Catheter ablation induced phrenic nerve palsy by pulsed field ablation—completely impossible? A case series

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Background
Pulsed field ablation (PFA) is a new feasible and safe method for the ablative treatment of cardiac arrhythmias, such as atrial fibrillation (AF). Through the use of electric fields, it causes pore-like openings in the cell’s wall, leading to cell death. The most appealing characteristic of this new technique is its selectivity for cardiomyocytes and consequently its low risk of collateral damage to extracardiac tissues. We present three cases of a PFA-induced transient phrenic nerve (PN) injury documented during pulmonary vein isolation (PVI).

Case summaries
Three patients aged 55–81 years underwent PFA for symptomatic AF. Cases 1 and 3 were affected by paroxysmal AF without evidence of structural heart disease. Case 2 had persistent AF and ischaemic cardiomyopathy with preserved ejection fraction. We observed a transient right hemidiaphragm palsy during the delivery of impulses in the right superior pulmonary vein (Cases 1 and 2) and in the right inferior pulmonary vein (Case 3). The palsy lasted <1 min and was followed by spontaneous full recovery in all cases.

Discussion
Transient PN dysfunction can be observed following PFA in AF ablation. According to our initial experience, a full recovery of the PN function can be expected within seconds. We hypothesize a hyperpolarization of neuronal cells or a depletion of acetylcholine in the motoric endplate to explain this event. Further studies are required to understand the exact pathophysiological mechanism.

Keywords
Atrial fibrillation • Pulsed electric ablation • Phrenic nerve dysfunction • Pulmonary veins isolation • Case report • Case series

ESC Curriculum
5.3 Atrial fibrillation • 5.4 Atrial flutter • 5.5 Supraventricular tachycardia

Learning points
• Pulsed field ablation is a new ablative technique highly selective for cardiomyocytes, nevertheless transient phrenic nerve palsy can be observed. Hyperpolarization of neuronal cells or depletion of acetylcholine in motoric endplate are possible explanations but further studies are needed to clarify this phenomenon.
• The phrenic palsy was followed by full spontaneous recovery with a few seconds, confirming the safety of this new technique.

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Introduction

Pulsed field ablation (PFA) technology is also called irreversible electroporation (IRE) and offers some potential advantages over traditional thermal ablative techniques.1 A major limitation of common ablation technologies refers to the non-selective nature of the destructive process with potential injuries to neighbouring structures. In particular, phrenic nerve (PN) palsy (PNP) is a typical complication during cryoballoon ablation. Conversely, IRE can create irreversible lesions, specifically in the myocardium, with a low incidence of damage to the extracardiac structures.2 This new modality consists of microsecond electrical pulses applied across the cells to generate an electric field, causing the formation of permanent defects in the cell membrane. The permanent permeability of the cell membrane leads to changes in the cells’ homeostasis and the death of the cell. Every tissue has a different specific field threshold that induces necrosis. Cardiomyocytes have the lowest threshold value (400 V/cm), and this may constrain collateral damage (e.g. oesophagus or PN). There is no coagulative necrosis, but evidence of myofiber disruption and inflammation, resulting in a more homogenous fibrotic remodelling than radiofrequency ablation. In addition, IRE is able to induce faster cell necrosis and therefore, the time of the procedures is shorter in comparison with other standard ablative techniques.3 Although nerve damage is not an expected complication during PFA, we can report the first three cases of transient self-recovering PNP related to this technique.

We performed pulmonary vein isolation (PVI) in the following cases with the PFA technique using our standard approach. The ablations were performed under conscious sedation with a bolus of midazolam and fentanyl followed by a continuous infusion of propofol, according to the position paper of the German Society of Cardiology on cardioanalgesia.4 After a single transseptal puncture, we performed the procedure and the access site was closed with a ‘Figure 8’ suture in all cases. Amiodarone was disrupted after the ablation.3

Case presentations

Case 1

A 55-year-old male was referred to our department for symptomatic paroxysmal AF. His body mass index (BMI) was 22 (190 cm x 78 kg) with no history of previous heart disease. The patient had a good clinical status with no findings of heart failure. Arterial blood pressure was 120/80 mmHg at admission. The echocardiogram showed a structurally normal heart, with preserved ejection fraction (EF) (60%), no significant valvular diseases and the left atrial diameter was 44 mm. The patient was treated with beta-blockers and no other antithrombotic drugs. The patient had a low cardioembolic risk (CHA2DS2-VASc Score 0) and he was treated with low-molecular-weight-heparin for 3 weeks before the ablation and for 2 months after the ablation.

During the first application of PFA at the RSPV with 35 mm Farawave catheter, we noted a loss of function of the right sided PN (Figure 2A and B; Supplementary material online, Videos 1 and 2). We still completed all planned energy deliveries to the RSPV checking fluoroscopically the diaphragmatic movements after each application. We observed full recovery 6 s after the last application (Figure 2C and D; Supplementary material online, Video 3), therefore, PNP lasted ca. 40 s overall.

The procedure ended without side effects. The PN dysfunction was resolved completely and the pulmonary veins were all correctly electrically isolated. The procedure took 36 min. After the ablation, the same medical therapy was prescribed.

Case 2

A 68-year-old male was referred to our department for symptomatic persistent AF. His BMI was 26 (178 cm x 83 kg). He was affected by arterial hypertension and Type II diabetes, complicated by peripheral neuropathy. He had a history of coronary artery disease with previous percutaneous coronary revascularization. The patient had no signs and symptoms of congestive heart failure and arterial blood pressure was 140/80 mmHg at admission. The echocardiogram showed a preserved EF (55%) without abnormalities of the regional wall motion and no significant valvular diseases. His left atrial size was 41 mm. The patient was treated with beta-blockers, amiodarone, and apixaban.

We performed PVI via the PFA technique with our standard approach. In this case, we used the 31 mm Farawave catheter, eight impulses were delivered (1 + 7) at all PVs that were isolated after the first impulse (2.0 kV). During the employment of PFA at the RSPV, we noted the same transient palsy of the right PN, self-recovering after few seconds. The procedure was successful and no complications were observed. Amiodarone was disrupted after the ablation.

Timeline

| Patient number | Before the procedure | During the procedure |
|----------------|----------------------|----------------------|
| 1              | The patient is a 55-year-old male affected by symptomatic paroxysmal atrial fibrillation (AF). Pulmonary vein isolation with PFA was performed. | Transient right hemidiaphragm palsy was observed after the first application of the impulses at the RSPV. We still continue to deliver the eight planned impulses. We observed the full recovery of the hemidiaphragm function 6 s after the last application. The PNP lasted overall around 40 s. |
| 2              | The patient is a 68-year-old male with symptomatic persistent AF and ischaemic heart disease. Ablation of AF was performed with the PFA technique. | During the delivery of the pulses at the RSPV, a transient PN dysfunction of few seconds occurred, followed by spontaneous recovery. |
| 3              | An 81-year-old male affected by symptomatic paroxysmal AF underwent isolation of the pulmonary vein with PFA. | A transient PN dysfunction was observed during the erogation of impulses at the RIPV. In this case, the phrenic palsy lasted few seconds as well, followed by spontaneous full recovery. |
**Figure 1** Farawave catheter in ‘basket’ (A) and ‘flower’ (B) configuration at pulmonary vein ostium.

**Figure 2** (A and B) Paradoxical movement of the right hemidiaphragm immediately after the application of the impulses. (C and D) Right and left hemidiaphragms contract synchronously after 6 s. FW, Farawave Catheter, CS Inquiry Catheter 10 mm in coronary sinus.
Case 3
A 81-year-old male with no history of heart disease was referred to our department for symptomatic paroxysmal AF. His BMI was 24 (192 cm × 87 kg). The patient had no signs and symptoms of congestive heart failure and arterial blood pressure was 130/70 mmHg at admission. His echocardiogram showed a structurally normal heart (the EF of the left ventricle was 60%) and left atrial diameter was 42 mm. The patient was treated with beta-blockers and edoxaban.

We performed PVI with the PFA technique according to our standard approach. With a 35 mm Farawave catheter, four impulses in the flower configuration and four in the basket configuration that were isolated after the first impulse (2.0 kV) were delivered in each pulmonary vein. During the delivery of pulses at the RIPV, a transient palsy of the right PN was documented, followed by a full recovery within few seconds. The same medical therapy was continued after the ablation.

Discussion
Pulsed field ablation induces selective myocytes necrosis, without collateral damages to other tissues such as the oesophagus, the PN, or the endothelial cells.2,3 In our case, we clearly documented a transient PN dysfunction causing a transient paresis of the right hemidiaphragm. This effect would not be expected, considering that application of PFE is highly selective for cardiomyocytes.

The response of nerve tissue to the IRE has not been clearly defined. In the last decade, different scientists studied the effects of IRE on nerve tissue. Onik et al.4 in their study on canine models, found that nerves were apparently preserved after the application of IRE pulses. Weil et al.5 performed a pilot study in this field with electrophysiological, histological, and functional analysis on sciatic nerve models of rats. They showed that nerves treated with IRE were damaged after immediate direct injury with a full recovery after 2 weeks. The pathophysiological changes were typical Wallerian degeneration and regeneration of axons.

According to the Seddon’s classification, the mildest level of peripheral nerve damage is ‘neuapraxia’ (characterized by focal segmental demyelination at the site of injury without disruption of axon continuity and its surrounding connective tissues) that requires at least several weeks to obtain a complete functional recovery.6

In our case, the dysfunction was transient and passed within few seconds, so we assume a different pathophysiological mechanism based on this phenomenon.

We assume that the transient nerve dysfunction could be related to hyperpolarisation of the nerve. This hypothesis is supported by previous experimental evidence. Rems et al. showed that IRE and its electric field create not only pores in the membrane lipid bilayer, but also electroconformational changes of the voltage-gated ion channels. An ionic imbalance through the membrane cell occurs, which later becomes hyperpolarized and unable to conduct action potentials (AP).8

A second possible explanation could be the effect of IRE on the motric endplate. Conventionally, upon the arrival of an AP at the axon terminal, voltage-dependent calcium channels open and calcium flows from the extracellular fluid into the cell. This intracellular current triggers a biochemical cascade that induces the release of acetylcholine into the synaptic cleft that depolarizes the muscle fibres, leading to force generation. In this scenario, a transient perturbation of the Ca2+-ions flow induced by the electric field of IRE could lead to a decreased release of acetylcholine and to the subsequent inability to generate an effective muscular activation.9

Some authors suggested to test the PN by stimulation of the SVC before, during, and after performing the PFA application at the sepal pulmonary veins.10 Due to the very short application time, this monitoring of the PN function was not performed in our series.

Conclusions
Transient PN dysfunction can be observed following PFA in AF ablation. According to our initial experience, a full recovery of the PN function can be expected within seconds. Pulsed field ablation appears to be a safe technique without long-term sequelae. Further studies and experience are needed to better evaluate the risk of PN damage as a clinical complication.

Lead author biography
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Supplementary material
Supplementary material is available at European Heart Journal—Case Reports online.

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Slide sets: A fully edited slide set detailing this case and suitable for local presentation is available online as Supplementary data.

Consent: The authors confirm that written consent for submission and publication of this case report, including images and associated text, has been obtained from the patients in line with the COPE guidance.

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