

ORIGINAL RESEARCH

Hypnosis intervention for the management of pain perception during cataract surgery

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Objective: To investigate the effectiveness of hypnosis in pain management during cataract surgery.

Methods: Male or female patients with bilateral age-related cataract who wished to have both eyes subjected to phacoemulsification surgery were preliminarily admitted. Immediately after the first-eye surgery, each patient was evaluated for pain using the visual analog scale (VAS), and patients with a VAS score > 1 were enrolled. By using block randomization, the enrolled patients were allocated to either the treatment group, which received a hypnosis intervention before the scheduled second-eye surgery, or the control group, which did not undergo hypnosis. The levels of anxiety, pain, and cooperation were evaluated independently by the patients and the surgeon. Results: During the intraoperative pain assessment, 5%, 34%, 38%, and 23% of patients in the control group reported experiencing no pain, mild pain, moderate pain, and severe pain, respectively. In contrast, in the hypnosis group, 18%, 56%, 15%, and 11% of patients reported experiencing no pain, mild pain, moderate pain, and severe pain, respectively, which showed significant differences between the groups (P<0.005). The evaluation of anxiety level showed that the mean score in the control group and hypnosis group was 11.77±0.32 and 6.64±0.21, respectively, revealing a highly significant difference between the two groups (P<0.005). The assessment of patient cooperation showed that only 5% and 18% of patients in the control group and 18% and 36% of patients in the hypnosis group showed excellent and good cooperation, respectively, while 47% of patients in the control group and only 24% of patients in the hypnosis group exhibited poor cooperation, revealing significant differences between the groups (P<0.005). **Conclusion:** Hypnosis may be considered as an auxiliary measure in cataract surgery, especially for patients who experienced obvious pain during the first-eye surgery.

Keywords: pain, patient satisfaction, cataract surgery, hypnosis

Introduction

Phacoemulsification is the most effective cataract surgery currently performed and is characterized by a smaller wound and reduced surgery time. Given the advancements in equipment and the proficiency of surgical skills, topical anesthesia is the preferred method in phacoemulsification surgery to avoid the risks associated with local (retrobulbar or peribulbar) anesthesia, such as chemosis, ecchymosis, retrobulbar hemorrhage, globe penetration, or perforation; however, phacoemulsification with topical anesthesia is not a completely painless procedure, with patients frequently experiencing pain during the surgery. In a study comparing topical anesthesia and local anesthesia for phacoemulsification surgery, about

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30.1% of patients with topical anesthesia complained of intraoperative pain, while only 6.7% of patients with local anesthesia reported intraoperative pain. In another study, 78.3% of patients reported experiencing pain during phacoemulsification surgery with topical anesthesia; when the intensity of pain was evaluated, the percentage of patients reporting mild, moderate, and intense pain was 35.9%, 25.0%, and 17.4%, respectively. Although the use of topical anesthesia may help avoid the potential complications of local anesthesia, the possibility of the eye moving or closing due to pain may represent a significant handicap; thus, preoperative administration of analgesics or the use of an additional sedative measure may increase patient comfort and cooperation.

Hypnosis is a method of psychotherapy that can modulate the phenomenological aspects of conscious experience, relieve pain, relax the body, and reduce anxiety. 8,9 A limited number of studies have demonstrated the efficacy of clinical hypnosis in the treatment of pain and anxiety in severe diseases. 10–12 In the context of anesthesia, hypnosis can be used alone or in combination with the drugs typically used in anesthetic regimens, and increasing evidence supports the value of hypnosis in reducing patient stress and anxiety and inducing relaxation. 13,14

In this study, we discuss the effectiveness of hypnosis in alleviating pain associated with cataract surgery.

Methods

Patients

Male or female patients, aged 50–85 years, with bilateral agerelated cataract and who wished to have phacoemulsification surgery in both eyes were preliminarily admitted. Immediately after the first-eye surgery, each patient was evaluated for pain using the visual analog scale (VAS), and patients with a VAS score >1 were enrolled. Using block randomization, the enrolled patients were allocated to either the treatment group, which received hypnosis intervention before the scheduled second-eye surgery, or the control group, which did not undergo hypnosis.

Treatment

Patients in the treatment group were managed by a hypnotherapist half an hour before the second-eye surgery. After explaining the hypnosis procedure, the hypnotherapist initiated a 10- to 15-minute short hypnosis session using a technique described by Erickson;15 after the hypnosis session, the patient's anxiety for the forthcoming surgery was evaluated, and a standard phacoemulsification surgery under topical anesthesia was performed by the same surgeon. The topical anesthesia used was 0.5% proparacaine hydrochloride drops (Alcaine, Alcon Pharma; Puurs, Belgium) which was applied to the ocular surface twice at 2- to 3-minute intervals. In cases where patients experienced unbearable pain and were uncooperative, an additional dose of 2% lidocaine was administered by sub-Tenon's injection. The patients in the control group underwent the same procedures except for the hypnosis session. Patients who had any of the following were excluded from the study: complicated surgeries, such as zonular dialysis or posterior capsule rupture; a history of previous eye surgery; evidence of injury or inflammation; nystagmus; communication or cooperation difficulties; any prior experience of using hypnosis; conditions of psychosis, such as hallucinations, delusions, or depression; or excessive smoking and drinking habit. The surgeon was blinded to the group allocations until the entire study was completed. Upon completion of the surgery, the patients received oral instructions to guide their return to consciousness, and pain and cooperation levels were subsequently evaluated by the patients and the surgeon separately.

Patient pain was evaluated after the surgery using a VAS ruler as follows: 0 point, no pain; 1–3 points, mild pain; and >3 points, moderate-to-severe pain (Figure 1).

Patient cooperation was graded from 0 (no event) to 3 (obvious eye and head movement and/or lid squeezing) by the surgeon after the surgery. The grade was classified as follows: 0, excellent cooperation (no events); 1, good cooperation (eyelid squeezing); 2, sufficient cooperation (globe movement and eyelid squeezing); 3, poor cooperation (head movement, globe movement, and eyelid squeezing).

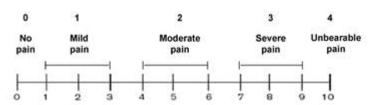


Figure I Pain score chart. 0, no pain; I-3, mild pain; 4-6, moderate pain; 7-9, severe pain; 10, unbearable pain.

Patient anxiety was evaluated before surgery using the Amsterdam Preoperative Anxiety and Information Scale (APAIS). The test involved a six-item questionnaire. Four items in the questionnaire were used to measure preoperative anxiety, and two items were used to measure the need for information. The items were rated on a 5-point Likert scale from "not at all" to "extremely". Each question was scored from 1 to 5. The total anxiety scale score ranged from 4 to 20. The information demand scale score ranged from 2 to 10.

This study was approved by the Third Affiliated Hospital Institutional Review Board, Third Military Medical University, Chongqing, China, and all patients provided written informed consent. The clinical trial registration number is ChiCTR1800014711.

Statistical analysis

Quantitative data were compared between the two groups via t-test (normal distribution) and the Wilcoxon rank sum test (non-normal distribution). The χ^2 test was used to compare categorical variables. The Spearman rank correlation was used for nonparametric data. Statistical analysis was performed using SPSS 13.0 (SPSS Inc., Chicago, IL, USA). The sample size for the study was based on the results from an earlier study; two-tailed t-tests were conducted with a=0.05 and β =0.1, and it was determined that at least 50 patients were required for each group. Taking patient discontinuation into consideration, as well as possible complications in the surgery, a total of 120 patients were chosen to ensure sufficient enrollment.

Results

A total of 120 patients were enrolled and formed the intent-to-treat population, with 60 patients randomized to each study group. Among the enrolled patients, three in the treatment group and two in the control group did not undergo their second-eye surgery for health or other reasons. In addition, two patients from each group were excluded from the study because complications occurred and lidocaine was used in the surgery. Hence, 111 patients (48 men and 53 women, 55 in the treatment group and 56 in the control group) between 50 and 82 years (median 64 years) were included in the final evaluation. Baseline characteristics were well balanced between the two groups (Table 1 and Figure 2).

During the intraoperative pain assessment using the VAS, 5%, 34%, 38%, and 23% of patients in the control group reported having no pain, mild pain, moderate pain, and severe pain, respectively. In contrast, in the hypnosis group, 18%, 56%, 15%, and 11% of patients reported having no pain, mild

Table I Patient characteristics

Parameter	Hypnosis	Control	P-value
Patients (n)	55	56	_
Mean age (years) \pm SD	64.11±5.74	64.66±5.07	0.59
Sex (male/female)	25/30	27/29	0.85

pain, moderate pain, and severe pain, respectively, which showed significant differences between the groups (P<0.005) (Table 2 and Figure 3).

Patient anxiety levels were evaluated using the APAIS. A highly significant difference was noted between the control group and the hypnosis group (P<0.005). The mean score of the control group was 11.77 (SD 0.32), and the mean score of the hypnosis group was 6.64 (SD 0.21). Results from the patient anxiety assessments are presented in Table 3 and Figure 4.

The cooperation assessment revealed that only 5% and 18% of patients in the control group exhibited excellent and good cooperation, respectively, whereas 47% of patients exhibited poor cooperation. In contrast, in the hypnosis group, 18% and 36% of patients exhibited excellent and good cooperation, respectively, whereas 24% exhibited poor cooperation, which showed significant differences among the four groups (P<0.005). The results of the patient cooperation assessments are presented in Table 4 and Figure 5.

There was a significant positive correlation between the VAS and cooperation, both in the hypnosis group and in the control group (r=0.414, P<0.005; r=0.596, P<0.005). However, there was no significant correlation between the VAS and the anxiety analysis in either group.

Discussion

In conventional cataract surgery, a skilled doctor can perform a successful phacoemulsification surgery in approximately 10 minutes, and topical anesthesia is thus often used to save time; however, topical anesthesia is not always completely effective for relieving pain and anxiety in phacoemulsification surgery. Most recent studies have focused on determining whether the pain experienced during the second-eye surgery is greater than or equal to that experienced during the first-eye surgery, ^{16–19} but few researchers have focused on reducing patient pain during the cataract surgery under topical anesthesia. In fact, if the patient experienced pain during the first-eye surgery, particularly obvious pain, he/she will inevitably experience pain and anxiety during the second-eye surgery if only local anesthesia is used without any auxiliary sedative measures.

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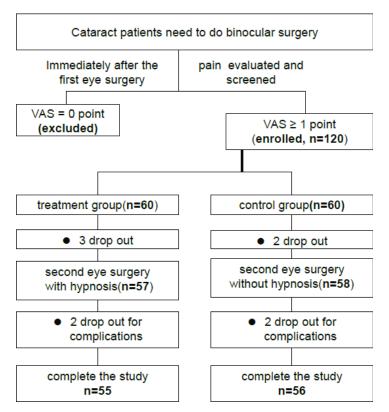


Figure 2 Study flow chart.

Abbreviation: VAS, visual analog scale.

Table 2 Patient VAS scores

Groups	n	VAS during	ohacoemulsificatio	χ²	P-value		
		0	I - 2	2–3	3–4		
Hypnosis	55	10 (18%)	31 (56%)	8 (15%)	6 (11%)	15.557	0.001
Control	56	3 (5%)	19 (34%)	21 (38%)	13 (23%)		

Abbreviation: VAS, visual analog scale.

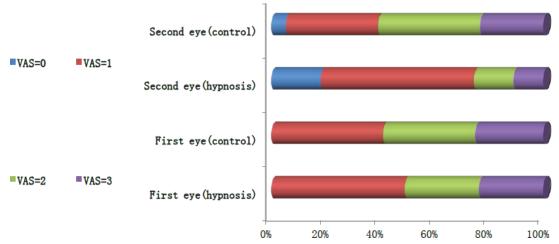


Figure 3 Extent of patient pain with or without hypnosis. **Abbreviation:** VAS, visual analog scale.

Hypnosis has been used in ophthalmic surgery for many years, and several cases of successful hypnosis application have been reported.^{20,21} However, these reports mainly focused on the ability of hypnosis to reduce medication use and increase patient satisfaction or described its use in high-risk patients in unusual situations.^{22–24} In the present study, we assessed whether hypnosis can assist in alleviating cataract patient pain and anxiety and improve cooperation in the second-eye surgery. In total, our results showed that patients in the hypnosis group exhibited remarkably improved results, though hypnosis combined with topical anesthesia was not able to eliminate pain for all patients, which may be reasonable given individual differences in the hypnotic effect and pain sensitivity. Overall, many patients in the hypnosis group presented with reduced pain, lower anxiety, and increased cooperation.

Our study revealed two main conclusions. First, with hypnosis therapy, patients with a higher VAS score in their first-eye surgery showed a remarkable decrease in VAS score in their second-eye surgery (data not shown). These

Table 3 Patient anxiety evaluation

Groups	n	Anxiety before the phacoemulsification surgery (mean ± SD)	t	P-value
Hypnosis	55	6.64±0.21	13.21	<0.0001
Control	56	11.77±0.32		

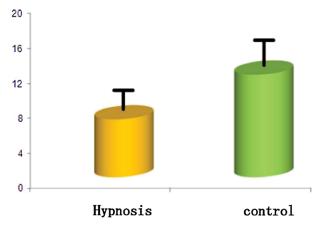


Figure 4 Anxiety levels in patients with or without hypnosis.

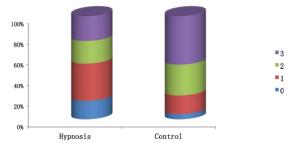


Figure 5 Cooperation levels in patients with or without hypnosis.

data suggest that hypnosis was able to help alleviate pain, especially for patients who experienced obvious pain during the first-eye surgery. Second, patients VAS score was positively associated with cooperation score, indicating that patients who experience less pain demonstrate better cooperation.

Conclusion

To the best of our knowledge, this is the first study investigating the effects of hypnosis on the management of pain during cataract surgery in the second eye. Although our data must be interpreted with caution given the high variability in hypnosis techniques, the lack of standard scales to indicate hypnotic susceptibility, individual differences to hypnotic susceptibility, and the small number of patients included in the study, our results clearly demonstrated that hypnosis may be considered as an auxiliary measure for cataract surgery in the second eye, especially for patients who experienced pain during the first-eye surgery.

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Author contributions

Chunlin Chen and Jian Ye designed the study and conducted the analyses. Xiaofan Chen, Rongdi Yuan, Xiao Chen, and Min Sun performed the experiments. Xiaofan Chen and Sen Lin wrote the manuscript. All authors read and approved the final manuscript.

Table 4 Patient cooperation scores

Groups	n	Cooperation during phacoemulsification surgery, n (%)				χ²	P-value
		0	I	2	3		
Hypnosis	55	10 (18%)	20 (36%)	12 (22%)	13 (24%)	12.944	0.005
Control	56	3 (5%)	10 (18%)	16 (30%)	27 (47%)		

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Disclosure

The authors report no conflicts of interest in this work.

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