Comparison of 2-wall versus 3-wall orbital decompression against dysthyroid optic neuropathy in visual function

A retrospective study in a Chinese population

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Abstract

To compare visual function of 2-wall (medial and lateral) versus 3-wall (medial, lateral, and inferior) orbital decompression in patients with dysthyroid optic neuropathy (DON).

A total of 52 eyes of 37 patients underwent orbital decompression for DON between 2013 and 2019 were retrospectively reviewed. Two- or 3-wall decompression was performed in 31 eyes of 23 patients and 21 eyes of 14 patients, respectively. We examined best-corrected visual acuity (BCVA), visual field mean deviation (MD) and pattern standard deviation (PSD), pattern-reversed visual evoked potential (PVEP) for P100 latency and amplitude at 60 and 15 arcmin stimulation checkerboard size, as well as proptosis using Hertel exophthalmometry.

Whether 2-wall or 3-wall decompression, all parameters of visual function were improved after surgery (all \( P < .05 \)). The improvement in BCVA, MD, and PSD was not statistically significant between groups (all \( P > .05 \)). Proptosis reduction was higher after 3-wall decompression (\( P = .011 \)). Mean increase in P100 amplitude after 3-wall decompression was statistically higher than that of after 2-wall decompression at 60 and 15 arcmin (\( P = .045 \) and \( .020 \), respectively), while the mean decrease in P100 latency was similar between the groups (\( P = .821 \) and \( .655 \), respectively). Six patients (66.67%) had persistent postoperative diplopia and 1 patient (20%) had new-onset diplopia in 3-wall decompression group, which were higher than in 2-wall decompression group (46.15% persistent postoperative diplopia and no new-onset diplopia).

Both 2-wall and 3-wall decompression can effectively improve visual function of patients with DON. Three-wall decompression provides better improvement in P100 amplitude and proptosis, however new-onset diplopia is more common with this surgical technique.

Abbreviations: BCVA = best-corrected visual acuity, DON = dysthyroid optic neuropathy, MD = mean deviation, PSD = pattern standard deviation, PVEP = pattern-reversed visual evoked potential.

Keywords: 2-wall orbital decompression, 3-wall orbital decompression, dysthyroid optic neuropathy, pattern-reversed visual evoked potential.

1. Introduction

Graves orbitopathy (GO) is an autoimmune disease involving intraorbital fat, extraocular muscles, and lacrimal glands. Its ocular signs and symptoms are complex, including proptosis, eyelid retraction, ocular dyskinesia, lagophthalmos, conjunctival congestion, diplopia, and exposure keratopathy. Dysthyroid optic neuropathy (DON) is a relatively uncommon but a serious complication of (GO), which can result in permanent vision loss if
not treated appropriately.\cite{11} It has an estimated incidence of 5% to 8.6% of patients with GO.\cite{12-14} While the exact mechanism of DON remains elusive, direct compression of the optic nerve by enlarged extraocular muscles\cite{15,16} stretching of the optic nerve by proptosis,\cite{17-19} orbital pressure,\cite{20,21} and inflammation\cite{22,23} have all been proposed.

There is controversial evidence regarding the optimum management strategy. Based on the current literatures, widely accepted treatments include systemic corticosteroids, orbital radiotherapy, and orbital decompression, in which intravenous methylprednisolone being the first line of treatment, and orbital decompression becoming indispensable if steroid therapy fails.\cite{13,14} Decompression surgery for DON is best performed after the disease has become inactive, but decompression may also be required in the active phase of GO for cases of DON that are refractory to medical treatment.\cite{13} DON has been managed with orbital decompression by various techniques.\cite{15-18} Orbital medial and inferior wall decompression for DON is the earliest approved surgical method, because the ethmoid sinus and maxillary sinus have huge space, which can alleviate the compression effect of medial and inferior rectus muscle on optic nerve.\cite{19} As the indications for surgical decompression for DON have expanded, new surgical approaches being developed for better postoperative cosmesis and fewer diplopia. Surgical techniques such as the deep lateral wall decompression,\cite{20,21} balanced decompression,\cite{22-25} and endoscopic medial decompression\cite{26,27} have all been widely used in the treatment of DON.

With the diversification of orbital decompression surgical methods, single surgical methods can no longer meet the requirements of surgery, and combined decompression of multiple surgical methods has gradually become a new development direction. However, there is no consensus among orbital surgeons about how much decompression should be performed to treat DON. Although orbital decompression is usually restorative in most cases, some patients do not respond well even with orbital decompression, and may still require either systemic immunosuppression or further decompression following initial surgery to improve optic nerve function.\cite{28,29}

Many studies compared different surgical techniques, such as lateral versus medial decompression, with or without inferior decompression or fat removal, and balanced decompression (medial and lateral) alone. But there are scanty evidences to elucidate the differences of visual function from a perspective of monitoring pattern-reversed visual evoked potential between 2-wall and 3-wall decompression. This retrospective study aimed to compare the therapeutic effects of 2-wall versus 3-wall orbital decompression on visual function in patients with DON.

2. Methods

2.1. Patients

Our study is a retrospective case series. The medical records of all the patients affected by DON and treated with 2-wall or 3-wall orbital bony decompression combined with orbital fat removal performed at Department of Ophthalmology, Union Hospital, Tongji Medical College, Huazhong University of Science and Technology between 2013 and 2019 were evaluated. This study was approved by the Ethics Committee of Union Hospital affiliated to Tongji Medical College, Huazhong University of Science and Technology, Wuhan, China, and was conducted in strict accordance with the Helsinki Declaration. All participants provided written informed consent.

Those patients that followed up for at least 3 months after the surgery were included in the study. One patient died and 5 were lost to follow-up. The last follow-up was either the last consultation in our clinic in those patients without further surgical interventions or the last consultation before the next surgical step in rehabilitative surgery (e.g., extraocular muscle surgery or lid surgery). The patients were excluded from study if they had the presence of any ocular disease affecting vision such as diabetic retinopathy, corneal ulcer, glaucoma, macular disease, and so on. Finally, the data from 37 cases (52 eyes) were recorded, including the age, sex, smoking status, thyroid function, previous therapy for GO, time elapsed from the first ocular symptoms to orbital surgery, presence of DON, orbital decompression technique, follow-up duration after orbital surgery, related surgical complications, and additional medical and/or surgical interventions after surgery.

All of the patients underwent full ophthalmic examinations both preoperatively and postoperatively, including clinical activity score (CAS), best-corrected visual acuity (BCVA), color vision, relative afferent pupillary defect (RAPD), fundoscopy, visual field by Humphrey 30-2 SITA-standard threshold strategy for mean deviation (MD) and pattern standard deviation (PSD), axial proptosis by Hertel exophthalmometry, and pattern-reversed visual evoked potential (PVEP) for P100 latency and amplitude at 60 and 15 arcm in stimulation checkerboard size. The diagnosis of DON was depended on the presence of any combination of visual deficits including visual acuity, visual field, and color vision. It was also supported by at least one of the apical crowding, optic disc edema, and afferent pupillary defect. Optic nerve dysfunction alone was responsible for all deficits.\cite{4,30}

2.2. Orbital decompression surgery technique

The surgery was performed under general anesthesia by the same ophthalmologist. The deep lateral wall decompression was performed via lateral canthal incision or extended upper eyelid crease incision. The subcutaneous tissue was separated to expose the outer rim of the orbit, and separation upwards to the lateral margin of the superior orbital margin and downwards to the superior margin of the zygomatic arch. The periosteum of the orbital margin was cut open by unipolar electrotome, and the anterior margin of the temporalis muscle attached to the orbital margin was cut back. Then, the lateral orbital bone flap was cut off with a chainsaw, with the anterior orbital bone cut off. After that, the anterior temporal muscle was dissected posteriorly to fully expose the temporal fossa. Combined with bone chisel, bone biting forceps and high-speed grinding drill, all the posterior bone of lateral wall of orbit was removed, deep to the transition of the dura and periosteum of the outer edge of the superior orbital fissure, up to the sphenoid crest and lacrimal fossa, and down to the upper edge of the inferior orbital fissure. Exposure of most of the anterior temporal lobe meninges was a sign of termination of lateral wall decompression. An average of 3 cm$^3$ of fat was removed from this region.

Medial orbital decompression was performed through a transcarnuncular incision. Blunt dissection was carried posteriorly and medially with scissors to expose the medial orbital wall just behind the posterior lacrimal crest. The orbital peristium was resected at 1.5 cm behind the crista lacrimalis posterior and the ethmoid paper template was bitten off by sinus forceps. It went up
to the level plate of the ethmoid bone, deep to the front of the inner edge of the optic canal, and down to the bone where the ethmoid bone meet the maxilla, namely orbital strut. The bone here was hard and about 3 mm thick, having a supporting effect on the orbital tissue, which can reduce the probability of diplopia caused by the muscle cone shifting inward and downward. It should be reserved. In severe cases, it could be removed or compressed to shift inward and downward. Separate the muscle spacers and shift the medial rectus and fat toward the ethmoid sinus.

If floor decompression was planned, a transconjunctival incision was performed which was extended by a conjunctival incision in the medial wall. The posterior 1/2 structure at the strut was removed, extending downwards to the posterior 1/2 of the inferior orbital wall, deep to the apex of the orbit, and beyond to the inferior orbital fissure, retaining the inferior orbital nerve groove.

After satisfactory eyeball retraction without obvious displacement was observed, the orbital outer margin bone flap was reset and fixed with titanium nail and titanium plate. Then, the transcaruncular incision, peristium, subcutaneous tissue, and skin incision were sutured successively. Postoperative compression bandaging was performed for 1 week, and intravenous broad-spectrum antibiotics and hemostatic drugs were routinely given at 3 days.

2.3. Statistical analysis

Data analysis was performed by using SPSS for Windows, version 20 (SPSS Inc., Chicago, IL). Best corrected Snellen acuity was converted to logarithm of minimal angle of resolution (LogMAR) units for statistical analyses. Eyes without formed visual acuity were assigned decimal visual acuity according to previous study. Wilcoxon signed rank test, independent t test, paired t test were used when necessary. If there were bilateral DON in 1 patient, data of left and right eyes were both used for analysis. P < .05 was considered statistically significant.

3. Results

3.1. Demographic characteristics

A total of 52 eyes of 37 patients who underwent orbital decompression for the treatment of DON were included in the study. All bilateral decompressions were of identical type on each side. All patients were previously treated either by radioiodine (n = 12) or thyroidectomy (n = 5) or antithyroid drug therapy (n = 36) and had normal serum FT3 and FT4 values at the time of decompression surgery as well as during follow-up. All patients previously received systemic steroids and 2 patients underwent additional orbital irradiation without sufficient or definite remission of GO prior to decompression. Because in most cases pretreatment was not coordinated by our hospital, both orbital irradiation and steroid therapy were administered at variable treatment and dosing regimes. There were 9 men and 28 women. The mean age at decompression was 48 years (range, 28–71 years). Two-wall decompression was performed in 31 eyes (23 patients) and 3-wall in 21 eyes (14 patients). The mean follow-up time was 22 months (range, 3–71 months).

There were no differences between the groups with respect to demographics, smoking habits, treatment, duration of GD, GO and DON, and thyroid function at preoperative visit (all P > .05).

3.2. Clinical outcome measures

Preoperative and postoperative clinical outcome measures were summarized in Table 1. There was no differences between the groups about preoperative value of LogMAR BCVA, MD, PSD (all P > .05), except for proptosis, which was greater in 3-wall decompression group (P < .001). Whether 2-wall decompression or 3-wall decompression, BCVA, MD, PSD, and proptosis were significantly improved after decompression surgery (all P < .001). Moreover, there were no significant differences between the 2 groups in improvement of these 3 parameters (all P > .05), except in proptosis (P = .011).

3.3. PVEP

The changes of PVEP after decompression surgery were shown in Table 2. Mean P100 amplitude increased significantly in 2-wall decompression group by 2.30 and 2.43 μV at 60 and 15 arcmin stimulation checkerboard size, respectively, corresponding to increases significantly in 3-wall decompression group by 4.89 and 4.49 μV at 60 and 15 arcmin stimulation checkerboard size, respectively. Mean increase in P100 amplitude after 3-wall decompression was statistically higher than that of after 2-wall.
decompression at 60 and 15 arcmin (P = .046 and .020, respectively) (shown in Fig. 1).

Mean P100 latency shortened significantly in 2-wall decompression group by 6.16 and 5.76 ms at 60 and 15 arcmin stimulation checkerboard size, respectively, corresponding to decreases significantly in 3-wall decompression group by 7.02 and 9.12 ms at 60 and 15 arcmin stimulation checkerboard size, respectively. Mean decrease in P100 latency was not statistically significantly different between the 2 groups (P = .821 and .655, respectively) (shown in Fig. 2).

3.4. Diplopia

Of the 23 patients with 2-wall decompression, 13 patients (56.52%) complained preoperative diplopia and of the 14 patients with 3-wall decompression, 9 patients (64.29%) complained preoperative diplopia, with no statistical difference between the 2 groups (P = .738). Six patients (66.67%) had persistent postoperative diplopia and 1 patient (20%) had new-onset diplopia in 3-wall decompression group, which were higher than in 3-wall decompression group (46.15% persistent postoperative diplopia and no new-onset diplopia). As to the resolution rate of diplopia, 2-wall decompression group (53.85%) was higher than 3-wall decompression group (33.33%).

4. Discussion

In this study, we demonstrated that both 2-wall and 3-wall decompression techniques could effectively improve proptosis, visual acuity impairment, visual field defect, and PVEP abnormality in patients with DON. However, 3-wall decompression provided better improvement of visual functions regarding the parameters of P100 amplitude. To the best of our knowledge, this is the first study that showed 3-wall decompression was superior to 2-wall in improving P100 amplitude. The results could provide theoretical guidance to the ophthalmologist for the choice of orbital decompression technique.

PVEP reflects the transmission function of visual signals from retinal ganglion cells to occipital visual cortex. Previous studies showed that PVEP was a valuable tool in the diagnosis and follow-up of DON, as DON was significantly associated with P100 amplitude loss and delayed deflection.\[4,34,35\] In addition, PVEP was able to identify even subclinical courses of DON with
high sensitivity.\textsuperscript{36} Several studies suggested that orbital decompression exert beneficial effects on improvement of P100 latency and amplitude in patients with DON.\textsuperscript{33,37,38} Moreover, Tsaloumas et al.\textsuperscript{33} argued that P100 latency was shortened and P100 amplitude was increased after orbital decompression, but the improvement of P100 amplitude was more obvious than that of P100 latency, providing parallel results with our works, which indicated that P100 amplitude was more sensitive to the change of orbital pressure. Our study demonstrated that both surgical techniques significantly improved PVEP in patients with DON, in which the increase of P100 amplitude at 15 and 60 arcmin stimulation checkerboard size after 3-wall decompression was significantly higher than that after 2-wall decompression ($P = .046$ and .020, respectively).

As for the mechanism of DON under the crowded orbital condition, it has been demonstrated that the compression lesions of DON could have double effects of demyelination and axonal damage.\textsuperscript{34,35} Because the loss of myelin sheath and the restored myelin sheath are extremely thin, uneven, and underdeveloped in insulation, the conduction of nerve impulse changes from jump conduction between Ranvier’s nodes to crawling conduction along the repaired myelin sheath, thus slowing down the conduction and prolonging P100 latency. Another reason for P100 latency delay is the lack of nerve fibers with thicker diameters and faster conduction. In addition, the degeneration and necrosis of optic nerve axons result in the decrease of axonal number, consequently triggering the decrease of electrical activity and VEP amplitude. Both 2-wall and 3-wall orbital decompression effectively enlarges the orbital volume and decreases the intra-orbital pressure, which is more distinct in 3-wall decompression. Orbital decompression relieves the compression of the remaining axons and is beneficial to the regeneration of myelin sheath, and also the reconstruction of impulse conduction, thus could improve the optic nerve function and also relieve PVEP abnormalities. To our delight, the present study showed for the first time that 3-wall decompression was superior to 2-wall in improving P100 amplitude.

The main pathological changes of GO are inflammatory cell infiltration, edema, or extracocular muscle thickening of the extracocular muscles, which leads to the increased orbital contents not compatible with the volume of the orbital cavity, with the proptosis and the increase of the orbital pressure appearing.\textsuperscript{39} Proptosis is an important index in the clinical diagnosis of GO, and also the evaluation of the therapeutic effect. The improvement of proptosis is dependent on the number of orbital bone walls decompressed and the amount of fat removed. It has been demonstrated that proptosis reduction was $7.2 \pm 1.9$ mm for 3-wall decompression, compared with $5.1 \pm 1.3$ mm for 2-wall decompression.\textsuperscript{40} This difference was found to be even more marked in the present study, with an average $7.53 \pm 3.40$ mm of proptosis reduction following 3-wall decompression versus $5.16 \pm 2.34$ mm after 2-wall decompression ($P = .011$). Although the main goal of orbital decompression is not to relieve proptosis but to reserve the visual function, better cosmesis, and more eyeball retrospect could be achieved by 3-wall decompression if the preoperative proptosis is severe.

According to the results of Korkmaz and Konuk,\textsuperscript{40} 3-wall decompression was superior to 2-wall in improving both MD and PSD, while the vision acuity improvement is comparable. However, we demonstrated that 2-and 3-wall decompression effectively improved the MD and PSD, and vision acuity in a similar manner. Visual field is a subjective examination, and there are some patients whose accuracy is affected by poor cooperation. This is similar to the visual acuity test, so there is no doubt that the results are different from other researchers. Orbital decompression effectively decreases the intra-orbital pressure, showing beneficial effect on improvement of visual field in patients with DON. This is parallel with several studies.\textsuperscript{37,40}

A large number of clinical evidences demonstrated that orbital inferior wall decompression, especially combined with medial wall decompression, was inclined to destroy the “orbital strut” of the supporting structure of the orbital inferomedial wall, resulting in the occurrence of diplopia and the displacement of the eyeball after surgery.\textsuperscript{25,31} It was suggested that balanced decompression technique was associated with more favorable results regarding the postoperative new diplopia rates.\textsuperscript{16,25} Our study also analyzed the occurrence of diplopia before and after surgery in both groups. We found that the resolved diplopia rate of 2-wall decompression (53.85%) was higher than that of 3-wall decompression (33.33%), and the new-onset diplopia rate of 3-wall decompression (20%) was significantly higher than that of 2-wall decompression (0%). In 3-wall decompression group, the development of postoperative significant new-onset diplopia could have been induced by inferomedial shifting of the orbital contents after simultaneous removal of the medial and inferior orbital walls. Preservation of the inferior orbital wall provides support for orbital soft tissues from below and prevents inferomedial shifting of the muscle cone and orbital fibroconnective septal system. This might be the reason for the absence of postoperative new-onset diplopia in 2-wall decompression group.

Interpretation of the results of this study must take into account its limitations. First, its retrospective nonrandomized study design results in incomplete and inaccurate data, such as the statistics of dyschromatopsia and subjective diplopia which is absent of objective data for color vision and diplopia assessment. Second, there were statistically significant differences in the amount of preoperative proptosis due to this design. While these preoperative differences make direct outcome comparisons difficult, we contend that the most critical determinant of operative success is the degree of postoperative change which we have showed in the results. Third, the sample size is relatively small. Because of the relative infrequency of DON, surgeon-dependent variations in surgical technique, and difficult follow up, the number of cases is relatively small. More cases can be accumulated unless the number of cases is selected for a longer period of time. To design a large multicenter prospective randomized comparative clinical trial is a pleasurable choice, which can also address the first problem above.

We conclude that 2-wall and 3-wall decompression effectively treated patients with DON. Both techniques significantly improved the visual acuity, visual field, and P100 latency in a similar manner. However, 3-wall decompression provides better improvement in P100 amplitude and proptosis, while the resolved diplopia rate is lower and new onset diplopia is more common with this surgical technique. Afterwards, preoperative ocular evaluation of patients with DON is essential and indispensable to the choice of surgical techniques and the optimal treatment.

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