· Original article ·

Pattern scan laser versus single spot laser in panretinal photocoagulation treatment for proliferative diabetic retinopathy

Shu Zhang, Guo-Fan Cao, Xiang-Zhong Xu, Cheng-Hu Wang

In – patient Department No. 2, Affiliated Eye Hospital of Nanjing Medical University, Nanjing 210029, Jiangsu Province, China

Correspondence to: Shu Zhang. In-patient Department No. 2, Affiliated Eye Hospital of Nanjing Medical University, Nanjing 210029, Jiangsu Province, China. zhangs60@ aliyun. com Received:2016-10-14 Accepted:2016-12-08

多点与单点激光治疗增殖性糖尿病视网膜病变 的疗效比较

张 舒,曹国凡,徐向忠,王成虎

(作者单位:210029 中国江苏省南京市,南京医科大学附属眼科 医院二病区)

作者简介:张舒,毕业于东南大学,本科,主任医师,副教授,研究 方向:眼底病。

通讯作者:张舒. zhangs60@ aliyun. com

摘要

目的:观察577-nm多点激光用于新近诊断的增殖性糖尿病视网膜病变(proliferative diabetic retinopathy, PDR)行全视网膜光凝(panretinal photocoagulation, PRP)的疗效。

方法:该前瞻性对照研究共纳入 32 例 40 眼 PDR 患者,随 机分为 2 组,每组 16 例患者(20 眼)。第 1 组采用多点激 光(pattern scan laser, PSL)行 PRP 治疗,第 2 组采用单点 激光(single spot laser, SSL)治疗。所有患者在 PRP 治疗 前与最后一次 PRP 治疗后 3mo 均行荧光素眼底血管造影 检查(fundus fluorescein angiography, FFA),以判断是否存 在无灌注区。激光点数、完成 PRP 的治疗次数、治疗时长 以及患者在治疗时的疼痛程度均作为判断指标。

结果:PSL 组患者需3次治疗以完成 PRP,而SSL 组需4次 完成。第1组每次治疗时间为7.3±2.3min,较第2组的 时间(13.2±4.1min)明显缩短(t₃₈=5.596, P<0.01)。第 1组的治疗疼痛指数较第2组明显降低(P<0.01)。最后 一次 PRP 治疗 3mo 后复查 FFA,第1组有5眼(25%)出 现无灌注区,而第2组有8眼(40%)出现无灌注区,需要 进一步治疗。

结论:多点激光用于 PDR 行 PRP 治疗较单点激光有着明显的优势。它具有效率高,治疗疼痛轻,疗效好的特点。 关键词:全视网膜光凝;多点激光;增殖性糖尿病视网膜 病变

引用:张舒,曹国凡,徐向忠,王成虎. 多点与单点激光治疗增殖性糖尿病视网膜病变的疗效比较.国际眼科杂志 2017;17(2):205-208

Abstract

AIM: To investigate the efficacy of 577-nm pattern scan laser in panretinal photocoagulation (PRP) treatment in newly diagnosed proliferative diabetic retinopathy (PDR).
METHODS: Prospective and comparative observation was performed in totally 32 patients with high-risk PDR. They were randomly divided into group 1 (using pattern scan laser, PSL) and 2 (using single spot laser, SSL), each containing 16 subjects to which totally 20 eyes received PRP. Non-perfusion region was identified with fundus fluorescein angiography (FFA) before and 3mo after final PRP. The advantage of PSL was verified in terms of the number and the duration of PRP sessions needed for satisfactory outcomes, and the pain score.

• RESULTS: Three PRP sessions were needed for each eye to complete the treatment using PSL, while 4 sessions were needed using SSL. The duration of each session with PSL in group 1 was 7.3 ± 2.3min, which was significantly shorter than that with SSL in group 2 (13.2± 4.1, t_{38} = 5.596, *P*<0.001). Treatment-related pain score was also significantly lower in group 1 than in group 2 (*P*< 0.01). Three months after the final PRP, the number of eyes with retinal non – perfusion regions that required additional treatment was 5 (25%) in group 1 and 8 (40%) in group 2.

• CONCLUSION: PSL showed clear advantages over SSL in the PRP treatment of PDR, not only in the improved efficacy, but also in the reduction of pain and the improvement of effectiveness.

• KEYWORDS: panretinal photocoagulation; pattern scan laser; proliferative diabetic retinopathy

DOI:10.3980/j.issn.1672-5123.2017.2.03

Citation: Zhang S, Cao GF, Xu XZ, Wang CH. Pattern scan laser versus single spot laser in panretinal photocoagulation treatment for proliferative diabetic retinopathy. *Guoji Yanke Zazhi* (*Int Eye Sci*) 2017;17(2):205–208

INTRODUCTION

 severe vision loss can be significantly reduced^[2-4]. The major benefit of PRP therapy is the reduction or cessation of neovascularization, resulting in a decrease of retinal blood flow^[9-13]. However, the PRP treatment is considered painful, time consuming and needs to be repeated in several sessions^[7,14]. In the present study, we evaluated the advantages of using pattern scan laser (PSL) over the single spot laser (SSL) that was used conventionally in PRP therapy.

SUBJECTS AND METHODS

Subjects A prospective and comparative study with a between-group design was conducted in the Affiliated Eye Hospital of Nanjing Medical University. Data of PDR treatments were collected between Jan. 2014 and Feb. 2015 from totally 32 newly found patients with high-risk PDR, 16 subjects each randomly (listed in advance by table of random numbers) allocated in group 1 with PSL and in group 2 with SSL. In each group, totally 20 eyes were treated. The subjects in the two groups were matched in age and gender and the duration/history of diabetes (Table 1). The patients all had diabetes (based on WHO criteria) and were eligible for focal laser treatment as defined by the ETDRS criteria^[15]. This study was adhered to the tenets of the Declaration of

Helsinki and was approved by the Medical Research Ethics Committee of the Affiliated Eye Hospital of Nanjing Medical University. Written informed consent was obtained from all participants.

PRP Treatment A 577 – nm laser photocoagulator (Supra Scan, Quantel Medical, France) was employed for PRP with both PSL and SSL. In 18 spot PSL, the major settings of the equipment include power of 450 - 800 mW and duration of 0.035–0.050s for each spot of 200-300 µm. In the classical SSL, the corresponding settings were power of 250-450 mW and duration of 0.25–0.30s for each spot of 200-300 µm. The settings were adjusted in above ranges so that the retinal spots showed moderate whitening immediately after being burned by the laser beam.

The interval between consecutive sessions was 2wk in both groups. Before every following session, visual acuity and macula OCT were examined. If more than 20% drop in visual acuity and/or 20% increases in the thickness of macula was/ were found, the following session would be postponed till these laser induced side reactions alleviated.

Evaluations The evaluations involve several measures. 1) Fundus fluorescein angiography (FFA) test was performed in each patient before and 3mo after the final PRP to verify retinal non-perfusion regions;2) The duration of each session was recorded from the time when the contact lens was placed to the time when it was taken off; 3) The pain score was quantified in ranking by each patient immediately after each session in a way reported previously^[16-17]: 0 = no pain, 1 = little pain, 2 = moderate pain, 3 = severe pain. A standard instruction was used in the ranking to avoid bias.

Statistical Analysis The quantitative data are presented as mean \pm SD. The treatment duration was compared between groups using Student's *t*-test. The pain score was compared 206



Figure 1 A fundus image taken immediately after the first treatment from a group 1 subject. The burned spots show grey to moderate white and spread evenly in lower posterior retina.

Table 1 Pretreatment data of patients

Patients	Group 1	Group 2
No. of patients (eyes)	16(20)	16(20)
M:F	8:8	7:9
Mean age (a, Mean±SD)	46.5±9.6	50.2±11.2
History of diabetes		
≤10a	8	8
>10a	8	8

Table 2	Pain scores	as indicated	by the	number of eyes
---------	-------------	--------------	--------	----------------

Groups	Pain scores					
	0	1	2	3		
Group 1 (eyes)	4	10	5	1		
Group 2 (eyes)	1	3	12	4		
Р	< 0.01					

P values are the results of post-hoc pairwise comparisons (Mann-Whitney Rank Sum Test) between groups.

using ranking comparison (Mann–Whitney Rank Sum Test). Differences are considered significant when P<0.01.

RESULTS

In group 1 (PSL group), three sessions were needed to complete PRP. In each session, approximately 400 spots were burned(18 spots at one shoot), which were spread evenly in lower posterior retina (Figure 1), upper posterior retina, and the third session, peripheral retina and subsidy. In group 2 (SSL group), four sessions of treatment were needed for each eye, and approximately 250 spots were burned in one of the 4 quarters of fundus in each session: lower nasal + macular temporal, upper temporal, lower temporal, upper nasal respectively. No treatment session was postponed due to the laser induced side reactions.

The duration of each PRP session was 7.3±2.4min in group 1, which was 44.5% shorter than that of group 2 (13.2±4. 1min, t_{38} = 5.596, *P*<0.001).

Table 2 showed the ranked pain perception score by the subjects. A rank test (Mann – Whitney method) showed significantly less pain in group 1 than in group 2 (P<0.01). Shown by FFA 3mo after final session of PRP, 5 eyes (25%) in group 1 were found with retinal non-perfusion regions, and this number was 8 (40%) in group 2.

DISCUSSION

PRP currently is generally acknowledged as the mainstay and gold standard therapy for PDR since the studies on diabetic Retinopathy were published^[3-6]. It is estimated that about 60% PDR patients respond to laser PRP with retinal neovascularization regression within 3mo^[18]. However, PRP is a destructive procedure, which is time consuming, often painful, and cannot be completed without multiple sessions of treatment. PRP is also often accompanied by a decrease in peripheral visual field and an increase in the risk of macular edema^[6]. In order to minimize the damage, laser spots must be evenly-distributed in each session in which hundreds of spots were burned rapidly over the whole retina. PSL technique was more suitable for this purpose^[12,14,19-22]. Using this technique, many more spots can be burned in one session. Therefore, fewer sessions are needed. The result of the present study is consistent with that reported by others^[23]. Furthermore, many more spots were burned in group 1 (-1200 points/eye) in a shorter period of time than in group 2 (-1000 points/eye). This was because 18 spots were burned in one shoot. Although more spots and larger total area were burned in one session using PSL, no more side reaction was found.

The primary effect of laser treatment is a thermal injury induced at the level of the retinal pigment epithelium (RPE). But concurrent damage to adjacent retinal photoreceptors and choriocapillaris occurs as a result of heat transmission. The evidence of thermal spread from the RPE is shown as the whitening of the retinal, which is taken as the index of successful laser burn. This damage may result in macular edema which could induce loss of central vision, decreased contrast sensitivity and reduced visual fields^[17-18,24-27]. In conventional PRP using SSL, laser pulse duration is 0.25-0.30s, which is much longer than the duration in PSL. Such long pulse duration is needed in order to produce visible whitening on the isolated spot. On the other hand, the effective burning can be achieved with shorter duration in PSL, in which 18 spots were burned in a close neighborhood. This makes it possible to reduce the pulse duration^[28-29].

The shorter duration likely confines the damage to the RPE/ choroid melanin granules. It has been shown that laser exposure of shorter duration affects mainly the RPE, and with little or no damage on the photoreceptors or choriocapillaris. In addition, the heat affected area in each spot would also be reduced when short duration is used. This is accompanied by reduced inflammatory cytokines in retina as compared with the results of long pulse duration in SSL^[28]. The shorter duration for each treatment and the less damage of the tissues around RPE are likely the reason for the low pain score in group 1.

Related to the larger number of spots burned in PSL, the treatment was extended to more peripheral area of retina. This appears to be beneficial in neovascularization clear – up and regression. Correspondingly, a longer lasting of the treatment effect using PSL was supported by less reoccurrence of non – perfusion region that required additional treatment in this group.

Although different laser beams, such as that with 532 - nm wavelength, had been used in PSL, here, we chose the 577-nm yellow laser beam, which was proved to be more effective than other available lasers^[30-31].

Several limitations exist in the present study, including an overall small sample size. We recommend further studies including a randomized treatment trial to make a clear comparison across different methods of PRP.

In conclusion, the present study demonstrated the advantage of 577-nm pattern scan laser on the management of patients with PDR over conventional single spot laser PRP. The evidence supports the choice of the pattern scan laser substitute to replace single spot laser in future.

REFERENCES

1 Chew EY, Kim J, Coleman HR, Aiello LP, Fish G, Ip M, Haller JA, Figueroa M, Martin D, Callanan D, Avery R, Hammel K, Thompson DJ, Ferris FL, 3^{rd} . Preliminary assessment of celecoxib and microdiode pulse laser treatment of diabetic macular edema. *Retina* 2010; 30(3): 459–467

2 Jung JJ, Gallego-Pinazo R, Lleo-Perez A, Huz JI, Barbazetto IA. NAVILAS laser system focal laser treatment for diabetic macular edema – one year results of a case series. *Open Ophthalmol J* 2013;7:48–53

3 Tapp RJ, Svoboda J, Fredericks B, Jackson AJ, Taylor HR. Retinal photography screening programs to prevent vision loss from diabetic retinopathy in rural and urban Australia: a review. *Ophthalmic Epidemiol* 2015;22(1):52–59

4 Demers – Turco P. Providing timely and ongoing vision rehabilitation services for the diabetic patient with irreversible vision loss from diabetic retinopathy. J Am Optom Assoc 1999;70(1):49-62

5 Sinawat S, Rattanapakorn T, Sanguansak T, Yospaiboon Y, Sinawat S. Intravitreal bevacizumab for proliferative diabetic retinopathy with new dense vitreous hemorrhage after full panretinal photocoagulation. *Eye* (*Lond*) 2013;27(12):1391–1396

6 Early photocoagulation for diabetic retinopathy. ETDRS report number
9. Early Treatment Diabetic Retinopathy Study Research Group. Ophthalmology 1991;98(5 Suppl):766-785

7 Avitabile T, Bonfiglio V, Castiglione F, Castaing M, Contarino F, Mistretta A. Severe proliferative diabetic retinopathy treated with vitrectomy or panretinal photocoagulation: a monocenter randomized controlled clinical trial. *Can J Ophthalmol* 2011;46(4):345-351

8 Gurelik G, Coney JM, Zakov ZN. Binocular indirect panretinal laser photocoagulation for the treatment of proliferative diabetic retinopathy. *Ophthalmic Surg Lasers Imaging* 2004;35(2):94-102

9 Yang CM. Surgical treatment for severe diabetic macular edema with massive hard exudates. *Retina* 2000;20(2):121-125

10 Zhu Y, Zhang T, Wang K, Xu G, Huang X. Changes in choroidal thickness after panretinal photocoagulation in patients with type 2 diabetes. *Retina* 2015;35(4):695-703

11 Flynn HW, Jr., Chew EY, Simons BD, Barton FB, Remaley NA, Ferris FL, 3rd. Pars plana vitrectomy in the Early Treatment Diabetic Retinopathy Study. ETDRS report number 17. The Early Treatment Diabetic Retinopathy Study Research Group. *Ophthalmology* 1992;99 (9):1351-1357

12 Alasil T, Waheed NK. Pan retinal photocoagulation for proliferative diabetic retinopathy: pattern scan laser versus argon laser. *Curr Opin Ophthalmol* 2014;25(3):164-170

13 Zhu Y, Zhang T, Wang KY, Xu GZ. Prognostic value of multifocal electroretinography and optical coherence tomography in eyes undergoing panretinal photocoagulation for diabetic retinopathy. *Invest Ophthalmol Vis Sci* 2014;55(10):6358-6364

14 Chappelow AV, Tan K, Waheed NK, Kaiser PK. Panretinal

photocoagulation for proliferative diabetic retinopathy: pattern scan laser versus argon laser. Am J Ophthalmol 2012;153(1):137-142. e2

15 Fundus photographic risk factors for progression of diabetic retinopathy. ETDRS report number 12. Early Treatment Diabetic Retinopathy Study Research Group. *Ophthalmology* 1991;98(5 Suppl): 823-833

16 Sauder G, Jonas JB. Topical anesthesia for penetrating trabeculectomy. *Graefes Arch Clin Exp Ophthalmol* 2002;240(9): 739-742

17 Kernt M, Cheuteu RE, Cserhati S, Seidensticker F, Liegl RG, Lang J, Haritoglou C, Kampik A, Ulbig MW, Neubauer AS. Pain and accuracy of focal laser treatment for diabetic macular edema using a retinal navigated laser (Navilas). *Clin Ophthalmol* 2012;6:289–296

18 Arnarsson A, Stefansson E. Laser treatment and the mechanism of edema reduction in branch retinal vein occlusion. *Invest Ophthalmol Vis Sci* 2000;41(3):877-879

19 Yun SH, Adelman RA. Recent developments in laser treatment of diabetic retinopathy. *Middle East Afr J Ophthalmol* 2015;22(2):157-163 20 Nagpal M, Marlecha S, Nagpal K. Comparison of laser photocoagulation for diabetic retinopathy using 532 - nm standard laser versus multispot pattern scan laser. *Retina* 2010;30(3):452-458

21 Velez – Montoya R, Guerrero – Naranjo JL, Gonzalez – Mijares CC, Fromow–Guerra J, Marcellino GR, Quiroz–Mercado H, Morales–Canton V. Pattern scan laser photocoagulation: safety and complications, experience after 1301 consecutive cases. *Br J Ophthalmol* 2010;94(6): 720–724

22 Yadav NK, Jayadev C, Rajendran A, Nagpal M. Recent developments in retinal lasers and delivery systems. *Indian J Ophthalmol* 2014;62(1):50-54

23 Muraly P, Limbad P, Srinivasan K, Ramasamy K. Single session of Pascal versus multiple sessions of conventional laser for panretinal photocoagulation in proliferative diabetic retinopathy: a comparitive study. *Retina* 2011;31(7):1359-1365

24 Ogata N, Tombran – Tink J, Jo N, Mrazek D, Matsumura M. Upregulation of pigment epithelium – derived factor after laser photocoagulation. *Am J Ophthalmol* 2001;132(3):427-429

25 Ogata N, Ando A, Uyama M, Matsumura M. Expression of cytokines and transcription factors in photocoagulated human retinal pigment epithelial cells. *Graefes Arch Clin Exp Ophthalmol* 2001;239(2):87–95 26 Shinoda K, Ishida S, Kawashima S, Wakabayashi T, Uchita M, Matsuzaki T, Takayama M, Shinmura K, Yamada M. Clinical factors related to the aqueous levels of vascular endothelial growth factor and hepatocyte growth factor in proliferative diabetic retinopathy. *Curr Eye Res* 2000;21(2):655–661

27 Spranger J, Hammes HP, Preissner KT, Schatz H, Pfeiffer AF. Release of the angiogenesis inhibitor angiostatin in patients with proliferative diabetic retinopathy: association with retinal photocoagulation. *Diabetologia* 2000;43(11):1404–1407

28 Ito A, Hirano Y, Nozaki M, Ashikari M, Sugitani K, Ogura Y. Short pulse laser induces less inflammatory cytokines in the murine retina after laser photocoagulation. *Ophthalmic Res* 2015;53(2):65–73

29 Muqit MM, Marcellino GR, Henson DB, Young LB, Patton N, Charles SJ, Turner GS, Stanga PE. Optos – guided pattern scan laser (Pascal) – targeted retinal photocoagulation in proliferative diabetic retinopathy. *Acta Ophthalmol* 2013;91(3):251–258

30 Hirano T, Iesato Y, Murata T. Multicolor pattern scan laser for diabetic retinopathy with cataract. Int J Ophthalmol 2014; 7(4): 673 -676

31 Hirano T, Iesato Y, Imai A, Toriyama Y, Kikushima W, Murata T. Effect of Laser wavelength on delivering appropriate laser burns through the opaque lens using a pattern scan laser. *Ophthalmic Res* 2014;51(4): 204–209