

World Journal of *Clinical Cases*

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- 1888 Endoscopic transluminal drainage and necrosectomy for infected necrotizing pancreatitis: Progress and challenges
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- 1903 Functional role of frontal electroencephalogram alpha asymmetry in the resting state in patients with depression: A review
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- 1918 COVID-19 related liver injuries in pregnancy
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- 1939 Laparoscopic common bile duct exploration to treat choledocholithiasis in situs inversus patients: A technical review
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- 1951 Airway ultrasound for patients anticipated to have a difficult airway: Perspective for personalized medicine
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- 1963 Clinicopathological features and expression of regulatory mechanism of the Wnt signaling pathway in colorectal sessile serrated adenomas/polyps with different syndrome types
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- 1974 Effects of individual shock wave therapy *vs* celecoxib on hip pain caused by femoral head necrosis
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- 1985 Very low calorie ketogenic diet and common rheumatic disorders: A case report
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- 2009** Transient ischemic attack induced by pulmonary arteriovenous fistula in a child: A case report
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ABOUT COVER

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Transseptal approach for catheter ablation of left-sided accessory pathways in children with Marfan syndrome: A case report

Zi-Yan Dong, Wei Shao, Yue Yuan, Li Lin, Xia Yu, Lang Cui, Zhen Zhen, Lu Gao

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Abstract

BACKGROUND

Left-sided accessory pathways (APs) can be accessed with either a transaortic (TA) or transseptal approach (TS). For children with Marfan syndrome (MFS) who have aortic disease, the use of TA can aggravate the disease, making TS the best choice for these patients.

CASE SUMMARY

A 10-year-old girl was hospitalized because of intermittent heart palpitations and chest tightness. She was diagnosed with MFS, supraventricular tachycardia, Wolff-Parkinson-White syndrome, and left-sided AP was detected by cardiac electrophysiological. Catheter ablation was successfully performed *via* TS under the guidance of the Ensite system. During the follow-up, no recurrence or complications occurred.

CONCLUSION

The TS for catheter ablation of left-sided APs can be considered in children with MFS. Adequate evaluation and selection of the appropriate puncture site are particularly important.

Key Words: Transseptal approach; Left-sided accessory pathway; Catheter ablation; Pediatric; Marfan syndrome; Case report

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Core Tip: Left-sided accessory pathways (APs) can be accessed with either a transseptal (TS) or transaortic approach (TA). For children with Marfan syndrome (MFS) who have aortic disease, however, the use of TA can aggravate the disease. We report a case of successful catheter ablation of the left-sided accessory pathway by TS in one such child. The TS approach for catheter ablation of left-sided APs appears to be a safe and effective therapeutic option applicable in older children with MFS, though it requires a certain level of training and experience to perform.

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INTRODUCTION

Catheter ablation is the main method for eradicating supraventricular tachycardia (SVT) in children, and left-sided accessory pathways (APs) can be accessed by using a transaortic approach (TA) or a transseptal approach (TS). Reports show that children with Marfan syndrome (MFS) have a higher risk of developing SVT than healthy children[1] and that they also often have aortic lesions. When left-sided APs are accessed in these children, TA for catheter ablation may result in aggravated aortic disease. Hence, TS may be a better method. However, due to the combination of mitral valve disease, the difficulty of catheter ablation in MFS, and the small amount of experience most providers have with atrial septal puncture in children, there are currently no relevant reports on this technique. We therefore report a case of successful ablation of the left-sided AP by TS in a child with MFS and review the current literature, providing new ideas for the treatment of MFS.

CASE PRESENTATION

Chief complaints

A 10-year-old girl was admitted to Beijing Children's Hospital in May, 2021 for a 4-year history of intermittent heart palpitations and chest tightness.

History of present illness

Symptoms started 4 years before the child was admitted to the hospital. She presented with intermittent episodes of heart palpitations and chest tightness with pallor. Each episode lasted from 15 min to 3 h, with a frequency of about twice a year. Paroxysmal SVT was diagnosed at a local hospital, and could be converted to sinus rhythm after application of propafenone. Four months before admission to Beijing Children's Hospital, her palpitations increased in frequency and had occurred 4 times in the last month. Antiarrhythmic drugs were needed to convert.

History of past illness

Pectus excavatum was found at birth. In April, 2014, Nuss pectus excavatum correction was performed, and congenital short esophagus and congenital diaphragmatic hernia were diagnosed. In April 2016, the corrective plate was removed.

Personal and family history

The child was born full-term to nonconsanguineous, healthy parents by normal vaginal delivery after an uneventful pregnancy.

Physical examination

The weight of the patient was 27.5 kg, and her height was 153 cm. A surgical scar of about 6 cm was observed under her right armpit, and she was noted to have dysmorphic features including long and thin extremities with hyperflexible joints, prior pectus excavatum, and scoliosis. Both wrist sign and thumb sign were positive. Marfan syndrome was suspected and additional tests were carried out.

Laboratory examinations

No abnormalities were found in routine blood or urine analyses.

Imaging examinations

Chest X-ray revealed a right diaphragmatic hernia and scoliosis. Echocardiography showed mild mitral and tricuspid prolapse and regurgitation, and the aortic sinus and ascending aorta were widened. Cardiac computed tomography (CT) scan and reconstruction demonstrated severe aortic root dilatation, with left-right diameters of 3.56 cm (z-score of +6.41), anteroposterior diameters of 3.32 cm (z-score of +5.33), and perforating septum thickness of approximately 13.9 mm (Figure 1). Ophthalmic examination revealed high myopia in both eyes and mild opacification of the posterior lens capsule in the right eye. Finally, genetic testing revealed a heterozygous mutation in the FBN1 gene, c.5841delC (p.C1947). Because she had an aortic root Z-score ≥ 2 , a pathogenic FBN1 gene mutation, and a systemic score of 7, MFS was diagnosed according to the 2010 revised Ghent Criteria[2] (Table 1).

Further diagnostic work-up

During hospitalization, the child still suffered from palpitation attacks. Electrocardiogram (ECG) during an episode revealed SVT with a heart rate of 214/min. After a few minutes, the episode terminated spontaneously. The ECG showed sinus rhythm with a ventricular pre-excitation pattern. Paroxysmal SVT and Wolff-Parkinson-White syndrome type A were subsequently diagnosed.

After general anesthesia, cardiac electrophysiological examination was performed under the guidance of Ensite NavX mapping system (Figure 2). Electrode catheters were placed in the coronary sinus (CS) and the right ventricle. Atrioventricular nodal jump phenomenon was not found during atrial program stimulation. Stable narrow QRS tachycardia could be induced by stimulation at 270 ms, with a ventricular rate of 200 bpm and an A/V ratio of 1. The A wave of the CS catheter was eccentrically distributed, and was the most advanced at 1-2 electrodes. Atrioventricular reentrant tachycardia (AVRT) with AP located on the left side was diagnosed.

FINAL DIAGNOSIS

Based on the child's medical history, her final diagnosis was MFS, SVT, Wolff-Parkinson-White syndrome, and left-sided AP.

TREATMENT

Considering that the child had aortic sinus disease, there was a greater risk of retrograde delivery of ablation catheter through the aorta, so we delivered the ablation catheter to the left side of the heart by transseptal puncture performed under X-ray fluoroscopy. The introducer sheath and dilator, followed with the guide wire, were introduced into right femoral vein and placed to the superior vena cava. After that, we pushed the transseptal needle (Brockenbrough BRK, 71 cm, St. Jude Medical/Abbott) to about 1 cm from the end of the dilator. The needle and sheath were rotated clockwise until the tip pointed toward 4 o'clock, and then were slowly retracted. After two "jumps" the tip of the sheath fell into the fossa ovalis. We then adjusted the direction of the introducer sheath tip in the right anterior oblique view, advanced the puncture needle, and injected contrast agent to confirm that the needle tip was in the left atrium. In the left anterior oblique view, the outer sheath had been guided to the left atrium.

The ablation catheter (Safire, 7-F, 115 cm, St. Jude Medical/Abbott) was then advanced into the left atrium along with the sheath, and the target was subsequently mapped. A satisfactory target was marked on the atrial side at a location of 3:30 (o'clock) of the mitral annulus. During sinus rhythm, the ablation was performed at 52°C and 30 W. After 2 s of discharge, the body surface pre-excitation pattern disappeared, and the V-A wave separated. Under ventricular pacing, the V-A wave of the ablation catheter separated, and the reverse A wave of the CS catheter changed to a centripetal distribution. Consolidation discharges were performed at the target for a total of 120 s, and repeated cardiac electrophysiological examination and intravenous isoproterenol failed to induce AVRT or other arrhythmias during the 48-h observation period after ablation. Ventricular pre-excitation was not restored, and echocardiography showed no aggravation of valve injury.

OUTCOME AND FOLLOW-UP

Regular follow-up was performed at 1, 3, 6, and 12 mo, during which ECG, echocardiography, and dynamic ECG were re-examined. No complication or recurrence of SVT were detected at any of these follow-ups.

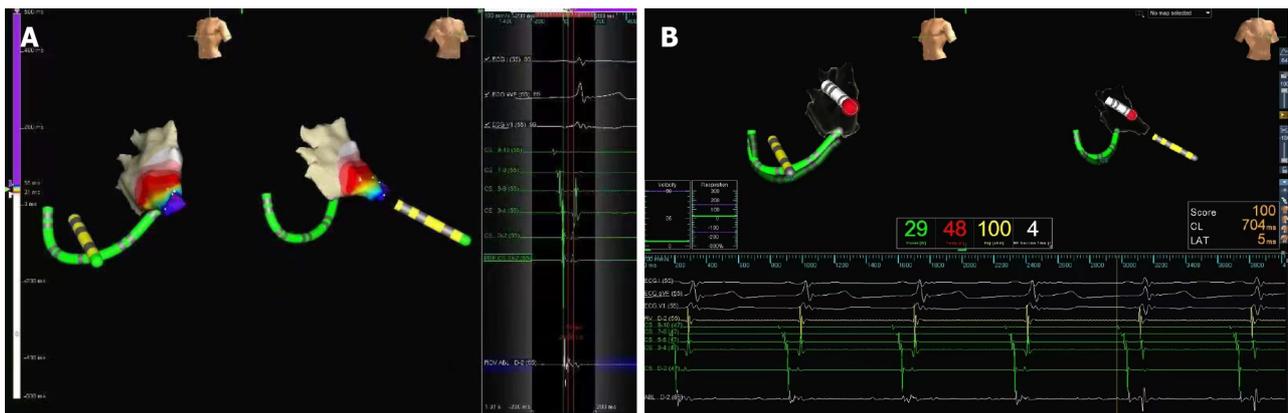
Table 1 Marfan syndrome score of the child

	Wrist and thumb sign	Pectus excavatum	Myopia > 3 diopters	Scoliosis	Mitral valve prolapse
Score	3	1	1	1	1



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Figure 1 Cardiac computed tomography scan and reconstruction showed severe aortic root dilatation resulting in a narrow septum.



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Figure 2 Cardiac electrophysiological examination and catheter ablation were performed guided by an Ensite NavX system. A: A satisfactory target was marked on the atrial side at 3:30 (o'clock) on the mitral annulus; B: The body surface pre-excitation pattern disappeared.

DISCUSSION

MFS is an autosomal dominant disorder of the connective tissue that usually occurs due to mutations in the FBN1 gene and has an incidence of 1.5–17.2 per 100000[3]. About 50% of MFS patients have aortic dilatation in early childhood that gradually progresses. Aortic root disease, which was initially considered to be the main cause of complications and death in patients with MFS, became the target of most treatments and clinical studies. However, the German Marfan Organization believes that some manifestations of MFS may seriously threaten the health of patients, including SVT, ventricular tachycardia, and repolarization abnormalities[4]. Wafa *et al*[5] showed that the burden of inpatient arrhythmia in MFS is higher than the general non-Marfan population and that SVTs were significant predictors of higher adjusted health care-associated cost. In addition, early aortic repair is recommended by the Ghent guidelines[2], and surgical treatment should also be considered in patients with aortic aneurysm or valvular insufficiency. Postoperative scarring both increases the incidence of arrhythmias and poses difficulties for catheter ablation, however. Demolder *et al*[6] demonstrated that heart failure and arrhythmia have become the primary causes of death for patients with MFS after aortic surgery. Therefore, eradicating arrhythmias in patients with MFS in childhood who have not undergone intracardiac surgery has the potential to save lives. Furthermore, with modern the improvements in aortic repair, the treatment of arrhythmias in children with MFS should be the focus of future research.

The TS is preferred for catheter ablation to avoid aggravated aortic injury when there is a left-sided AP. The TS does not enter the left side of the heart through the aorta, which effectively avoids the risks of aortic valve injury and ascending aortic aneurysm rupture, among others. Moreover, for children with APs located in the left posterior lateral or left posterior septum, the operation of the ablation catheter is more flexible by a TS[7]. In adults, a TS is associated with a higher success rate and lower recurrence and complication rates, and the safety and feasibility of this method have also been demonstrated by recent studies. Yoshida, Rami, and Koca showed that TS for ablation was safe and effective even in children weighing less than 30 kg[8-10]. However, considering of the need for special training and the risk of serious potential complications (cardiac tamponade and pericardial effusion), TS is still not recommended by most cardiovascular pediatricians at present[11].

Our center at Beijing Children's Hospital is a national children's medical center, and our staff are experienced in ablating left-sided AP in children using a TS. The child in this report was the only one with left-sided AP who had confirmed MFS, however. She had mitral and tricuspid regurgitation, pectus excavatum, previous history of thoracic surgery, and may also have had a potential delay in vascular healing and repair that presents in most patients with MFS. These factors not only increase the risk of arrhythmia but also pose difficult challenges for atrial septal puncture and radiofrequency ablation[12,13].

The complications of atrial septal puncture include pericardial effusion and cardiac tamponade, as well as others. Cardiac ultrasound, cardiac CT scan and reconstruction must be performed to understand fully the shape and degree of aortic dilatation and cardiac chamber size. Although the child had mitral and tricuspid regurgitation, fortunately the atrial size was normal and had little change at the puncture site. However, her noncoronary sinus, located in the anterosuperior aspect of the interatrial septum, was significantly dilated and pushed against the septum, resulting in a narrow septum. In a case like this the needle tip should therefore be slightly oriented away from the aortic root during puncture (moved backward and downward compared to normal children) to avoid perforating the aorta. In addition, the child weighed < 30 kg, leading to a small heart size. To reduce the risk of penetrating the heart chamber we increased the curvature of the puncture needle.

For safety, a small amount of radiation is required during the puncture. Transseptal puncture under fluoroscopy provides real-time images of surrounding organs (trachea and spine) and CS electrodes, which is helpful in locating the puncture site. However, zero-radiation transseptal puncture can be achieved with the assistance of other imaging tools, such as transthoracic echocardiography, transesophageal echocardiography, or intracardiac echocardiography, but these methods have rarely been used in children[14].

Children are more sensitive to radiation, which may increase their risk of developing hematologic or intracranial tumors[15]. Thus, our subsequent ablation procedure was guided by 3D mapping to minimize the radiation dose while ensuring safety and efficacy. Radiofrequency ablation complications can include valve injury, atrioventricular block, cardiac tamponade, and thromboembolism. Furthermore, valve disease associated with MFS can increase the risk of thromboembolism. Because of the patient's tricuspid regurgitation, to avoid aggravation of injury, gentle operation low movement amplitude of the ablation catheter at the valve site were necessary. Additionally, in the presences of mitral regurgitation, careful mapping should be performed to determine the ablation site in order to reduce the potential damage of repeated discharges to the valve. For the patient in this case report, echocardiography was performed after catheter ablation and showed that valve damages were not aggravated.

Since the operation, there has been no recurrence of SVT, and no complication have been found during the 1-year follow-up period. The child has gone to school and lived a normal life. For children with aortic diseases such as MFS, who cannot undergo catheter ablation by TA, a TS can be attempted in experienced medical centers. However, infantile MFS is more severe than classical MFS[16]. Severe mitral valve disease and small heart size make ablation much more difficult, and atrial enlargement also causes changes in the puncture site, increasing the risk of accidental atrial or aortic penetration. Weight < 15 kg and age < 4/5 years have also been used as criteria for high risk of catheter ablation in previous studies and guidelines[17]. The weight of this child was large enough so that our conclusions are not applicable to infantile MFS; subsequent studies are needed for using a TS in infantile MFS patients specifically.

CONCLUSION

In conclusion, it is necessary to perform catheter ablation for MFS patients in order to eliminate SVT in childhood before intracardiac surgery. TS for catheter ablation of left-sided APs, appears to be safe, effective and applicable in older children with MFS, can certainly be attempted in experienced medical centers. Increasing the curvature of the puncture needle and moving the tip posteriorly and inferiorly may be safer for children weighing < 30 kg who have significantly dilated aorta. Gentle manipulation is needed to avoid damaging the valve during ablation.

FOOTNOTES

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