




Survey of Intravitreal Injection Techniques Among Retina Specialists in Mexico

This article was published in the following Dove Press journal:
Clinical Ophthalmology

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Purpose: To report technique preferences for intravitreal injections among retina specialists in Mexico.

Methods: Cross-sectional survey. Ophthalmologists with a two-year retina training, active members of the Mexican Retina Association, were contacted through email to answer a survey consisting of 37 items regarding their IVI application technique.

Results: A total of 133 retina specialists participated, with a response rate of 78%. Forty-five percent applied the intravitreal injections in an operating room designated for the procedure. Sixty-three percent reported never injecting both eyes on the same day. Ninety-six percent wore a face mask during the procedure and 91% wore gloves. Eighty-two percent used a lid speculum. Tetracaine drops were the anesthetic method employed by 97% of participants. All participants utilized povidone-iodine for antisepsis. Eighty percent measured the puncture site with a caliper. Superotemporal quadrant was the one chosen to place the injection by 63% of participants. Fifty-nine percent indicated post-injection antibiotic drops for several days. Post-injection counting fingers visual acuity was verified by 53% of the participants. Fifty-six percent of the participants placed an eye-patch after the procedure.

Conclusion: There are different practices regarding the application of intravitreal injections among retina specialists in Mexico. Performing this type of survey periodically could show changes in preferences, as new evidence is incorporated into clinical practice.

Keywords: intravitreal injections, retinal diseases, anti-VEGF agents, topical antibiotics

Introduction

Currently, intravitreal injections (IVI) are accepted as the standard form of drug administration for a large number of retinal diseases. Despite the frequency with which they are applied and the numerous guidelines that have been published in this regard,¹⁻³ issues such as drug preparation, injection technique, indications, follow-up and office vs operating room setting may vary according to surgeon preference and/or country. These practices are always aimed at preventing endophthalmitis and other complications such as retinal detachment, cataract formation and increased intraocular pressure,^{4,5} as well as reducing patient discomfort during the procedure.

Previous studies have reported the preferences for IVI application among retina specialists from countries like the USA, Canada, Israel, India and Brazil.⁶⁻¹¹ To the best of our knowledge, there are no reports about these preferences in Mexico.

The aim of this study was to report the variety of practices regarding IVI application technique and pre/post-injection procedures performed by retina specialists in Mexico, members of the Mexican Retina Association, who underwent a formal two-year retina subspecialty training.

Materials and Methods

All active members of the Mexican Retina Association (n=170) were invited to participate in an anonymous survey sent by e-mail on March 2nd, 2020. The questionnaire was based on previously published surveys^{6–11} plus other items that were considered relevant to our practice. It consisted of 37 questions written by the authors, regarding the practices before, during and after the application of IVI and was distributed using Google Forms (Google Inc. Mountain View, CA, USA). Those who did not answer the first email received a second email during May 2020 and some of them were reached through phone call. The final results were gathered on June 3rd, 2020 for analysis. This study adhered to the tenets of the Declaration of Helsinki, and the ethic committee approval was unnecessary as the anonymous questionnaire was conducted via emails.

Results

A total of 133 (78%) out of 170 retina specialists participated in the survey. Regarding the time elapsed since they completed their specialty training, 19.5% (26/133) had finished it in a period between 1 and 5 years, 22.6% (30/133) between 6 and 10 years, 20.3% (27/133) between 11 and 15 years, 16.5% (22/133) between 16 and 20 years and 21.1% (28/133) more than 20 years ago. Forty-four percent (59/133) had an established medical practice in Mexico City and its metropolitan area and the remaining 55.6% (74/133) in the rest of the country.

About the weekly physician volume of IVI, 45.1% (60/133) referred to apply between 5 and 10 injections, 36.1% (48/133) less than 5 injections, 12.8% (17/133) between 10 and 20 and 6% (8/133) more than 20 injections. Among anti-VEGF (vascular endothelial growth factor) agents, Ziv-Aflibercept (Zaltrapziv, Sanofi-Aventis) was applied by 40.6% (54/133), being the most frequently used, followed by Aflibercept (Wetlia, Bayer) used by 35.3% (47/133), then Ranibizumab (Lucentis, Novartis) used by 13.5% (18/133) and Bevacizumab (Avastin, Roche) applied by 10.5% (14/133).

More than half of the participants (58.6% [78/133]) requested the patient to sign an informed consent form for each injection, while 31.6% (42/133) asked it only for the first injection and 9.8% (13/133) did not request it at all.

Regarding the setting where the IVI were applied, 45.1% (60/133) mentioned performing them in a room

specially dedicated for the procedure, 33.8% (45/133) in the operating room (69% due to hygiene measures in the operating vs the office and 31% by request of the medical insurer) and 21.1% (28/133) in their medical office.

About the application of IVI in both eyes, 63.2% (84/133) referred never inject both eyes on the same day, 24.8% (33/133) inject both eyes on the same day only under special circumstances and 12% (16/133) inject always both eyes on the same day. Among the participants who referred to inject both eyes on the same day, 57% (28/49) referred to employ different vials and batches of medication for each eye and 43% (21/49) used the same vial for both eyes.

Pre-injection antibiotic drops were prescribed by 21.8% (29/133) of participants. Seventy-six percent (22/29) used quinolones and 20.7% (6/29) aminoglycosides, of which 17.2% (5/29) reported using any of these antibiotics in combination with a steroid. Three percent (1/29) did not specify the antibiotic they prescribed.

Pre-Injection Procedures

To obtain the dose of the drug to be injected into the vitreous cavity: 47.4% (63/133) used aliquots that were previously divided from a vial, 39.8% (53/133) used a vial from which they took multiple doses to use in different patients, 12.8% (17/133) took the dose from the vial and injected it, using only one vial per eye.

Forty-eight percent (46/96) cleaned the vial cap with povidone-iodine, 38.5% (37/96) with alcohol, while 13.5% (13/96) reported not cleaning the vial cap.

To inject the medication, 16% (16/100) utilized the same needle for drawing the medication from the vial and applying the IVI, while 84% (84/100) used a needle to draw the medication and a different one to perform the injection.

Regarding the use of a face mask, 96.2% (128/133) of the participants reported to wear a face mask during the procedure. Among them, 54% (69/128) reported that the face mask was only worn by the physician, while in the remaining 46% (59/128) the face mask was worn by both the physician and the patient. In 3% (4/133) of cases, masks were not worn by neither the physician nor the patient and in 0.75% (1/133) the patient was the only one wearing the face mask. Eighty-six percent of participants asked the patient not to talk during the procedure, despite the use of a face mask.

Ninety-one percent (121/133) of participants referred the use of gloves for the IVI (97% [117/121] sterile gloves and 3% [4/121] nonsterile gloves). A lid speculum was

used during the IVI by 82.7% (110/133) of the participants. Regarding the position of the patient during the procedure, 48.9% (65/133) preferred a supine position, while 36.1% (48/133) used a semi-sitting position and 15% (20/133) a sitting position.

For anesthesia, 97% (129/133) of participants used tetracaine drops, while 2.3% (3/133) utilized sub-conjunctival lidocaine and only 0.8% (1/133) applied lidocaine as a gel. Considering all anesthetic options, 63.9% (85/133) referred to administrate the medication from 5 to 10 minutes before the procedure, 26.3% (35/133) right before the injection and 9.8% (13/133) more than 15 minutes before.

All participants performed antisepsis using povidone-iodine before the IVI. Seventy-four percent (99/133) used the povidone-iodine at a concentration of 5%, 13.5% (18/133) at a concentration of 10% and 12% (16/133) at a concentration below 5%. Regarding the area where the povidone-iodine was applied, 67% (89/133) referred to apply it on periocular skin, eyelids and the conjunctival sac, 24% (32/133) only on eyelid margins and the conjunctival sac and 9% (12/133) only on the conjunctival sac.

IVI Technique

Eighty percent (107/133) measured the distance from the corneal limbus to the site of the injection. In order to obtain the measure, 84% (90/107) used a caliper and 15.9% (17/107) the needle cap. Regarding the quadrant to place the injection, 63.2% (84/133) preferred to perform it in the superotemporal quadrant, followed by 34.6% (46/133) in the inferotemporal and 2.3% (3/133) in the superonasal.

Concerning the gauge of the needle, 43.6% (58/133) of participants used 30 G, 28.6% (38/133) 31 G, 15% (20/133) 27 G, 9.8% (13/133) 32 G and 3% (4/133) 25 G. After the injection, 80.5% (107/133) made pressure with a cotton swab to avoid vitreous leaking.

Post-IVI Procedures

Twenty-three percent of participants (30/133) did not prescribe antibiotic drops after the injection. Fifty-nine percent (79/133) recommended antibiotic drops for several days after the injection and 18% (24/133) applied only one dose immediately after the injection. Among those who used post-injection antibiotics drops: 70.9% (73/103) utilized quinolones, 17.5% (18/103) aminoglycosides

and 19.4% (20/133) of them referred to use antibiotics combined with steroids.

Seventy-four percent (99/133) of participants did not utilize a hypotensive agent before or after the IVI, 18% (24/133) applied this medication only under special circumstances (ocular hypertension/glaucoma), while 7.5% (10/133) reported to use it on every procedure.

Fifty-three percent (70/133) affirmed to check counting fingers visual acuity right after the injection. Only 2.3% (3/133) referred to perform a paracentesis on a regular basis after every procedure. Fifty-six percent (74/133) placed an eye-patch after the injection.

Regarding follow-up examinations after IVI, 38.3% (51/133) of participants performed an examination at their office 1–5 days after the procedure, 36.8% (49/133) did not examine the patient until the appointment for the next IVI, 12.8% (17/133) performed an examination between days 7–15 post-IVI and 12% (16/133) did not perform an examination, but they did a follow-up phone call to the patient a few days after the IVI.

Discussion

This survey shows the diversity that exists in the different phases of the application of intravitreal injections among retina specialists in Mexico, and that in many aspects these practices differ from what has been previously reported in other countries.

Regarding the practice of applying bilateral injections on the same day, our study reported less bilateral injections compared to other countries such as the USA, where the frequency of this practice ranged between 45% and 71.5%.^{6,7} Other countries with a higher frequency were Canada (57%), Israel (56%), India (46%) and Brazil (36%)^{8–11} (Table 1).

Previous studies have reported that bilateral injections performed on the same day are well tolerated and can save time and money to the patients, without increasing the risk of endophthalmitis¹² or other systemic adverse events.^{13,14} The WHO, along with the Royal College of Ophthalmologists from the United Kingdom, suggest that the second injection should be considered as a totally new procedure, independent from the first one. This means that it requires the use of a new set of sterile gloves, a fresh preparation of povidone-iodine for antisepsis, a new and sterile caliper, lid speculum and surgical sheets in order to reduce the risk of infection or cross-contamination.^{15,16}

Most of the participants referred to apply the IVI in a room specially dedicated for this procedure. Data from

Table 1 Comparison of Intravitreal Injection Practice Patterns Between This Study and Previous Studies

	Canada 2014 Xing et al⁸	USA & others 2019 Uhr et al⁷	USA 2019 Chaturvedi et al⁶	Israel 2016 Segal et al⁹	India 2020 Sundar et al¹⁰	Brazil 2015 Shiroma et al¹¹	Europe 2016 Huang et al²⁰	Mexico 2020 Henaine- Berra et al (current study)
Number of participants (Survey response rate)	75 (64%)	399 (14%)	281 (17%)	52 (66%)	741 (73%)	352 (38%)	113 (90%)	133 (78%)
Pre-IVI procedures								
Setting								
Physician's office	72%	72%	N/A	N/A		5%	3.5%	21%
Procedure room	N/A	23%			8%	7%		45%
Operating room	28%	5.5%			91%	88%	68%	34%
Simultaneous bilateral injections	57%	71.5%	45%	56%	N/A	36%	N/A	12%
Pre-IVI antibiotics	20%	11%	21%	N/A	60%	N/A	67%	22%
Face mask	29%	33%	36%	37%	98%	N/A	N/A	97%
Gloves	39%	55%	73%	79%	98%	95%	75%	91%
Lid speculum	91%	72%	66%	98%	96%	95%	81%	83%
Anesthesia	Drops combined with other methods: 90% Drops only: 29% Gel: 25% Swab: 23% SC: 23%	Drops: 63% Gel: 32% SC: 27% Swab: 19% Others: 3%	Gel with or without drops: 31% SC with or without drops: 27% Drops only: 16% Drops + swab: 14%	Drops + gel: 67% Drops only: 25% Gel only: 8%	Local anesthesia: 91%	Drops only: 67% Drops + gel: 20% Peribulbar: 3%	N/A	Drops: 97% SC: 2% Gel: 1%
Antisepsis	PI: 100%	PI: 97.5% Clorhexidine: 2%	PI: 100%	N/A	N/A	PI: 99%	PI: 90%	PI: 100%
Injection technique								
Quadrant								
Superotemporal	43%	OD 32% OS 30%	27%	33%	N/A	N/A	N/A	63%
Inferotemporal	63%	OD 62% OS 61%	70%	17%	N/A	N/A	N/A	35%
Superonasal	7%	OD 1.5% OS 4%	0%	6%	N/A	N/A	N/A	2%

(Continued)

Table I (Continued).

	Canada 2014 Xing et al ⁸	USA & others 2019 Uhr et al ⁷	USA 2019 Chaturvedi et al ⁶	Israel 2016 Segal et al ⁹	India 2020 Sundar et al ¹⁰	Brazil 2015 Shiroma et al ¹¹	Europe 2016 Huang et al ²⁰	Mexico 2020 Henaine- Berra et al (current study)
Inferonasal	7%	OD 5% OS 5%	4%	4%	N/A	N/A	N/A	0%
Measurement of distance from corneal limbus to site of puncture	56%	30.5%	20%	69%	N/A	80%	N/A	80%
Needle gauge for intraocular injection								
25 G	N/A	N/A	NE	0	N/A	N/A	N/A	3%
27 G	N/A	N/A	1%	4%	N/A	N/A	N/A	15%
30 G or more	N/A	N/A	89%	90%	N/A	48%	N/A	82%
Method for preventing vitreous reflux								
Swab	41%	N/A	37%	N/A	N/A	N/A	N/A	80.5%
Angled scleral tunnel technique	7%	N/A	N/A	N/A	N/A	30%	N/A	N/A
Conjunctival displacement	11%	20.1%	N/A	21%	N/A	58%	N/A	N/A
None	40%	N/A	N/A	N/A	N/A	N/A	N/A	19.5%
Post-IVI procedures								
Use of antibiotics post-IVI	20%	17% Always or often instills antibiotics immediately after injection, 9% Always or often prescribes antibiotics for home use	28%	92%	89% for several days	89% for several days	89.5%	59% for several days, 18% one drop immediately after injection
AC paracentesis (pre or post injection)	51%	14% in patients with advanced glaucoma	10% performed 25% of the time	27% in cases of end- stage glaucoma	N/A	Always performed: 11% Performed when necessary: 54%	N/A	2%
Verify optic nerve perfusion	48%	56%	4% checked for visual acuity	44%	N/A	25%	N/A	53% checked counting fingers visual acuity

Abbreviations: N/A, not available; PI, povidone-iodine; AB, antibiotics; SC, sub-conjunctival, AC, anterior chamber.

other countries reveal relevant differences about the place for IVI application. In Canada, 72% used their office, on the other hand, in India, 91.4% performed the procedure in the operating room¹³ (Table 1).

Previous publications have shown an incidence of endophthalmitis between 0.029% and 0.09%^{10,13} when the IVI is performed at the physician office. These data suggest that this practice does not have an increased risk compared to the procedures performed in the operating room. A meta-analysis revealed that the incidence of endophthalmitis, when the IVI were performed in the operating room, was 0.07% compared to the 0.04% when the injection was applied at the physician's office. It is worth to mention that when the procedures were performed at the physician's office, the physicians did not use gloves, surgical sheets or face mask, but did use a lid speculum and povidone-iodine.¹⁷

All locations mentioned (operating room, procedure room or office) are safe and reliable to perform the IVI.¹⁸ The particular choice of setting could be related to different factors such as availability, cost and logistics within the different health systems.

Use of Antibiotics

Twenty-two percent of our participants confirmed the use of pre-IVI antibiotic drops. This percentage is higher compared to data from the USA, where only 10.9% of physicians prescribed them. However, in the United Kingdom and other European countries, the proportions were higher with 30% and 66.7%, respectively.^{19,20}

More than half of our participants (59.4%) prescribed post-IVI antibiotics for several days, while in the USA they were prescribed only by 9.1%² and in the United Kingdom by 74%.¹⁹

Our study revealed that 18% prescribed only one dose of antibiotic drops immediately after the IVI. This percentage is similar to that observed in the USA (16.6%).⁷ In the United Kingdom and Europe, this practice was carried out by 90% and 89.5%, respectively.²⁰

Surveys from Canada and Asia-Pacific showed that prophylactic antibiotic drops were applied by 43% and 84.9 of participants, respectively.^{8,21} Regarding post-IVI antibiotics, Brazil (89.21%),¹¹ India (89.3%)¹⁰ and the United Kingdom (74%)¹⁹ reported a higher frequency compared to ours.

Quinolones were the preferred antibiotics used by participants and the same behavior was observed in retina specialists from Canada.⁸ A previous study showed that

in 87.5% of cases the conjunctival flora was resistant to quinolones after 4 days of treatment.²²

Currently, there is sufficient evidence to support the fact that the use of topical antibiotics is not associated with a reduction in the incidence of endophthalmitis in the context of intravitreal injections,^{23–26} both post-injection^{27–29} and pre-injection,^{23,30} and it can contribute to microbial antibiotic resistance.^{22,31–34} The use of antibiotics before IVI plus the application of povidone-iodine does not confer an advantage over the use of povidone-iodine alone.^{23,24,30,35} Several studies and clinical practice guidelines affirm that the use of antibiotics before or after the IVI is unnecessary and reinforce that a proper antisepsis technique is the best prophylactic practice to avoid endophthalmitis.^{36–38} Furthermore, a higher risk of endophthalmitis was found in the group that used antibiotics in some studies,^{27,28,39,40} suggesting that repeated exposure to antibiotics may promote the development of resistant strains and this may lead to this paradoxical finding.^{2,22} Accumulated evidence suggests that the use of antibiotics before or after IVI does not reduce the risk of infection and it must not be used routinely.^{23,27,29}

Pre-IVI Procedures

In our study, when off-label medications were used, several doses for different patients were taken from the same vial. When the medication was approved for intra-ocular application, the dose was directly taken from the vial and one vial was used for only one patient, or alternatively aliquots were previously prepared and stored under sterile conditions. Surveys carried out in other countries did not mention if the vial cap was cleaned before taking the dose.

There were no previous reports mentioning if physicians use one needle for taking the medication and a new needle for applying the IVI. General practice to avoiding infections suggests using a new, sterile needle for intravitreal medication delivery, different from the one used to draw the medication.⁴¹

Regarding the use of face mask, data obtained from surveys around the world have evidenced that the use of face mask is a very variable practice, ranging from 37% in Israel to 98% in India.^{9,10}

Several retrospective studies have demonstrated a positive correlation between the use of face mask plus the habit of not talking during IVI with a reduced risk of developing endophthalmitis, based on the reduction of potential contamination from the oropharyngeal flora.^{18,42–44} In a study published in 2011, in the setting of a simulated intravitreal injection, significantly greater

bacterial dispersion was demonstrated when speaking without a mask than when speaking using a mask or remaining silent.⁴⁵ It is worth to mention that this survey was carried out before the COVID-19 pandemic, so it is probable that the percentages of physicians and patients wearing a facemask has changed radically ever since.

There are no publications demonstrating that the use of gloves (sterile or nonsterile) correlated with a reduced risk of endophthalmitis.¹⁸ Nevertheless, some clinical practice guidelines recommend the use of gloves for the prevention of infections when working at the physician's office.¹ Surveys from the USA, Canada and Brazil reported the use of gloves by 55%, 39% and 95% of physicians, respectively.^{7,8,11}

In a survey from Israel, 79% of participants affirmed to use gloves (95% of them used sterile gloves),⁹ whereas in India 98% of physicians used sterile gloves and a face mask during the procedure.¹⁰ On the other hand, 71% of the retina specialists in Israel utilized sterile surgical sheets and only 37% wore a face mask.⁹

Regarding the use of a lid speculum, surveys have shown a percentage of use of 98%, 96% and 66% in Israel, India and the USA, respectively.^{6,9,10} Recently, the use of the lid speculum has been replaced by manual retraction. Uhr and cols reported that the use of lid speculum dropped from 92% in 2011 to 72.3% in 2019.^{7,46} In addition, Fineman and cols⁴⁷ have demonstrated that assisted manual retraction is a safe technique. Their retrospective study found an incidence of endophthalmitis of 0.03%, similar to the incidence associated to the use of lid speculum.

Anesthesia

The use of ophthalmic drops as anesthesia for the application of IVI was the preferred method in Canada (90%) and Brazil (67%).^{8,11}

Regarding the application of sub-conjunctival anesthesia, this method provides a very effective analgesia for the application of IVI. However, the injection necessary to apply lidocaine in the subconjunctival space is usually more painful than other available methods or the procedure itself. Furthermore, a previous study showed that this injection provoked a sub-conjunctival hemorrhage in more than 50% of the cases.⁴⁸ In some other surveys,^{7,8} sub-conjunctival anesthesia was mentioned as the preferred method for more than 20% of the participants.

The use of anesthesia as a gel is a less frequent practice among the retina specialists in our country. However, in

USA, Canada and Israel⁶⁻⁹ more than 25% of the participants referred the use of the gel as the election method, either alone or combined with ophthalmic drops. The use of gel anesthesia is controversial, since it has been reported that if the gel is applied before povidone-iodine, it may interfere with its antiseptic action and increase microbial survival.⁴⁹

Surveys from other countries do not mention the time elapsed between the application of the anesthesia and IVI. In our survey, 63.9% referred to apply the anesthesia 5–10 minutes before the procedure, 26.3% immediately before the injection and 9.8% 15 minutes or more before the procedure. The anesthetic effect of proparacaine drops starts 20 seconds after its application and last for 10–15 minutes, whereas the effect of tetracaine drops starts 1 minute after its application and last for 15–20 minutes.⁵⁰

Antisepsis

All of our participants referred to perform antisepsis with povidone-iodine before IVI. This behavior is similar in the USA, Canada and Brazil, where more than 99% of the physicians carry out antisepsis in the same way.^{6,8,11} Several publications have demonstrated that application of IVI without previous antisepsis importantly increases the risk of endophthalmitis.⁵¹⁻⁵³

Most of the participants (74%) used povidone-iodine at a concentration of 5%. It has been demonstrated that povidone-iodine at a 5% concentration takes 2 minutes to eliminate 100% of the micro-organisms of the surface where it is applied. In addition, at this concentration, it causes less irritation of the ocular surface compared to 10% povidone-iodine. Furthermore, povidone-iodine at 10% takes 4 minutes to eliminate 100% of the micro-organisms.⁵⁴ In conclusion, there is no advantage for using povidone-iodine at 10%.

Regarding the area of application of povidone-iodine, a survey among retina specialists from Canada performed in 2014 revealed that 48% applied it on both the periocular skin and the conjunctival cul-de-sac.⁸

Other alternatives for antisepsis found in different surveys include the use of chlorhexidine and antibiotics either alone or in combination with povidone-iodine.⁷

Injection Technique

Both in our survey and those carried out in Israel⁴ and Brazil,¹¹ more than 80% of participants referred to measure the exact distance from the corneal limbus to the site of injection using a caliper. On the other hand, retina

specialists from the USA revealed that more than 60% of them did not measure the distance from the corneal limbus to the spot of injection.^{6,7}

Regarding the quadrant where the injection is placed, surveys from Canada and the USA reported a preference for the inferotemporal quadrant in more than 60% of the participants⁶⁻⁸ (Table 1).

About the gauge of the needle used for the puncture of the eye globe, most of the participants in our survey used a 30G needle. Our observation agrees with data obtained from surveys performed in Israel and the USA, where a 30G needle was used by 90% and 61%, respectively.^{6,9}

The use of a cotton swab as a method to prevent vitreous reflux through the site of puncture was a very common practice among our participants. Leaking is prevented by applying pressure with the cotton swab immediately after the puncture. Only two of the surveys (USA and Canada)^{6,8} reported its use, with around 40% of physicians practicing it, which contrasts with 80% in our survey.

Post-IVI Procedures

Only 7.5% of our participants used hypotensive drops routinely. This percentage is similar in Brazil,¹¹ where it is used by 5% of retina specialists; in addition, 10% of Brazilian physicians prescribed oral acetazolamide. Eyes with no vitreous reflux at the time of injection have risk of suffering intraocular pressure elevation, which decreases without intervention in the following 30 minutes. Eyes with vitreous reflux present a reduction of the intraocular pressure which returns to normal in the following 10 minutes.⁵⁵

In our survey, 52.6% corroborated immediate post-injection counting fingers vision, which, if present, is associated with adequate optic nerve perfusion. This practice is also performed in the USA (21%-56%),^{6,7} Canada (48%)⁸ and Israel (44%)⁹ (Table 1). In Canada and Israel, specialists referred to perform pupil dilation for an adequate examination after the injection.

Regarding anterior chamber paracentesis, in the Canadian⁸ and Brazilian¹¹ surveys, 5% and 11% of their respondents performed paracentesis in all their cases.

The routinely use of an eye-patch after the procedure was employed by 77.3% of the physicians in India.¹⁰ The use of an eye-patch is not recommended on the guides for the management of IVI published in 2014 in the USA.¹ A study conducted in Hong Kong in 2015⁵⁶ reported the use of eye-patch routinely with antibiotic ointment and the patch was removed when the patients arrived home.

The risk of adverse events related to IVI is relatively low. Therefore, there is no consensus regarding the examination after the procedure. In countries like India¹⁰ and Brazil¹¹, a follow-up examination at the office was performed within the first 2 weeks after injection by the 85% and 79%, respectively.

Conclusion

Our study shows that there are different practices regarding the application of intravitreal injections among retina specialists in Mexico. The response rate of 78% was high, compared to surveys conducted in other countries. The limitation, as in any survey, was the bias in the selection of participants, in this case being answered only by active members of the Mexican Retina Association. Performing this type of survey periodically could show changes in preferences, as new evidence is incorporated into clinical practice.

Abbreviations

IVI, intravitreal injections.

Data Sharing Statement

The datasets used and/or analysed during the current study are available from the corresponding author on reasonable request.

Consent for Publication

Not applicable.

Ethics Approval and Informed Consent

Not applicable.

Author Contributions

All authors contributed to data analysis, drafting or revising the article, have agreed on the journal to which the article will be submitted, gave final approval of the version to be published, and agree to be accountable for all aspects of the work.

Funding

This study is partially funded by the Mexican Retina Association.

Disclosure

Gerardo Garcia-Aguirre reports personal fees from Alcon, Allergan, Bayer, Liomont, Novartis, Roche/Genentech,

and Zeiss, outside the submitted work. The authors declare that they have no other potential conflicts of interest for this work.

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