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1 **COVER SHEET**

2

3 **TITLE**

4 Percutaneous electrochemical debridement of the Plantaris tendon: a novel option in the treatment of midportion Achilles
5 tendinopathy.

6

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22

23 **Authors' contribution:** GM and CM were responsible for the study concept and design. GM drafted the manuscript.

24 All authors approved the final manuscript.

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**ACCEPTED
MANUSCRIPT**

2
3 **TITLE**

4
5 Percutaneous electrochemical debridement of the Plantaris tendon: a novel option in the treatment of midportion Achilles
6 tendinopathy.

7
8 **ABSTRACT**

9
10 Plantaris tendon disorders are a well-known source of midportion Achilles tendinopathy. Plantaris tendon thickening and
11 fibrous tissue formation between the tendons are the histological abnormalities which are typically observed. Surgical
12 approaches (scraping of the Achilles medial and ventral paratendinous tissues and excision of the Plantaris tendon) have
13 already shown good clinical outcomes; despite this, cost-benefit ratio of these interventions may be unfavourable and
14 their accessibility is limited. Percutaneous needle electrolysis is a minimally invasive ablative technique of increasing
15 consideration in the treatment of tendinopathies and associated conditions. The purpose of this article is to introduce a
16 novel procedure to treat Plantaris tendon-related midportion Achilles tendinopathy. The procedure starts with the insertion
17 of a non-coated needle (diameter: 0.30-0.40 millimetres; length: 30 millimetres) between the Plantaris and Achilles
18 tendons, under ultrasound guidance. Subsequently, galvanic current (intensity: 2 mA) is locally transferred. This, in turn,
19 induces instant non-thermal electrochemical ablation of the intertendinous tissues in close proximity to the needle, finally
20 debriding the Plantaris tendon. In order to further promote its release, second part of the procedure involves partial
21 tenotomy of the lateral peripheral aspects of the Plantaris tendon. Usually, the total duration of the session does not exceed
22 thirty minutes. Percutaneous needle electrolysis may be considered as a valid alternative to surgery. The out-patient
23 procedure presented in this article is, in fact, safe and quick to perform. Additionally, long suspension of working or
24 sporting activities after the treatment is not required. Future investigations are needed to ascertain the short- and long-
25 term therapeutic outcomes in the treatment of Plantaris tendon-related midportion Achilles tendinopathy, in particular by
26 comparing them with those obtained with other mini-invasive interventions.

27
28 **KEYWORDS**

29
30 Ablative techniques – Electrolysis – Interventional ultrasonography – Tendon injuries

1 **INTRODUCTION**

2

3 The role of the Plantaris tendon in the etiopathogenesis of midportion Achilles tendinopathy has been certified
4 over the last decade [1–3]. However, research of highly specific diagnostic modalities and optimal treatment strategies is
5 still ongoing. Plantaris tendon-related midportion Achilles tendinopathy is clinically characterised by debilitating pain
6 and swelling, which are typically localised in the medial aspects of the Achilles tendon body [2–3]. Friction and
7 compression traumas between the tendons are likely to be the biomechanical disorders that lead to the histological changes
8 observed in many researches, such as Plantaris tendon thickening, fibrous tissue formation between the tendons and
9 alteration of the vascularisation and innervation patterns of the Achilles paratenon [1–4].

10 Surgical scraping of the Achilles medial and ventral paratendinous tissues has shown good clinical results in many
11 trials, especially when associated with excision of the Plantaris tendon [5–8]. Despite this, cost-benefit ratio of these
12 interventions may be unfavourable and their accessibility is relatively limited. Additionally, post-surgery rehabilitation
13 protocols may last several months. As a consequence of this, there is need to identify new therapeutic solutions which
14 may be as effective as surgery but without having these relevant weak points.

15 Thus, the main purpose of this article is to introduce the debridement of the Plantaris tendon via electrochemical
16 ablation, induced by cathode-centred percutaneous needle electrolysis. The latter is an ultrasound-guided and minimally
17 invasive technique which may be considered a valid alternative to surgery or, at least, a treatment option to contemplate
18 before performing it. In support of this hypothesis, minimally invasive techniques (needle scraping or sclerosing
19 polidocanol injections) have already shown encouraging clinical results in the treatment of Plantaris tendon-related
20 midportion Achilles tendinopathy [9] and, on the other hand, percutaneous needle electrolysis is generally of increasing
21 consideration in the treatment of tendinopathies and associated conditions [10–13].

22

23 **PERCUTANEOUS NEEDLE ELECTROLYSIS**

24

25 This study was designed and conducted according to national and international standards and in compliance with the
26 Helsinki Declaration and the International Principles governing research on humans. Considering the typology of this
27 article (clinically illustrated – technical note), *material and methods* and *results* sections are neither required nor
28 presented.

29

30 *Equipment*

31 The Authors apply the technique using a specifically developed and medically certified device (EPI Advanced

1 Medicine®, Barcelona, Spain; directive 93/42/EEC). This instrumentation permits intratissue galvanic current transfer,
2 at settable intensities, through an appropriate non-coated needle (diameter: 0.30-0.40 millimetres; length: 30 millimetres;
3 same manufacturer as above). While the needle acts as the cathode, the anode can be handled by the patient or applied on
4 the skin. The cathodic flow is the only one that is used during the procedure (cathode-centred electrolysis). When the
5 current is transferred, the basic electrochemical process of saltwater electrolysis instantly develops, inducing the non-
6 thermal ablation of the tissue in close proximity to the needle. The latter is inserted under ultrasound guidance in order to
7 precisely treat the target tissue, without involving other structures. For this purpose, the Authors use the GE Healthcare®
8 Logiq S7 Expert ultrasound equipped with the ML6-15 (50mm; 6-15 MHz) and L8-18I-D (25mm; 8-18 MHz) linear
9 probes.

10

11 *Preliminary ultrasound investigation*

12 The patient lies on his or her side, with the medial aspects of the Achilles tendon directed upwards. The region is
13 shaved and disinfected by applying a proper protocol. A preliminary ultrasound is carried out in order to accurately detect
14 the portions of the Plantaris tendon in anatomical relationship with the medial aspects of the Achilles tendon body. It may
15 be helpful to delimit the region to be treated, marking its distal and proximal limits with a sterile dermatographic pencil. It
16 is also advisable to mark the points at which the patient complains about having more pain and swelling (“critical areas”)
17 and where the major anatomical alterations are discernible (figures 1–2).

18

19 *Description of the procedure*

20 The procedure is graphically represented in figure 3. First, the needle is inserted between the Plantaris and Achilles
21 tendons, under ultrasound guidance (figure 4). Subsequently, the galvanic current is transferred (intensity is pre-set to
22 2mA). Doing this, the local ablation of the fibrous intertendinous and Achilles paratenon tissues is instantaneously
23 obtained, anatomically debriding the Plantaris tendon (figure 5). The single applications of current last 2-3 seconds.

24 Then, the needle is partially withdrawn and pointed toward the Plantaris tendon. In order to further promote its
25 release, a partial tenotomy of the lateral peripheral aspects of the tendon is performed (figure 6). To this effect, the single
26 application can have a variable duration, between 2-6 seconds, depending of the mechanical resistance offered by the
27 tendinous tissue to the needle penetration (the lower the resistance, the shorter the application).

28 All the actions presented above are repeated approximatively every five millimetres (or less, in the “critical areas”;
29 see above), in distal-proximal direction, throughout the region previously skin-marked. Typically, the total duration of
30 the session does not exceed thirty minutes (including disinfection and dressing processes).

31

1 *Tolerability of pain and side effects*

2 The insertion of the needle typically cause minimal discomfort. By contrast, the patients may experience moderate
3 strong pain during the applications of galvanic current. Anyway, anaesthetics are usually not locally injected before
4 percutaneous electrolysis, since the procedure is generally well-tolerated by the patients (the single galvanic current
5 application can be stopped at any time if the pain is not bearable) and because the use of syringes would substantially
6 increase the overall invasiveness of the intervention. Anyway, use of anaesthetics remains a considerable option. Relevant
7 vagal reactions during and immediately after the intervention are possible [14]. Bleeding in area of needle insertion and
8 intervention-related discomfort in the treatment area (up to 48 hours) are the most common side effects. Infection-related
9 issues are extremely rare, as the technique is minimally invasive and the electrolytic process has a substantial bactericidal
10 effect.

11

12 **DISCUSSION**

13

14 The main purpose of this article is to introduce the debridement of the Plantaris tendon, via electrochemical
15 ablation induced by cathode-centred percutaneous needle electrolysis, to treat Plantaris-related midportion Achilles
16 tendinopathy. This novel procedure permits to eliminate the fibrous tissue interposed between the Plantaris and the
17 Achilles tendons, debriding the Plantaris tendon and improving the local biomechanics.

18 The main practical value of this technique is the possibility of performing it in out-patient clinics, reducing
19 considerably the costs, the waiting lists-related issue and the other implication and side effects of Achilles tendon surgery,
20 such as suture reactions, incisional neuromas, and granuloma formation [15]. Furthermore, it should be reminded that
21 post-surgery protocols may last several months. On the contrary, consistent with our experience, in particular with
22 professional football (soccer) player, it is not necessary to completely suspend the sporting (or working) activities for
23 more than 24-48 hours, after percutaneous needle electrolysis treatment. In fact, the side effects tend to be very moderate.
24 However, many authors that use this technique, including us, find it helpful to implement a complementary protocol of
25 active physical therapy [10–13]. Preliminary studies carried out by our research group indicate that the short-term clinical
26 results (not yet published) after percutaneous needle electrolysis treatment in professional athletes are very promising but
27 that it commonly necessary to conduct at least 3-5 sessions (one per week) to obtain long-lasting results. However, to
28 date, clinical or imaging predictors of outcome are substantially unknown.

29

30 **CONCLUSION**

31

1 Percutaneous needle electrolysis is an ultrasound-guided and minimally invasive technique that allows specific
2 treatment of the anatomical alterations that cause Plantaris tendon-related midportion Achilles tendinopathy. Since it is
3 safe and quick to perform, it may be considered as a valid alternative to surgery. Future investigations are needed to
4 ascertain the short- and long-term therapeutic outcomes in the treatment of Plantaris tendon-related midportion Achilles
5 tendinopathy, in particular by comparing them with those obtained with other mini-invasive interventions.

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15

16 **FIGURES**

17

18 **Figure 1.** Plantaris tendon morphological abnormalities are commonly observed in case of mid-portion Achilles
19 tendinopathy. The transversal, grey-scale, high resolution ultrasound image presented shows increasing of the thickness
20 and width of the Plantaris tendon (indicated by the *dashed arrow*). In order to complete an accurate investigation, it is
21 advisable to compare the tendon dimensions with those of the contralateral one and to take into high consideration the
22 data concerning the normal morphology of the Plantaris tendon presented by Olewnik et al [1]. *Ach* = Achilles tendon; *K*
23 = Kager's fat pad. Probe used: ML6-15 (50mm footprint).

24

25 **Figure 2.** Transversal, high resolution ultrasound image showing high blood flow (*filled arrows*) in the medial aspects of
26 the Achilles tendon (*Ach*) and paratenon, between the Achilles and Plantaris tendon (indicated by the *dashed arrow*) and
27 around the latter. In case of mid-portion Achilles tendinopathy, these signs may be discernible also in the ventral portions
28 of the Achilles tendon [2]. Probe used: ML6-15 (50mm footprint).

29

30 **Figure 3.** Schematic diagram of the percutaneous needle electrolysis procedure, for the debridement of the Plantaris
31 tendon. The needle is initially inserted between the Plantaris (*dashed arrow*) and the Achilles (*Ach*) tendons, as indicated

1 by the *line 1*, and, subsequently, within the lateral peripheral aspects of the Plantaris tendon, as indicated by the *dashed*
2 *line 2*. *Asterisk* = tibial neurovascular bundle.

3

4 **Figure 4.** Ultrasound-guided insertion of the needle between the Achilles and the Plantaris tendons. Considering their
5 anatomical relationship, the preferred approach is in posterior-to-anterior and medial-to-lateral direction. Inclination of the
6 needle is variable and depends upon the specific morphological features of the tendons.

7

8 **Figure 5.** An iperechoic area (*arrows*) is typically observable around the needle immediately after the application of
9 galvanic current and consequent development of the electrochemical process. This should confirm that only the
10 intertendinous tissues have been treated, without involvement of other structures (first part of the procedure). *Dashed*
11 *arrow*=Plantaris tendon; *Ach*=Achilles tendon. Probe used: L8-18I-D (25mm footprint).

12

13 **Figure 6.** After the ablation of the intertendinous fibrotic tissues, the needle (*asterisks*) is partially withdrawn and inserted
14 in the lateral peripheral aspects of the Plantaris tendon (second part of the procedure). *Dashed arrow*=Plantaris tendon;
15 *Ach*=Achilles tendon. Probe used: L8-18I-D (25mm footprint).

16

17 **COMPETING INTERESTS:** none.

18

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