Autologous Platelet-Released Growth Factor and Sexual Dysfunction Amendment: A Pilot Clinical Trial of Successful Improvement Sexual Dysfunction after Pelvic Irradiation

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Abstract

Sexual dysfunction (SDF) is a common sequel to cancer treatment which affects the quality of life in women treated with pelvic radiotherapy. The aim of this study was to evaluate the safety, symptom resolution and objective improvement the injection of autologous platelet released growth factor (APRGF) for treatment of SDF in cited patients. This prospective pilot study enrolled 10 cancer-free patients with SDF who underwent pelvic radiotherapy at least 5 years ago, randomly. Each patient was received 1-2 cc APRFG within four weeks and all patients were re-evaluated at eight weeks and six months. CD34 immuno histochemistry and Masson’s trichrome staining were performed on vaginal biopsy section for angiogenesis and fibrosis assay respectively. Sexual satisfaction after the injection of APRFG was clinically difference and the entire patient had sexual satisfaction. In the patient’s follow-up, none of them needs to repeat the treatment. Our results declared that APRFG injection was effective and symptoms were disappeared in the entire patients. Significant objective improvements in vaginal diameter (mean before injection, 6.5 cm vs 7.1 cm after injection) (p-value = 0.001) and vaginal flexibility (mean before treatment, 0.72 cm vs 1.85 cm after injection) (P-value = 0.026) were observed. Characteristics of discharge before the injection in 60% of patients were included dry vagina and 40% had mild discharge but after injection 40% of patients had moderate and also 60% had mild and sufficient discharge (P-value= 0.190). Overally, our patients reported better sexual function and showed better vaginal function indexes, after APRFG injection.

Keywords: Autologous-platelet-growth factor- sexual dysfunction- pelvic irradiation

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Introduction

Sexual dysfunction is a common sequel to cancer treatment, affecting the quality of life (QOL) in women treated with pelvic radiotherapy (Jensen et al., 2003; Andreyev, 2007; Dunberger et al., 2010; Incrocci and Jensen, 2013). Pelvic radiation may be delivered as the external beam and/or brachytherapy. The accumulated radiation dose to the pelvic organs is important for acute bowel, bladder, and genital toxicity (Jensen and Froeding, 2015). The occurrence of complications is dose and fractionation dependent. Radiation effects are progressive and may become symptomatic after a latent period, but there may be a continuous progression from acute edema, mucosal and sub-mucosal inflammation and persistent ulceration and necrosis to fibrosis. On the longer term, vaginal wall thinning, adhesions atrophy, and fibrosis may occur often followed by decreased vaginal elasticity, narrowing, shortening and ultimately total vaginal stenosis (Schover et al., 1989; Bergmark et al., 1999). A healthy sexual response is described having four phases; desire, excitement, orgasm, and resolution, whereas female sexual dysfunction (FSD), includes desire, arousal, orgasmic and sexual pain disorders (Basson et al., 2000). Dyspareunia (pain during sex) is the most usual ailment in these patients. It’s often linked to alterations in the vaginal tissues (stenosis, vaginal fibrosis or atrophy), vaginal size (total vaginal length, genital hiatus) or vaginal dryness which results from in loss of adequate lubrication during intercourse. These changes manifest as superficial or deep dyspareunia. Current studies have evaluated various treatment choices for female dyspareunia and have
recommended hormone replacement therapy, the use of vaginal dilators and also surgery.

The cited treatment strategies have not been successful, because of the concern around the risk of fistula after surgery and cancer recurrence following estrogen intake (Phillips, 2000; Kingsberg et al., 2010). Therapeutic alternatives for solving these problems are very special because of the median age of adult females who became radiation-induced cancer was so high and the tone of sex life was not a priority for patients (Carter et al., 2017). During the past decade by changing the shape of cancer outbreaks in younger women and progress in developing new radiation methods for cancer treatment, life expectancy has increased and quality of sexual relations has been spotlighted. Over the last decades, the role of platelet-derived growth factors has been highlighted in regenerative medicine (Mostafaei et al.). Platelet-derived growth factors are biologic active compounds and also acting in different mechanisms and pathways including activation or induction of chemotaxis, cellular proliferation and angiogenesis (Mohammadi et al., 2016; Mohammadi et al., 2017b). Platelet related products have been a satisfactory outcome in the treatment of chronic skin and soft tissue lesions and plastic surgeries (Serra-Mestre et al., 2014; Mohammadi et al., 2017a). In new strategies for treatment of FSD, following the cited facets should be considered (Farmer et al., 2016; Carter et al., 2017)

- Do not contain hormonal combinations
- Able to reduce fibrosis and enhances vaginal flexibility
- Can stimulate vaginal secretion and natural lubrication by angiogenesis
- Had a good price

We focused for the first time to identify, describe, and evaluate the safety, symptom resolution and objective improvement the injection of APRGF for treatment of FSD in patients after pelvic irradiation.

Materials and Methods

Patients and methods

Demographics data

Ten patients with a history of the dyspareunia after radiation were recruited, in the Tehran University of Medical Sciences, Shariati hospital from July 2017 to December 2017. The age of these patients ranged from 30 years to 56 years. Patients were selected from women who had a history of radiotherapy due to cancer (bladder, uterine, cervix or rectum) with vaginal problems. A summary of patients’ characteristics was presented in Table-1.

Patient selection

This prospective pilot, single arm clinical trial enrolled 10 cancer patients with sexual dysfunction who underwent pelvic radiotherapy at least 5 years ago, randomly. All patients were cancer-free according to their clinical and para-clinical evaluation. An exclusion criteria were defined as follows: lower platelet count of <10*10^12 µL, hemoglobin <10g/dL, anti-proliferative therapies such as chemotherapy in the past 3 months, anti-inflammation medications in the past 2 weeks, radiation in the past 3 months, growth factor therapies in the last 2 weeks, history of diabetes, history of mal-absorption (celiac disease) and any contra-indication in peripheral venues access (Mohammadi et al., 2017b). All the patients were asked to sign a written consent form freely after explanation of all the aspects of study according to their level of knowledge. The cases were referred to the Medical Ethics Committee in order to avoid any conflict of interest. The current study was approved by Tehran University Medical Ethics Committee (IR.TUMS.VCR.REC.1397-4915) and with the Helsinki Declaration of 1975, revised in 1983.

Patients’ history and medical data

Pap smear, abdominal and pelvic CT scan and complete examination are done for all of the patients. Vaginal examination was performed during the screening on those who fulfilled the inclusion and exclusion criteria. The vaginal exam includes a POP-Q, speculum and manual, to evaluate the labia, genital skin, vaginal mucosa, cervix, uterine size (if not surgically removed); mass perception or touch, vaginal flexibility, stenosis, and atrophy was performed. The vaginal flexibility was measured by speculum examination and width of opening blades.

In two patients, we were not capable to place a speculum because of severe stenosis and zero flexibility reported. In three patients, we could just go into the speculum and there was no possibility of opening the blades, we covered the flexibility 0.5 in these patients. Another item that was evaluated at the beginning of the study and follow-up visits was the TVL, which was measured during the POP-Q examination with a graded swab; because shortness of TVL due to radiotherapy fibrosis causes deep dyspareunia. Patient report of vaginal discharge was one of the important points that asked about its quality and quantity in every visit. A natural vaginal discharge is a clear, odorless liquid that facilitates penetration and reduces dyspareunia, by creating lubrication. Its origin of transudate vaginal capillaries is a sign of vaginal refreshment and youth. Vaginal dryness is one of the significant causes of dyspareunia in patients due to lack of hormonal support and fibrosis. This component was evaluated qualitatively through a patient’s history and during the examination. It was reported to be severe dryness (never), mild to moderate and satisfactory (adequate) discharge. Intercourse frequency and patient report of sexual satisfaction were evaluated through biographies and special questionnaires at each visit, and the score was recorded. All details were documented and assessed at the end of the survey. Finally, in patients who were satisfied with the sampling, one biopsy of the lateral vaginal wall (0.5 * 0.5 cm) was taken to compare before and after treatment.

Study protocol and PRP preparation

Each patient received 1-2 cc PRFG within four weeks (4 times injection) period. For APRGF preparation, first 15 ml peripheral blood was drawn from a donor using a 20 ml syringe containing 1.5 ml anticoagulant: sodium citrate
(ACD-A1) (Terumo BCT, Inc., USA). Then, the blood was shaken gently 5 times. Afterward, it was transferred into two 10 mL tubes (Standard kit) (Noavaran Salamat Arzhang, Inc., Iran) and centrifuged at 150×g for 10 min at 24°C. After first centrifugation, the 2 fold rich platelet in the supernatant plasma was achieved. This platelet-rich plasma (PRP) was then transferred to the second tube for 340g (5min) centrifugation. APRGF was obtained from concentrated PRP by freeze and thaw method. Obtained products were centrifuged three times at 900 g for 30 min to remove platelet bodies (Mohammadi et al., 2016). Finally autologous APRGF injected sub-dermally and/or intra-muscularly, infiltrating the areas involved of the vaginal walls (eg. Anterior vaginal wall into a space between the vagina and urethra, into the clitoris and lateral vaginal walls), Then the patients were compared for changes in color and appearance of the vaginal tissue and vulva, vaginal discharge and dryness, vaginal flexibility, trans vaginal length, intercourse frequency and sexual satisfaction to determine the need to repeat the treatment.

Lateral wall vaginal biopsy

According to the signed consent forms taken from the patients, a lateral wall vaginal biopsy must have been done before starting APRGF injection and after final injection in order to evaluate the histological differences. The tissue was evaluated for neovascularization using immunohistochemistry (IHC) assay of CD34+ (BD PharmingenTM, purified rat anti-CD34) positive cells in order to evaluate the angiogenesis. Additionally, Masson’s trichrome staining was ordered to investigate the fibrosis before and after treatment.

Treatment assessment

All ten patients were re-evaluated at eight weeks and six months later to rule out vulvar carcinoma, relapse of the symptom and the need for re-injection.

Statistical Methods

All data were analyzed using Statistical Package for Social Sciences, SPSS, version 18.0 (SPSS, Inc., Chicago, IL, USA). Continuous variables are presented as mean± standard deviation and categorical variables are presented as N (%). For comparing the central tendency (e.g. mean for normal and median for non-normal gene expression) between before and after intervention, paired t-test or Wilcoxon signed rank non-parametric test was performed. Any P-value less than 0.05 considered as statistically significant.

Results

Patients’ data

In this study data were obtained from ten Iranian women aged 30 to 56 years old (mean= 46 years old), regarding parity (nulliparous 30%, multiparous 70%), sexual activity and frequency of sexual relations (50 % inactive , 50% Rare), type of cancer (rectum 20%, cervix 40%, bladder 20% and endometrial cancer 20%), radiation type (brachtherapy 40%, external radiotherapy 60%), history of hormone replacement therapy (40% of

<table>
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<th>Table 1. Summarizes Patients’ Characteristics</th>
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<tr>
<td><strong>Age (year)</strong></td>
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<td><strong>Cancer Type % (Number)</strong></td>
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<tr>
<td>Rectum cancer</td>
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<td>Uterine cancer</td>
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<td>Bladder cancer</td>
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<td>Cervical cancer</td>
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<td>Radiation Type % (Number)</td>
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<tr>
<td>External radiotherapy</td>
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<td>Brachtherapy</td>
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<tr>
<td>Sexual activity % (Number)</td>
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<tr>
<td>Nulliparous</td>
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<tr>
<td>Multiparous</td>
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<tr>
<td>Sexual frequency % (Number)</td>
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<tr>
<td>Inactive*</td>
</tr>
<tr>
<td>Rare**</td>
</tr>
<tr>
<td>History of HRT % (Number)</td>
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<tr>
<td>Yes</td>
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Activity and frequency of sexual relations: *Inactive: Not having sexual intercourse in the last year, **Rare: Less than 4 times a year

APRGF reduces fibrosis and improvement vaginal flexibility, through increasing the diameter of genital hiatus and vaginal length.

Data showed that symptoms had disappeared in all of the patients with significant objective improvements in vaginal diameter (mean of TVL before PRFG injection was 6.5 cm vs 7.1 cm after PRFG injection), p-value = 0.001. Vaginal flexibility (mean of flexibility before injection was 0.72 cm vs 1.85 cm after injection), P-value = 0.026, showed significant improvement after APRGF injection. Figure 1 and the summarized results in Table 2 were declared that PRGF reduces fibrosis and improvement vaginal flexibility, through increasing the diameter of genital hiatus and vaginal length.

Intercourse frequency and sexual satisfaction improvement after APRGF injection is related to vaginal discharge

Form discharge points of view, 60% of patients had a dry vagina and 40% had mild discharge that did not provide the effective lubricant during sexual intercourse. After APRGF injection, 40% of patients had mild to moderate and 60% had adequate discharge (P-value= 0.190). The color and appearance of the vagina were significantly different before and after treatment. Patient’s intercourse frequency after APRGF injection increased and sexual satisfaction improved after treatment. Before study, patients were disagreeing while after APRGF injection 60% strongly agreed and the entire patients had sexual

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satisfaction which was related to TVL, vaginal flexibility and discharge. During the follow-up visit, any patient needs to repeat the treatment. Patient report of vaginal discharge and sexual satisfaction before and after treatment with PRGF were summarized in Table 3.

**Histopathological assessment**

Comparison of vaginal wall biopsy showed improvement of atrophy, fibrosis, and angiogenesis after PRGF injection (Figure 1). The reticulin staining demonstrated a significant decrease in fibrosis after PRGF injection and also showed subepithelial loose fibroconnective tissue without fibrosis (Figure 2A). The micro-vessels formation (angiogenesis) was evaluated in all the patients through a lateral vaginal biopsy. There was no significant difference before and after APRGF injection in IHC staining for CD34 in this biopsy section after four weeks (4 doses injection) (Figure 2B).

**Discussion**

Sexual dysfunction is a common sequel to cancer treatment, in women treated with pelvic radiotherapy. This complication increased over the time and was greater for cancer patients who were female, older, married or in a committed relationship (Acquati et al., 2017). The cohort

| Table 3. Patient Report of Vaginal Discharge and Sexual Satisfaction before and after Injection of APRGF |
|---|---|---|---|
| Intercourse frequency | Patient report of vaginal discharge | Sexual satisfaction |
| Before treatment |  |  |  |
| Inactive | Dry | 50% | Disagree | 50% |
| Rare | 50% | Mild | Strongly agree | 0% |
| Active | 0% | Adequate | Agree a little | 40% |
| After treatment |  |  |  |
| Inactive | 0% | Dry | Agree a little | 10% |
| Rare | 30% | Mild | Strongly agree | 40% |
| Active | 70% | Adequate | Disagree | 50% |
Autologous Platelet-Released Growth Factor and Sexual Dysfunction Improvement

A profile was designed for the first time by the Women’s Health Clinic (WHC) at Mayo Clinic in Rochester, Minnesota, medical information. They were evaluated 8688 women from 2005 to 2017, to determine the prevalence of the sexual dysfunction and its relationship with health, disease, age and so on. The study showed a high need for curative and preventive sexological care to improve quality of life of women with infertility (Faubion et al., 2018). Available therapeutic options such as vaginoplasty and hormone replacement therapy are not suitable. The risk of fistula and secondary reconstruction vaginal stenosis following surgery and a relapse of the tumor due to hormonal administration are the common concern of these therapeutic strategies (Kingsberg et al., 2010; Carter et al., 2017). Owing to the many probable etiologic factors involved in FSD and the inconsistency in the response to current treatment modalities, this part needs research to introduce effective therapeutic options (Buster, 2013; Milbury et al., 2013).

In new strategies for treatment of FSD, ability in fibrosis reduction enhances vaginal flexibility and stimulation of vaginal secretion via angiogenesis induction should be considered (Farmer et al., 2016). Based on the ability of PRP in the induction of angiogenesis and tissue regeneration, it seems to be a suitable choice in this area (Mostafaei et al.; Mohammadi et al., 2017a; Mohammadi et al., 2017b). Thus, the effort to use low-complicated treatments that could rejuvenate the patient to pre-radiotherapy conditions led us to use APRGF. The result of the present study for the first time reported better sexual and vaginal function indexes after APRGF injection in patients undergoing pelvic irradiation. Our histopathology data showed that injection of APRGF into the anatomic regions involved in sexual responsiveness via growth factors and cytokines might promote differentiation of pluripotent stem cells and induction of neoangiogenesis, glandular proliferation (Skene’s glands), fibrosis reduction and new neuronal growth. These changes eventually could be improved physiologic responsiveness. Hence, we presumed that amended vascularity and neuronal regrowth in the vagina and in the clitoral area could augment sexual responsiveness through enhancing blood flow to the area, particularly in cases where hormonally independent vaginal atrophy contributes to FSD. In addition to increased blood flow, collagen and sensory nerve regrowth might relieve coital discomfort as well as enhance vaginal sensitivity.

Also, increased blood flow in the clitoris, if induced by PRP injections, could also lead to improved arousal and orgasm. Our recent studies demonstrated that PRP induces regrowth of new tissue through activation of pluripotent stem cells and also angiogenesis in average 5 weeks after injection. These cells are capable of differentiating into several tissue types when stimulated by growth factors produced by activated platelets. (Mostafaei et al.; Mohammadi et al., 2017a; Mohammadi et al., 2017b). Platelets contained granules and growth factors, such as TGF-β, PDGF, IGF, VEGF, EGF and FGF-2 (Christgau et al., 2006), which stimulate the inflammatory cascade and healing process. PDGF plays an important role in cell proliferation, angiogenesis tissue and mucous regeneration (Kim and Yun, 2007; Alio et al., 2012; Pal et al., 2012; Scalfani and McCormick, 2012). Angiogenesis can be effective in the production of natural vaginal discharge and lubrication during intercourse. PDGF is a required element in the cellular division for fibroblasts, a type of connective tissue cell that is the most effective factor in reducing vaginal fibrosis and stenosis and increasing...
vaginal flexibility (Song et al., 2005; Anitua et al., 2009). Since based on our previous study angiogenesis need more than 4 weeks, we hypothesized that this no significant differences in CD34 IHC might be related to the time of taking biopsy (Mohammadi et al., 2017b). In line with our findings, Charles Runels and colleagues in their pilot study demonstrated that specifically placed intra-vaginal and intra-clitoral PRP injections could be an effective method to treat certain types of FSD, especially in the areas of desire, arousal, lubrication, and orgasm. In spite of our findings, improvement in satisfaction and pain were not statistically significant in their study (Runels, 2014).

The result of Behnia et al; showed that PRP represents a potential alternative to topical steroids for treatment of vulvovaginal autoimmune conditions such as lichen sclerosus. They hypothesized growth factors released by platelets, monocytes, and nutrients have an important role in phagocytosis of fibrotic tissue, inflammation reduction, angiogenesis stimulation, and collagen III synthesis (Behnia-Willison et al., 2016).

In conclusion, to sum up, in this study, ten patients had dyspareunia and extraordinarily vaginal stenosis and dryness mucous were treated with the APRGF as prescribed. Our patients reported better sexual function and showed better vaginal function indexes, after APRGF injection. However results were positive but mandate further investigation due to limitation of our small pilot study,

Conflicts of Interest

The authors report no conflicts of interest.

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