and occlusion of previous deployed stents (*arrow*). **i** Salvage transvenous dural sinus occlusion by coils was performed. **j,k** Bilateral common carotid artery shows obliteration of fistula after embolization. **l** Follow-up MRI 79 months after initial diagnosis shows normalization of engorged cortical and medullary veins. Besides occasional dizziness, the patient was in independent functional status

Angiographic studies revealed that all patients had cortical venous reflux. Cortical venous reflux from an isolated sinus was noted in four patients, and the corresponding DAVFs were classified as Cognard type IV fistulas. The remaining patients had a visible flow in the sinus, and the corresponding DAVFs were classified as Cognard type IIB or IIA+B fistulas. In 16 patients, the fistulas were located in the lateral sinus, comprising the sigmoid, transverse sinus, and torcular region. One fistula was located in the posterior superior sagittal sinus, and one was in the falcotentorial region. Aggressive symptoms were observed in 13 patients (72%). The remaining five patients (28%) had headache, dizziness, tinnitus, and chemosis. Intracranial hemorrhage occurred in one patient (patient 2) at the time of diagnosis. Table 1 shows individual patient data.

Endovascular Treatment Strategy and Initial Angiographic Result

Because VSS was used as adjunct therapy, endovascular embolization, either via transarterial, transvenous, or combined routes, were planned in single or staged fashion. In 11 patients, embolization and stent placement were performed simultaneously. A patient (patient 5) only had minimal residual AVF after stent implantation and thus did not require further embolization. In patients with staged endovascular treatment, VSS was performed before embolization in four patients; in these patients, the interval between embolization and stenting ranged from 3 to 183 days and 4 patients received planned staged embolization after stenting within 1 month.

Stents were implanted in 10 previous angiographically occluded sinuses and 9 stenotic sinuses. Fig. 1 shows an illustrative case with occluded sinus (patient 5). Patient 10 received bilateral lateral sinus stents (Fig. 2). Angiographic complete obliteration was observed immediately in three patients (17%), subtotal obliteration (trace amount of residual Cognard type 1 flow without significant early venous drainage) in six patients (33%), and significant residual flow (substantial Cognard type 1 and type 2A flow) in nine patients (50%). During the procedure, acute stent occlusion occurred in two patients (patients 1 and 6), and thrombectomy was performed successfully in them. Patient 18 had subdural hemorrhage immediately after embolization, and underwent decompressive craniectomy. This patient received stent placement 6 months after embolization uneventfully. No new hemorrhage, venous infarcts, or focal neurological deficits occurred after stent implantation in any of the 18 patients.

Follow-Up Results and Retreatment

One patient was lost to follow-up after 2 months, and the remaining patients had at least 5 months of follow-up (median 59.5 months; interquartile range, IQR 18-84 months). Among the 17 patients who received adequate clinical follow-up, 16 underwent vascular imaging studies (median 25 months; IQR 10-51 months). Among patients who underwent follow-up vascular imaging 5 (31%) had AVF obliteration and 11 (69%) had stent patency. On the other hand, eight (50%) had progressive recurrent AVF as observed through radiological assessment, including one (patient 1) with previous complete obliteration. Cortical venous reflux occurred in five patients (63%) with progressive recurrent AVF. The remaining three patients (19%) had stable disease status. In four patients with Cognard type IV DAVFs (isolated sinus), AVF obliteration was persistent in all patients during follow-up; patent stents were noted in three patients (75%).

In clinical assessment six patients were asymptomatic during follow-up (35%), particularly patients with Cognard type IIA + B or IIB DAVFs (5 patients, 83%). In other patients, the presenting symptoms improved, but they still had minor symptoms or pre-existing neurological deficits. As for functional status, the mean mRS score at follow-up was 1.28 (SD: 1.45). Patient 4 had a mRS score of 3 and was lost to follow-up because of a psychiatric disorder. Patient 16 had left M1 middle cerebral artery large vessel occlusion stroke 3 years after DAVF treatment, with a final mRS score of 5 after the event; however, other patients had favorable functional outcomes (mRS score ≤ 2).

Retreatment was conducted in all of 8 patients with progressive AVF. In patients 11 and 17 the diseased sinuses with permanent stent occlusion were expunged to achieve complete AVF obliteration. In patient 10 cerebral congestion was noted 4 years after stenting. The angiographic study revealed stent occlusion, and angioplasty was performed on the stent in addition to AVF embolization. In patient 13 Onyx embolization via puncturing to skull intraosseus vascular channel was performed due to limited access route. The remaining patients all received standard endovascular embolization. Angiographic complete obliteration was observed in three patients (38%), subtotal obliteration (trace amount of residual Cognard type 1 flow without significant early venous drainage) in four patients (50%), and significant residual flow (type 2A flow) in one patient (12%). No periprocedural complication was found. The clinical symptoms and functional status were stable during further follow-up.

Discussion

Patients with high-grade intracranial DAVFs often experience an unfavorable natural history if untreated. Our results revealed that the dural sinus could be restored through stent implantation with acceptable complication rates. During the long-term follow-up in this study, stent patency was noted to be maintained in 69% of the patients; however, only 38% of the patients could achieve angiographic AVF obliteration, and up to 50% of the patients needed retreatment. Notably, all patients had symptomatic improvement compared with the preoperative condition. Therefore, we believe that adjunct dural sinus stenting is a reasonable treatment approach when carefully planned, especially in previously occluded sinuses; however, this approach would require long-term follow-up.

In DAVFs with cortical venous reflux, the annual hemorrhage rate is 6% in patients with Borden type II and 10% in those with Borden type III fistulas [9, 10]. Therefore, in such patients, treatment is mandatory with an endovascular or surgical approach being the preferred method for immediate risk reduction. The worst complication of endovascular treatment is hemorrhage, which typically occurs because of venous hemorrhage from compromised cerebral venous drainage. Japanese registry data revealed the occurrence of complications in 7.7% of cases [11]. Therefore, sinus restoration could be a more physiological method in treating this disease than sinus sacrifice. In our study, only one patient had intracranial hemorrhage after embolization without stenting. Because the hematoma was located in the subdural space after initial embolization, it was considered a procedure-related complication of transvenous embolization. Overall, the complication rate is comparable to that in a large cohort study; however, prudency related to technical issues is warranted in this approach, especially with highgrade stenosis or occlusion.

Nevertheless, the relatively low angiographic cure rate based on our results is a concern for this approach. Ertl et al. revealed that a sinus-preserving approach had a lower cure rate than a sinus occlusive approach (71% vs. 93%) [12]. The likelihood of angiographic cure rate by sinus-preserving embolization can be enhanced by dedicated techniques, such as transarterial dual-lumen scepter balloon or transvenous balloon-assisted Onyx injection [13, 14]. A study by Vollherbst et al. reported that combined venous sinus balloon-assisted protection and transarterial Onyx injection can achieve cure rate of 86.4% [15]. Our results reveal a relatively low angiographic cure rate. The possible reason for this inconsistency could be because we did not use supplementary technique during liquid embolic agent injection. Notably, a recurrence after initial complete angiographic occlusion during follow-up is not uncommon. Ambekar et al. reported that asymptomatic recurrence could occur in 12.3% of patients during follow-up after angiographic cure with Onyx [16]. Hence, clinical follow-up is always required. In our patients, the decision for retreatment was based on imaging findings, and we did not wait until symptom progression. There is possibility that some recurrence might be occult on MRI; but they are usually clinically silent; however, no patient needed emergency retreatment. Nonetheless, although the presenting symptoms seldom disappeared, patients generally experienced symptomatic improvement and had excellent functional status unless burdened by other unrelated diseases. Our results suggest that this treatment approach could maintain long-term efficacy despite the low initial cure rate.

Notably, dural sinus stenting has several applications in the treatment of cerebrovascular disease [17, 18]. In DAVFs, a sinus stent is used for diseases occurring at the lateral sinus [5, 19]; however, several reports have indicated the application of this approach at various locations, including the torcular, superior sagittal, and straight sinuses [20-22]. A series by Levrier et al. reported poststent insertion outcomes in 10 patients, and all patients achieved cure or symptomatic improvement [23]. In their series, all diseases were located in the lateral sinus, and only 30% of patients had cortical venous reflux; the mean follow-up period was 21.1 months. They concluded that stent placement is a promising technique for DAVF treatment. Furthermore, Liebig et al. reported four low-grade lateral sinus DAVFs treated using only angioplasty and stent deployment, achieving satisfactory results [6]. Our results are concordant with their conclusions and we think that stent treatment could be efficient even in patients with cortical venous reflux during a longer follow-up period. Notably, stent placement in chronic stenotic or occluded sinuses is technically demanding in some cases. Nonetheless, crossing a chronically occluded sinus is frequently achievable, which is the foundation of transvenous embolization [24]. Advancements in neurovascular guiding catheters and peripheral or carotid self-expandable stents have facilitated the tracking of a stent to cross the tortuous dural sinus.

Notably, only one patient in our series had an initial presentation of intracranial hemorrhage. In patients who already had a severe hemorrhage, neurosurgical decompression was often required immediately, thereby creating a concern for the use of dual antiplatelet agents after stenting; however, whether patients with hemorrhage are suitable for sinus stents is still inconclusive. Nevertheless, a stepwise approach, wherein AVF is embolized first and the stenting is delayed, might prove to be a feasible strategy for long-term sinus restoration. We do not consider hemorrhage to be an absolute contraindication for dural sinus stent placement, and in certain circumstances acute dural sinus stenting might be helpful [25]; however, there is requirement for further investigation on criteria for sinus stent placement on

the basis of morphological and technical parameters. Evidence of impaired venous collateral routes might help in the selection of the appropriate treatment strategy [26]. Therefore, quantitative measurements, such as digital subtraction angiography or sinus pressure gradient measurement, delineating the severity of sinus restriction could be a useful tool in patient selection and follow-up [27, 28]. Nevertheless, further investigation is required to refine the criteria for patient selection.

Our study had some limitations. First, the retrospective nature of this study could have exerted a risk of selection bias; however, because of the relatively low prevalence of DAVFs and complex anatomical variations, a prospective design was not feasible. Second, our study cohort sample size was small. Nevertheless, the number of patients in our series is larger than those previously reported. Third, relatively heterogeneous treatment strategies were applied to our patients. Because the period was relatively long, the experience and device advancement during embolization treatment might have changed over time. We believe that embolization outcomes have improved in more recent cases. Fourth, the criteria for sinus stent placement were somehow subjective. Nevertheless, all procedures were performed by the same neurointerventionalist, thereby minimizing the variability between patients.

In conclusion, the use of stenting in addition to embolization for treating DAVFs with outflow restriction is safe and durable in most cases; however, the angiographic cure rate is low, and AVF progression is common. Moreover, longterm surveillance is required, and any retreatment can be performed accordingly.

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Author Contribution All authors made (1) substantial contributions to the conception or design of the work or the acquisition, analysis or interpretation of data for the work; (2) drafting of the work or revising it critically for important intellectual content; (3) final approval of the version to be published; and (4) are in agreement to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved.

Declarations

Conflict of interest Y.-H. Lin, C.-W. Lee and H.-M. Liu declare that they have no competing interests.

Ethical standards The work was approved by research ethics committee of National Taiwan University Hospital (Approval number: 202010071RIND). Individual informed consent was waived.

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