Evaluation of the benefit of using a combination of autologous platelet rich-plasma and hyaluronic acid for the treatment of Peyronie’s disease

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Abstract

Background: Platelet-Rich Plasma (PRP) has been successfully used for different medical conditions, to help regenerate fibrotic or inflammatory tissues. Peyronie’s disease (PD) could be a target for such treatment. In this study, PRP in combination with hyaluronic acid (HA) is proposed as an innovative treatment for this condition.

Objectives: To assess the effects of repeated intralesional injections inside the tunica albuginea (TA) with a combination of PRP and HA in PD patients.

Design, setting, and participants: 90 patients, naive of any prior interventional treatment, were injected with 8 ml of PRP combined with HA directly in the diseased area of TA under ultrasound (US) guidance. Injections were repeated 4 times within 2 months. Additional monthly sessions were done, if necessary.

Outcome measurements and statistical analysis: Penile deformation, TA thickening, presence and size of calcifications, and questionnaires evaluating both PD and sexual function. Global results were classified into satisfactory improvement (I+) unsatisfactory (I-) and unchanged (U).

Results and limitations: Angulation and thickening were both significantly improved after 4 sessions and continuously improved with the number of sessions, to achieve a mean final improvement of 39.65 +/-24.83% (p<0.001) (CI 14.26;18.82) and 26.23+/-10.29% (p<0.001) (CI 0.99;1.22), respectively. Younger patients achieved better results. 73.3% of the patients belonged to I+ group; 67.8% felt an improvement, 62.2% a decrease of their deformation, 46.7% had easier sexual intercourses and 70% judged the treatment as positive. Limitations included the absence of a control group.

Conclusions: Repeated intralesional injections of PRP-HA improve significantly penile deformation and thickening of the TA. This fully natural treatment may stand as a first line treatment of PD.

Patient summary: We evaluated here an innovative treatment of Peyronie’s disease using intralesional injections of Platelet-Rich Plasma mixed with hyaluronic acid. We observed a significant improvement of the deformation in 70% of the cases without any serious adverse events.

Introduction

The treatment of penile deformities first described in 1743 by King Louis XV’s first surgeon François Gigot de Lapeyronie [1] has gained much interest especially for non-surgical procedures [2]. An increasing knowledge of the pathophysiology [3] of the disease has urged clinical researchers to propose intralesional treatments [4] such as verapamil [5] and interferon alpha 2B [6] which remain off label, or collagenase clostridium histolyicum (CCH) (Xiapex®) which obtained an FDA approval [7] for Peyronie’s disease (PD). The interesting improvement rates observed with these treatments are challenged by side effects such as corporeal ruptures for CCH, flu-like symptoms for interferon alpha 2B, or penile bruising, dizziness and nausea for verapamil.

Moreover, a wider use of Xiapex® is to date limited by the high cost of the compound. More recently, Hyaluronic Acid (HA) (7) and Botulinum toxin [8] have also been proposed with the purpose of counteracting tunica fibrosis, the final issue of the pathological process of PD, before the deposit of calcifications [4]. We hypothesized that regenerative medicine could help stop the process and restore function. Although stem cell therapy for erectile dysfunction (ED) [9] and PD [10] has become a wide subject of research in animal studies with rising hopes for future treatments with adipose derived stem cells (ADSC’s), we did not find in the literature any mention of the potential use of autologous platelet-rich plasma (PRP) in both ED and PD. PRP is prepared from autologous blood after centrifugation with an adequate medical device giving a plasma with a higher platelet concentration than baseline. PRP is a source of platelets growth factors (GFs). In situ activation of platelets induces the orchestrated release of several growth factors such as FGF(fibroblast growth factor), PDGF (platelet-derived growth factor), IGF (insulin-like growth factor), VEGF (vascular endothelial growth factor), and TGF-b (transforming growth factor) [11]. Platelet GFs are signals that regulate cell growth and division, as well as synthesis of extracellular matrix for tissue regeneration. On the other hand, plasma is a source of proteins such as fibrinogen, vitronectin and fibronectin which play a key role in the

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regulation of cell-cell interaction and cell spatial organization [12].
Autologous PRP has been widely and safely used in many disciplines
including orthopaedics, maxillofacial surgery, plastic and reconstructive
surgery [13]. Adding thrombin to PRP produces a platelet gel which
can be used as biological glue. One of us (RV) has been using it during
graft procedures for PD surgery (unpublished data). From these early
experiences, we became interested in an intramuscular non-surgical use
of PRP for PD. Okabe et al. [13] have shown experimentally that the
addition of HA to PRP enhanced its regenerative potential, acting as a
biological scaffold. The combination of PRP and HA, now facilitated by
the availability of the new medical device CellularMatrix, encouraged
us to conduct a first trial on 13 patients with PD [14]. Preliminary data
from this case series indicated that patients experienced a significant
improvement of their condition, both functionally and anatomically.

Stimulated by these early encouraging results, we designed a new
study using extensive clinical (the Peyronie’s disease questionnaire
PDQ), echographic and photographic assessments as outcome
measures in 90 patients whose results we report here are, to the best of
our knowledge, the first in the English literature.

Patients and methods

Between July 2014 and November 2016, 90 patients (aged 25 to 77y,
median 56) complaining of penile deformations and/or curvatures
linked to the presence of plaques palpated in the tunica albuginea (TA),
and who had not received any previous local treatment were
enrolled in this study. All patients gave written informed consent.

Patient’s evaluation criteria

Patient’s eligibility was assessed on the basis of the onset and
duration of the disease, initial local trauma, and presence of ED assessed
by the IIEF-5 [15], PDQ questionnaires [16] evaluate symptom’s bother.
Deformation, consisting of curvatures (angulation in degrees) and/or
shaft strictures was measured on photographs of fully errected penis [7].

Duplex ultrasound (DUS) (General Electric Logic F8 with an L12
RS probe) measurements of the thickening of the TA and presence
of calcifications were assessed by one of us (HS) with a fixed protocol
at each session of treatment (see appendix 1). In addition, patients
responded to a self-administered questionnaire.

Treatment protocol

It is an office procedure under a 3-step penile local anesthesia: -
suprapubic application of lidocaine and prilocaine (EMLA, Astra Zeneca) 20 minutes prior to the procedure; - penile block with 10
ml of 2% lidocaine; - additional local application of a cream mixing
lidocaine, tetracaine and tetracaine hydrochloride (Receptura
Arzneimittel) on the injection sites. Four ml of PRP and 4 ml
of a combination of PRP/HA were prepared with a Regen BCT tube
and a BCT HA tube (Regen Lab SA, Switzerland). Autologous venous
blood was collected in both tubes and centrifuged for 5 minutes at
1500 g. At the end of the centrifugation, red blood cells were trapped
under a separator gel, while platelets were settled on its surface. After
appropriate resuspension of platelets in plasma, both preparations were
collected in a syringe and mixed together. Eight ml of this preparation
(containing 6 ml of PRP and 2 ml of HA) were injected into the TA’s
diseased sites under DUS control (Figure 1). Prior to injections,
mechanical action was applied to the fibrotic and/or calcified plaques
with 22G and/or 18G needles (BD Microlance™ 3). At the end of
the procedure, a slightly compressive dressing was applied. Intercourse
was prohibited for 48 hours. Injections were repeated 4 times every 15
days. A first evaluation took place 3 months after the first treatment
session. According to the outcomes, two additional sessions were
scheduled monthly, and repeated each three months, if necessary. All
patients had at least 5 sessions.

Main outcomes measures

Co-primary endpoints consisted of the percent changes in penile curvature and/or deformity, and in maximum thickness, between
baseline and one month after the fourth treatment session and after a
variable additional sessions (mean 7). The co-primary endpoints were
assessed both globally and according to the severity of deformation
(below 30°, between 30° and 59° and over) and the presence or not of
calcifications (Table 1).

Secondary endpoints were the changes in the 3 domains of the PDQ
and the IIEF-5 and answers to a self-administered 5-item questionnaire
evaluating treatment efficacy, deformation, erection, sexual intercourse
and global assessment of the treatment (for details, see appendix 3).

Age, trauma during intercourse, PD duration, number of treatment
sessions were tested as possible parameters influencing the results. By
grouping these data, we could finally classify the results into 3 groups:
• satisfactory improvement (n=66) I • unsatisfactory improved (n=11)
and bad B (n=13) according to the degree of improvement, thickening
and scores achieved by the self-administered questionnaire.

Statistical analysis

Statistical analysis was performed using SPSS 18.0 for Windows.

One-sample paired t test was used to compare the within-group
change in penile curvature and/or deformity and in maximum thickness (change versus baseline).

The changes in the 3 domains of the PDQ, the IIEF-5 and answers
to the self-administered questionnaire were studied by one-sample
paired t test.

The results were evaluated in relation to age, trauma during
intercourse, PD duration, number of treatment sessions, and presence
of calcifications using a Chi-square test.

Results

Average duration of the disease was 20.8 +/- 19.2 months. 84
patients suffered from a mean curvature of the penis of 44.37 +/-
15.93° and 4 had an isolated stricture of the shaft, while 7 had both
deformations. For 13.3 % of the patients, angulation was below 30°; for
68.9% of them, it was between 30° and 59°, and for 13.3% of them it
was over 59°. ED was present prior to PD onset in 34.4% of the patients.
The maximum average thickening was 4.14 +/- 0.8 mm. All but 6 PD’s
plaques (three lateral, 2 ventral and one septal) were localized in the
dorsum of the TA. Calcifications were present in 31.3% of the patients
(n=28). A direct trauma to the penis was reported by 15 patients
(16.7%). Dupuytren’s disease (2 cases) and ankylosing spondylitis (2
cases) were also recorded. Average number of treatment sessions was
7.09 +/- 2.54.

Both deformation and thickening of the TA one month after
4 sessions (n = 84) and at the last follow-up visit (n = 54) showed
a significant decrease (Table 1). The mean final improvement
was respectively 16.54 +/- 10.51° and 1.11 +/- 0.52 mm with a
mean gain percentage of 39.65 +/- 24.83% and 26.22 +/- 10.29% (Table
1). Presence of calcifications (n=28) did not change significantly the
results but 6 patients (21.4%) experienced a decrease of their US density
(Figure 2). These improvements were already significant after 4 sessions.
Virag R (2017) Evaluation of the benefit of using a combination of autologous platelet rich-plasma and hyaluronic acid for the treatment of Peyronie's disease

Table 1. Percent change in penile curvature and/or deformity and thickness between baseline and after the fourth treatment and additional treatment sessions globally and according to the severity of deformation and the presence or not of calcifications

| | Mean | SD | P value | CI |
|---|---|---|---|---|
| Angulation (after the fourth treatment session) | 279970 | 2101525 | P<0.001 | 9.74 ;14.55 |
| Angulation (additional treatment sessions) | 396546 | 2483378 | P<0.001 | 14.26 ;18.82 |
| Thickness (after the fourth treatment session) | 183582 | 910997 | P<0.001 | 0.67 ; 0.86 |
| Thickness (additional treatment sessions) | 262276 | 1029582 | P<0.001 | 0.99 ; 1.22 |

Presence or not of calcifications

| | Thickness (after the fourth treatment session) | Thickness (additional treatment sessions) | Angulation (after the fourth treatment session) | Angulation (additional treatment sessions) |
|---|---|---|---|---|
| No | Mean 188264 | 274160 | 298156 | 433948 |
| | SD 923874 | 1013316 | 198978 | 2473753 |
| Yes | Mean 173382 | 234850 | 236779 | 308650 |
| | SD 890173 | 1033799 | 2339249 | 2336245 |

Severity of deformation

| | Thickness (after the fourth treatment session) | Thickness (additional treatment sessions) | Angulation (after the fourth treatment session) | Angulation (additional treatment sessions) |
|---|---|---|---|---|
| Below 30˚ | Mean 215625 | 270283 | 296936 | 504425 |
| | SD 951791 | 1159128 | 1899753 | 3171063 |
| Between 30˚ and 59˚ | Mean 192361 | 272862 | 284790 | 384166 |
| | SD 857144 | 904433 | 2203684 | 2439523 |
| Over 59 | Mean 131058 | 217283 | 237155 | 347518 |
| | SD 910650 | 1335961 | 1819088 | 1706414 |

Figure 1. A: Longitudinal view of the dorsal TA and below cavernous body with one artery. In white the needle inserted in the fibrotic TA. B: Imaging of the injection of autologous PRP-HA

Figure 2. Distal Transversal echographic image of dorsal TA in 37 year old patient with PD. A: Initial evaluation. Dorsal TA thickness between 3.6 and 4.2mm with a calcified plaque of the deep layer of TA. B: 10 months later after 10 sessions of PRP-HA: decrease of the average thickening to 2.8mm and the calcified plaque has almost completely disappeared

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(p< 0.001) and increased with the number of sessions (p<0.001). The three domains of the PDQ questionnaire showed significant changes (Figure 2). Considering only patients suffering from ED prior to the onset of PD (n=11), IIEF-5 improved significantly (13.82+/- 3.7 vs 17.91 +/- 3.2) (p=0.008) (CI -6.85;-1.33). Finally, 62.2% of the patients felt a decrease in their deformation, 43.3% judged that their erections were better, 46.7% thought that sexual activity was easier and 67.8% considered that the treatment improved their initial condition. Only one patient stopped the treatment after 4 sessions because of worsening symptoms and was offered surgery. At the end, 73.3 % of the patients belonged to group I with satisfactory results, 12.2% to group II with unsatisfactory results and 14.4% to group III, classified as bad results. In addition, the influence of several factors such as the age, severity of initial angulation and thickening, duration of PD and presence of trauma on the outcomes was also tested. It appears that the number of sessions was higher in Group I (7.32 vs 6.18 and 6.11 for Group I, II and III, respectively). In Group I, mean basic angulation and thickening were lower. Decrease of TA thickening was significantly correlated to the age (p=0.026). It was also found a positive correlation between the decrease of angulation (%) and the TA thickening, after 4 sessions (p=0.015) and at the end of treatment (p=0.011). Neither duration of PD, nor the presence of direct trauma had significant influence on the outcomes.

Complications

Ecchymosis was present in 16.7% of the patients and marked hematoma in 10%.

Two patients suffered from transient hypotension after local anaesthesia.

No serious adverse events were recorded.

Discussion

This study on 90 patients confirms the positive initial conclusions of our preliminary trial [14]. Objective endpoints collected one month after 4 sessions and at the end of an average of 3 to 5 additional sessions showed significant improvement of PD, assessed by the 3 domains of the PDQ questionnaire, as well as by the responses to an additional self-administered questionnaire and to the IIEF-5 (Figure 3). We achieved a mean reduction of the angle of 16.54 °, which represents an average reduction of 39.65%. These results were even better -while non-significant, due to an insufficient number of cases- for the group of patients without calcifications (18.07 +/-10.89°, corresponding to a reduction of 43.59 %)

According to current regulations, PRP must be prepared from fresh blood and used during the same medical session and in the same room (extemporaneous procedure), making it difficult to set-up placebo-controlled studies. Nevertheless, available previous placebo-controlled studies [7] show a maximum angle reduction of 21% with Impress I and of 15.2% for the placebo group. Furthermore, the comparison of our results with those of untreated patients, as in Berneski’s trial [17] conducted on 176 patients to study the natural history of PD in the absence of medical or surgical therapy, further supports a beneficial effect of the combination of PRP and HA, as only 12% of untreated patients had an improvement of their deformation vs 62% in the present study.

A comparison of our data with other intrallesional injection trials, especially both Impress I and II series of the Xiapex® trial [7], shows slightly better results, particularly if only the patients without calcifications as those enrolled in the Xiapex® trial are considered (Figure 4). Moreover, contrary to Xiapex® [18], where it seems that after three cycles of treatment there is no more improvement, addition of sessions of PRP combined with HA after the first cycle of four, improves the results. The reason might lie in the biological process induced by the combination of PRP and HA. Contrary to CCH, which main aim is to destroy the plaque, PRP combined with HA is expected to restore normal tissue after digging of small tunnels into TA’s fibrotic tissue with a needle (see below). HA alone has been shown to be active on PD in a placebo-controlled study [8]. Preliminary data seems to indicate that a combination of PRP and HA provides better outcomes than HA alone for the treatment of gonarthrosis [19, 20] and wound defects [21,22]. PRP and HA are also expected to increase the potential

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**Figure 3.** Comparison of IIEF-5 (baseline and of study visit) and PDQ questionnaire results (before and after each visit). The results showed significant difference in mean PDQ questionnaire in PDQ1 with p= 0.002 (CI 0.720;2.968), PDQ2 with p= 0.021 (CI 0.275;3.225) and PDQ3 with p= 0.001 (CI 1.520;3.220).

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efficacy of stem cells, as suggested by in vitro and animal studies [13,23]. PRP rich in growth factors and other compounds initiates a reconstructive process in TA, whose global effects are visible through echographic controls. Although widely used for diagnostic purpose [24,25], US technique is very specific to this proposed technique of treatment with two advantages. First, it allows taking precise measurements of the whole albuginea, providing comparison data all along the procedure. From this, we have learned that the diseased TA extends largely beyond what is manually palpated, and also that the reduction of the thickening precedes the decrease of angulation, while both are correlated (p= 0.015 after 4 sessions and p=0.011 at the end of each patient’s treatment). Second, injections under US control are much more precise and safer. They also allow applying a mechanical action on the heavily fibrotic and/or calcified areas. Needles are used to dig channels into the fibrotic layers before injecting PRP and HA. Imaging avoids any major complications such as tunica ruptures or intracavernous injection of the product (Figure B in appendix 4), which have been reported with Xiapex [26]. Complications were limited to transient ecchymosis even in patients under anticoagulant treatment. Based on the use of autologous blood extemporaneously processed, the combination of PRP and HA is an office treatment complying with all regulations concerning fresh blood manipulations. It is economically viable, being much cheaper than Xiapex and Interferon alpha. Patients’ satisfaction, whatever measurements are used to quote it, was very high. Interestingly, patients themselves, even if the results were not tangible, asked to continue the treatment.

If confirmed in larger controlled studies, the use of a combination of PRP and HA obtained with CellularMatrix BCT-HA and Regen BCT devices for PD will stand as the first autologous blood-based treatment used for a sexual dysfunction. We are currently planning a similar study aiming at assessing the improvement of ED, based on the fact that 37% of the patients of the present study claimed better erections, independently of the improvement of PD symptoms.

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Appendix 1

US study of PD

All measurements were carried out by the same person (HS). Thickness of the TA was measured automatically in the dorsum, laterally and ventrally, not only where the plaques were felt but also at three different levels of the penile shaft: proximal, median and distal. The probe was applied transversally for the dorsal and ventral part of the TA and longitudinally for its lateral part. A previous investigation (yet unpublished) conducted on 50 patients naive of PD provided normative data for the TA (Figure A) useful for the comparison with data recorded in the present study. For better comprehension and statistics, for each patient the maximum thickness was elected in the study. The intralesional treatment was systematically injected under US guidance.

Figure A. Comparison of the average tunica albuginea thickness (in millimeters) between the control group and PD patients

Appendix 2

A BCT-HA tube (CellularMatrix BCT–HA Kit, Regen Lab SA, Switzerland) and a Regen BCT tube (RegenKit-BCT, Regen Lab SA, Switzerland) were used to prepare a combination of PRP/HA and PRP, respectively. Both tubes are evacuated tubes that contain an anticoagulant solution and an inert thixotropic separating gel that forms a physical barrier between blood components after centrifugation. Additionally, the BCT-HA tube contains 2 ml of a 2% hyaluronic acid solution. It is a non-crosslinked hyaluronic acid (1550 kDa) obtained by bacterial fermentation. Once prepared according to the Instructions for Use provided with the kits, both products were collected using a transfer device ensuring a close-circuit preparation procedure and mixed together in a 10 ml-syringe. As the Regen Lab’s technology allows the preparation of PRP with a platelet concentration 1.6-fold above the value in venous blood, platelet concentration in PRP is expected to be between 300'000 and 350'000 platelets/μl.

Appendix 3

Additional questionnaire

5 different items were asked after 4 sessions and after the treatment has been completed:

1 - Do you feel cured (5 points) improved (3 points) or unchanged (1) ?
2- Has the deformation disappeared (5) diminished (3) or unchanged (1)
3- Is sexual intercourse easier (5) identical (3) or more difficult ( -1)
4- In the overall, do you find the treatment positive (5) or without result (-1)

To obtain the global score mentioned in the text, sum up the scores of the four items

A fifth item was asked in the same questionnaire concerning erections: are they improved, unchanged or worse?

Appendix 4

Intracavernous accidental injection of Collagenase clostridium histolyticum in a 32Y old patient with Peyronie’s disease. Echography one month after intralesional injection (Figure B).
Figure B. 1) Site of CCH injection. 2) Channel inside TA. 3) Accidental Intracavernous injection of CCH