Amgar Gilbert and Pierre Bouhanna discuss the use of microneedling along with platelet rich plasma for the treatment of androgenic alopecia

ABSTRACT

The use of Platelet Rich Plasma (PRP) in cosmetic medicine is now well established and documented. Hair applications became available over the last 8 years. The two main fields of application for PRP are hair regrowth (androgenic alopecia, alopecia areata) and surgery (to improve hair transplants). Indeed, some of the treatments that exist in this field provide quantifiable results. In this study, the authors conducted an objective evaluation of androgenic alopecia for over 2 years. The study consisted of

70 patients and each case is documented by digital parameters (hair density, diameter, ratio terminal vellus) and standard photography. Follow-up 3 months after treatment, the authors observed an increase in hair density of 19%, which reached 38% at 12 months. The authors believe it would be suitable to consolidate these results with other evaluations of the same type, carried out by other teams and gradually define the best protocols in order to maximise the results.

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KEYWORDS

PRP Platelets Rich Plasma. androgenic alopecia, phototrichogram, Tropocells® kit

NDROGENIC ALOPECIA extremely common disease that affects both men and women. In normal hair loss, less than one hundred hairs fall each day and are replaced by new thick hair. In the evolution of male and female pattern baldness, the new hair is fine and thin (intermediate hair or miniaturised hair). Male and

female baldness usually progresses in a defined pattern.

The most publicised medical treatment available for male pattern baldness is 5% minoxidil lotion and 2% minoxidil for female baldness. Side-effects of minoxidil are minimal but include itching, eczema, and hypertrichosis in females. For male baldness, finasteride taken orally and daily (1 mg) inhibits the 5α reductase from forming dihydrotestosterone (DHT) and may lead to decreased libido. Spironolactone (in US) appears to be a competitive inhibitor of DHT-receptor

Cyproterone acetate (in Europe) can effectively block increased levels of male hormones that cause hair loss in some women

A new interest in preventing baldness has been stimulated by cellular therapy with platelet rich plasma (PRP) injected directly into the scalp.

Aiming for better tolerance, an easier process, and high satisfaction rates, researchers and doctors have been looking for improved treatments in this field for a number of years. In this context, PRP takes a prominent place. PRP is defined as an autologous concentrated preparation of platelets and their associated growth factors in a small volume of plasma. The revitalisation qualities of the platelets are well known and have been used for 30 years in various specialties such as plastic surgery, dermatology, rheumatology, and dentistry. The first publication on this subject goes back to 19751 and publications of its use on the scalp date back to 2007. PRP eliminates concerns ▷

66 A new interest in preventing baldness has been stimulated by cellular therapy with platelet rich plasma (PRP) injected directly into the scalp.





Figure 1 (left)

Figure 2 (right)



Figure 3



Figure 4

for disease transmission and minimises possible side-effects.

The challenge remains to evaluate the results with objective parameters².

Epidemiology

The prevalence and severity of pattern hair loss increases with age and the greatest incidence is in caucasian people. In general, 30% of caucasian males are affected by the age of 30, 50% by age 50, and 80% by age 70. In women, 13% of premenopausal women have some evidence of pattern hair loss. However, the incidence increases in women around the time of the menopause and may affect 70% of women over the age of 65 years³.

Physiological consideration

The process begins by performing a blood sample on an anti-coagulated tube with separating gel. The tubes are then centrifuged for a few minutes (speed and time depend on the kit protocol). Owing to their density the red cells fall to the bottom of the tube. On the upper-part of the tube the plasma and the platelets can be found.

Once in the dermal layer, platelets are activated, they inflate and the growth factors are released. Platelet activation is a natural reaction when a traumatic process occurs in tissue (injection or injure). Activation can also be provided by adjunction of CaCl2 or thrombin. In this series, the authors' did not use any activator. The most important growth factors for hair application are:

- Platelet derived growth factor (PDGF) stimulates the growth of dermal mesenchyme⁴
- Vascular endothelial growth factor (VEGF) creates neoangiogenesis, as well as increases and improves hair growth and hair size⁵⁻⁸
- Epidermal growth factor (EGF) stimulates mitosis of epithelial cells and fibroblast, and improves the rate of anagen, the active growth phase of hair follicles 9.10
- Insulin like growth factor (IGF) slows down apoptosis¹¹
- Fibroblast growth factor (FGF) stimulates the proliferation and differentiation of keratinocytes and endothelial cells¹²
- Nerve growth factor (NGF) strongly stimulates hair growth and slows down apoptosis. NGF has a modulating effect on the hair depending on the receptor with which it interacts. NGF also acts as stress-mediators. This is certainly a good way to explain the correlation between stress and hair loss¹³.

Previous studies

There are three applications in the hair-loss field: androgenic alopecia, alopecia areata, and hair graft. Regarding PRP evaluation in androgenic alopecia, there are five published studies.

The first study from Sorbellini et all exhibited an in vitro study conducted on 50 patients. Twelve follicles were taken from each patient, four follicles were induced with PRP, four in Ringer's solution, and four in standard solution. The mitotic activity was then measured. According to the results, the PRP group showed a significant increase of mitotic activity and a reduction of apoptosis.

The purpose of the next study by Takikawa et al¹⁵ was to determine whether there was a difference between simple PRP injections and PRP containing deltaparine and protamine microparticles, which acted as carriers for growth factors in the PRP. The study was conducted on 26 volunteers with thin hair who received five local treatments. PRP improved hair density in 16% at 12 weeks. There was no significant difference on the hair density improvement with or without deltaparine.

In the third study, Greco and Brandt¹⁶ demonstrated that infusing growth factors into the scalp did reverse miniaturisation over an 8 month period when compared to control. They observed an increase of 9.7% in average hair shaft diameter at 4 months and then 6.1% at 8 months in the treatment group.

In the fourth study by Sciavone et al, researchers explored the possible clinical benefit of injecting platelet-derived growth factors into the scalp of patients using a specific autologous blood concentrate. Two injections of a leukocyte platelet-rich plasma (L-PRP) with the addition of concentrated plasmatic proteins were administered at baseline and after 3 months in 64 patients. Macrophotographs were taken at baseline and after 6 months, and two independent evaluators rated them using the Jaeschke rating of clinical change. Some improvement was seen in all patients by one evaluator and in 62 by the other $^{\rm pr}$.

In the fifth study, Li et al showed that injection of mice with activated PRP induced faster telogen to anagen transition than in control mice¹⁸.

Objective of the evaluation

This article intends to retrospectively evaluate a cohort of 70 patients with progressive androgenic alopecia. These patients underwent a combination therapy with injection of platelet extracts prepared using a PRP kit, and a rotary single-use multipuncture instrument (dermaroller).

Materials and methods

PRP was prepared using the PRP kit (*Figure 1*) through a one step procedure. A total of 30ml of blood was transferred to PRP tubes containing acid-citrate dextrose modified solution (additive/blood ratio of 1:567) and separating gel (1:5). Two tubes have been used for each procedure (60cc). The PRP tube was then centrifuged (1500g for 10 minutes at room temperature) resulting in blood separation into three layers: an erythrocyte and granulocytes layer at the bottom of the tube beneath the gel, a PRP layer in the middle, and a platelet-poor plasma (PPP) layer at the top of the tube. The PPP layer was drawn out (13±1 ml) and the remaining plasma (5±1 ml) was gently mixed onto the gel surface where 85%±5% of the platelets are present, creating the PRP.

The remaining materials used in this study include:

- A dedicated centrifuge
- A 1.5mm dermaroller, (*Figure 2*) designed to make micropunctures on the surface of the scalp
- Digital phototrichogram (Figure 3) with capillary measurement software developed by the authors' team
- Photographs (Figure 4) with six incidences: front, 45°, 90°, 135°, 180°, and side.

Inclusion and exclusion criteria

The inclusion criteria consisted of:

- Men and women
- 20 to 80 years old
- Androgenetic alopecia stages II-VI of the Norwood-Hamilton scale, or 1-3 from the Ludwig scale. The exclusion criteria consisted of:
- History of cancer
- lacksquare Use of anticoagulants or blood products
- Scalp active skin infection
- Haematological diseases

■ Pregnancy.

Methodology

Seventy patients (31 males and 39 females) were recruited for the study. Digital measurements (tattoo mark, hair density, diameter, and ratio) and standard photographs were taken from each participant. In all, 60 cc of blood was drawn and 10 cc of PRP was processed. Following local anaesthesia, the scalp was first traumatised with a 1.5 mm microneedling roller and then injected with the PRP. Patients underwent two sessions 3 months apart. No external activation has been provided.

Schedule

Before the procedure could take place, a prior consultation with haematologic analysis and informed consent was required.

First treatment on day 0

- Signature of informed consent
- Photographs taken from six different angles
- Digital phototrichogram for measurement of capillary density and the diameter of the hair in an area identified by two micro-tattoos
- Local anaesthesia, application of the dermaroller and injection of PRP.

Second treatment on day 90

- Photographs taken again from the same six angles
- Digital phototrichogram
- A search for side-effects
- Local anaesthesia, application of the dermaroller, and injection of PRP.

Examination on day 180, 270, and 360

- Photographs taken again from the same six angles
- Digital phototrichogram
- A search for side-effects.

Treatment with PRP and dermaroller

First, a local anaesthesia injection of 1% lidocaine must be given prior to use of the dermaroller or intradermal injections of PRP.

In order to allow better diffusion of the PRP, multiple perforations were conducted on scalp areas affected by the process of androgenic alopecia using a single-use dermaroller instrument with peaks of 1.5 mm.

Next, treatment with PRP involves the injection of 10 cc of PRP obtained by centrifugation of the PRP kit with the centrifuge pre-programmed for this purpose. The injections will cover the area of the affected scalp. The preparation of the PRP was made in accordance with the operating instructions of the kit manufacturer.

Review of side effects

Common side-effects associated with PRP injections and dermaroller include the following:

- Pain
- Redness
- Pruritis

- Oedema
- Haematoma
- Other.

Descriptive statistics for the samples:

A. Normality test

1. Age

Seventy patients, 31 men and 39 women, with ages of 17-77 years old were enrolled in this evaluation.

	I N	Sean	ECTYP	Ch1:	CNI, IIW	PYCCA	Normality
Age	70	44,7	14,31	9,443	11,07	<0,093	Normality

2. Density

1	N	Mean.	EcType	Ch1'	Chi* lim	Proba	Normality
	Density 70	151,3	47,25	3,494	7,815	<0,32	Acceptant

3. Diameter

							Normality
Diameter	70	40,29	11,17	6,200	11,07	<0.28	Accept

The above analysis demonstrated that the size of the groups chosen is acceptable and can statistically represent the population, as tested by chi² test (StatEl). Chi² refers to Pearson's chi-squared test and StaEl calculates the probability and confirms that the relevant variables are normally distributed.

Table 1

Results at 3 month Density

At day 90, the authors observed an improvement of 19.7% in hair density, which they believed to be a very interesting result coming 3 months after the first session. The patients reported no more hair loss after 2 or 3 weeks after their first session. However, an initial acceleration of the fall due to the telogen expulsion was observed in some patients.

The repartition of the improvement was found to be very close between terminal and vellus hair (terminal +30.8% and vellus +28.3%). Hairs are considered as vellus if the diameter is <40 microns.

The percentage of improvement is defined as: (density at 3 months minus initial density) divided by initial density. It is not the number of new hairs.

Regarding sex, (*Figure 5*) the authors observed a small advantage for the men in density variation (21.3% against 18.5% for the women).

Regarding age, better results on density change were witnessed in patients over the age of 60 (30.6% against 17% for those under 60 years of age) (Figure 6).

Regarding stage, results were not as responsive for men in stages II and VI of the Norwood-Hamilton scale (*Figure 7*). Statistically, a clear correlation cannot be established between the results and the stages of hair loss for men. On the other hand, the female group showed higher values on stage 3 (25.3%) vs stages 1 and 2 (17%).

Diameter

At day 90, the mean diameter does not change (+0.3%). Regarding sex, the authors observe a little advantage for the women (+1.96% against -1.82% for the men).

Regarding age, results are not so positive on diameter after 60 years (-4%). It seems that there is a correlation

between hair density and the growth of new thin hair, so the mean diameter decreases.

Regarding stage, the authors could not establish a correlation regarding the stage for men. For women, they clearly observe a correlation on stage (stage 1:-5.31%, stage 2:+2.05%, and stage 3:+10.05%).

Telogen/vellus ratio

The results concerning the ratio are comparable with the diameter evolution. $\;$

Over the 90 day period, the authors observed many differences as illustrated by the digital hair image in Figure 8. As can be observed in Figure 8, the two images were taken exactly in the same place. The highlighted area shows the variation in hair density, which is changing from 3 to 5, 0 to 1, 2 to 3, and 2 to 3. This is a very important result that confirms clearly the effect of PRP on density.

The results can be seen in the macrophotographs before and 3 months after one PRP session (Figure 9).

Results at 12 month

Following 12 months from the first treatment, only 46 patients (out of the intial 70) were still involved in the study due to several causes:

- Distance (7): The PRP procedure is not often used, so recruitment is not local but rather national or international. This explains the monitoring difficulties
- Pain (2): This is not a painless procedure and pain evaluation is between 2 and 5 on a 10 point scale
- Loss of tattoo (1): Despite a standardised technique, sometimes the skin does not keep the pigment.
- Not enough of a result and the patient did not return
 (4)
- Unknown (10).

Density

At 12 months, after two sessions of PRP, an improvement of 38% in density variation (*Figure 10*) was observed. This is a very important evolution. For most of the patients, the improvement was observed at about 6-8 months.

Another way to evaluate the evolution is to observe the density in absolute value (*Figure 11*). The initial mean was 128 hairs/cm² and it increased to 161/cm² at 12 months.

However, the more accurate way is to evaluate the result is by using base 100 (*Figure 12*). This formula is often used in financial statistics to evaluate the yield of an investment. Values are converted as if every patient had an initial density at 100 hairs/cm². In this study, we obtained the following values: 100 - 115 - 122 - 121 - 123.

Looking at *Figure 12*, the yield is 23% and it seems to be a good reflection of the clinical reality. Looking at the curve, it is not linear as you would expect with the density variation. The curve has a logarithmic aspect and has a progression stage (6 months) and then a stabilisation stage. This is the translation of the physiological function. The procedure does not create new follicles. If it was so, a linear curve should be evident. Instead, the procedure stimulates a lot of follicles that are not strong enough to

provide hair but are not yet dead. 'New' hair grows but not new follicles. Usually, one session every year seems to be enough to keep the benefit. In a few cases, when a decrease is observed at 9 months, the authors provide two sessions every year.

The result can be seen on the macrophotographs before and 12 months after two PRP sessions (*Figures 13-14*).

Regarding sex, at 3 months women witnessed a slight delay. At 12 months, they recovered a clear advantage with 55% improvement (only 40% for the men) (Figure 15).

Regarding age, better results on density after 60 years. (126%, against 36% under 60 years of age) (Figure 16).

Again regarding stage, the authors could not establish a correlation regarding the stage for men. For women, the authors clearly observed a correlation on stage 1: 16.55%, stage 2: 28.23%, and stage 3: 233.63% (Figure 17).

Diameter

Globally, no significant changes were observed. However, focusing on women in stage 3, the authors observed a very positive evolution stage 1: -14%, stage 2: -1%, and stage 3: +36%

The authors looked to see if there was a correlation between improvement and initial value. No statistically valid correlation with linear regression for either men or women was found.

Side-effects

Pain is always present (from 2 to 5 on a 10 point scale). It is not a painless procedure and it could be useful to use analgesic gas. For one patient a preauricular lymph node occurred on the second day and disappeared spontaneously 2 days later.

Limit of this evaluation

It is impossible to take measurements all over the scalp. Therefore, the authors had to choose an evaluative area to see the changes brought about by the procedure. Nevertheless, the results can be different between areas. For example, an 18% improvement was observed on the border line for a man in stage 5, while no changes were detected on the vertex area.

Using adhesive also posed an issue. Before taking the digital phototrichogram, the authors cut the hair very short and used an adhesive tissue to take away hair fragments in order to get a clear picture. The hypothesis cannot be excluded that some telogen hair was present in the area and was subsequently removed.

The authors' protocol used two therapeutic parameters: PRP and microneedling. Even if clinical experience shows that the most important effect is due to the PRP, this evaluation is unable to prove it. The goal was to get the best result for the patient and the authors are evaluating other studies with different protocols (without microneedling, with activation, and with less quantity) and wll soon be able to quantify the result and improve the protocol.

The number of patients who dropped out of the study before the 12 month follow-up was high but this is not a

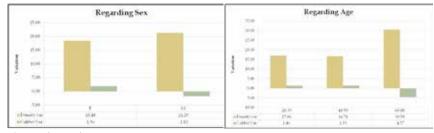
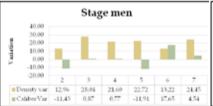


Figure 5 (Top left)
Figure 6 (Top right)



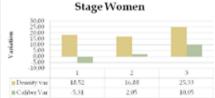


Figure 7



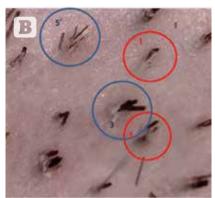


Figure 8 (A) digital image taken on day 1, and (B) at the same place on day 90 showing the change in hair density





Figure 9 (A) Before, and (B) 3 months after one PRP session

prospective study, this is a retrospective evaluation.

Conclusion

In conclusion, the authors evaluated 70 cases of androgenic alopecia over a period of 3 months, and 46 cases over 12 months post-treatment with PRP. They observed an improvement of 19.7% for the density

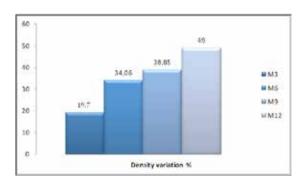


Figure 10

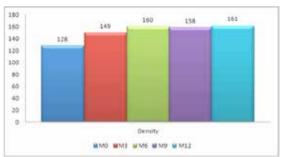


Figure 11 mean density

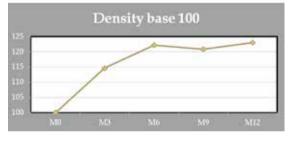


Figure 12



Figure 13 (A) Before and (B) 12 months after two PRP sessions



Figure 14 (A) Before and (B) 12 months after two PRP sessions

variation at 3 months that reached 38% at 12 months. The aspect of the curve suggests that new follicles were not being produced, and the authors believe that PRP treatment stimulates many live follicles that are not strong enough to provide hair. Therefore, 'new' hair is observed, but not new follicles.

Globally, concerning the diameter, no significant changes, except for advancing age (60-80 years) and women in stage 3, were observed.

It would be useful to consolidate these results with other evaluations of the same type, provided by other teams and gradually define the best protocols for the best results.

▶ Declaration of interest None

► Figures 1-17 ©

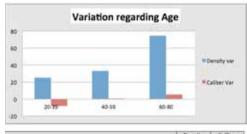
Figure 15

Variation regarding Sex # Density var 20 10 0 W Caliber Var

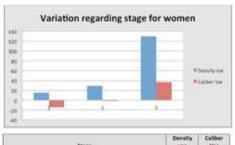
Sex	Density	Caliber Var
į.	44,53	3,43
M	31,37	-7,57
Mean	37,95	-2,07

Figure 16

Figure 17



	Age	Density	Caliber
20-39	11,123	25,05	-8,19
40-59		32,91	0,17
60 80		74,39	5,42



Stage	Density var	Caliber
1	15,35	-14.49
2	28,78	41,43
1	129,39	36,30
Mean	44,53	3,43

O Key points

- This article intended to retrospectively evaluate a cohort of 70 patients with progressive androgenic alopecia.
- These patients underwent a combination therapy with injection of platelet extracts (without any activation) and a rotary single-use multipuncture instrument. Digital measurements (tattoo mark, hair density, calibre, and ratio) and standard photographs were taken from each participant.
- An improvement of 19.7% for the density variation at 3 months, reaching 49% at 12 months was observed
- Occerning the caliber, no significant changes, except for advancing age (60-80 years) and women in stage 3 were observed

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