Combination of ranibizumab and indomethacin for neovascular age-related macular degeneration: randomized controlled trial

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Purpose: The aim of this study was to evaluate whether indomethacin eye drops and intravitreal ranibizumab (IVR) injections would provide additional benefit over ranibizumab alone in the treatment of choroidal neovascularization (CNV).

Participants and methods: This was a randomized, prospective pilot study of eyes with new-onset CNV. Fifty-eight patients were randomized 1:1 into a ranibizumab monotherapy (RM) group and a ranibizumab plus indomethacin (RI) group. All patients received monthly 0.5 mg IVR injections for 3 months, followed by monthly injections administered as needed. RI group patients also self-administered one drop of 0.5% indomethacin three times a day for 12 months. All patients were followed up for 12 months.

Results: At 12 months, both groups showed significant improvement in best-corrected visual acuity (BCVA) and central retinal thickness (CRT). The mean BCVA change from baseline to 12 months was −0.12±0.04 LogMAR and −0.20±0.04 LogMAR in the RM and RI groups, respectively, with the degree of change being significantly different between the two groups (P=0.04). At 12 months, the mean CRT in the RM group (316±41.2 μm) was significantly higher than that in the RI group (287±31.5 μm; P=0.004). The mean required number of IVR injections was 7.38±0.78 and 6.34±0.67 in the RM and RI groups, respectively (P<0.001).

Conclusion: Compared to IVR monotherapy, combination therapy with indomethacin eye drops and IVR provides superior anatomical and visual outcomes in patients with naive CNV lesions. Moreover, topical indomethacin might reduce the frequency of IVR injections, which is very beneficial considering the chronic and expensive nature of IVR therapy.

Keywords: central retinal thickness, choroidal neovascularization, indomethacin, inflammation, ranibizumab

Introduction

Age-related macular degeneration (AMD) is still the primary cause of visual impairment and blindness in patients older than 60 years in developed countries.1 In the neovascular form of AMD, the choroidal neovascularization (CNV) beneath the macula leads to fibrous metaplasia, permanent loss of photoreceptors, and disciform scarring, which often results in the loss of central vision.2

Large-scale clinical trials have shown that monthly or bimonthly intravitreal injection of vascular endothelial growth factor (VEGF) antagonists prevents vision loss and may even improve visual acuity in patients with neovascular AMD.3,4 However, it is important to note that VEGF is not the only causative factor in CNV. In particular, free radicals and oxidized lipoproteins in the aging retina are major local triggers of parainflammation, which is the chronic status responsible for the initiation and progression of age-related
chorioretinal damage.\textsuperscript{5,6} Thus, inflammation itself plays an important role in the pathogenesis of CNV and administration of a topical nonsteroidal anti-inflammatory drug (NSAID) has been shown to supplement the effectiveness of anti-VEGF agents in reducing central retinal thickness (CRT)\textsuperscript{7,8} and the rate of re-injection in CNV.\textsuperscript{9} A recent assessment of the vitreous penetration of NSAIDs and their effects on prostaglandin E\textsubscript{2} (PGE\textsubscript{2}) revealed a significant reduction in vitreous PGE\textsubscript{2} levels in response to indomethacin, bromfenac, and nepafenac.\textsuperscript{10} Considering these results, in this study, we prospectively evaluated whether the addition of 0.5\% indomethacin eye drops three times a day (TID) to the standard treatment of intravitreal ranibizumab (IVR) injections would provide better efficacy than that achieved with ranibizumab alone in patients with CNV.

**Participants and methods**

**Study design**

This was an open-label, pilot study in eyes affected by new neovascular AMD. This study was conducted in the Ophthalmic Center of “Spedali Civili di Brescia”, according to the ethical principles of the Declaration of Helsinki. The institutional review board of the “Spedali Civili di Brescia” hospital approved the study protocol (registered with ClinicalTrials.gov, identifier NCT03261635). All study participants provided written informed consent.

**Participants**

Sixty consecutive participants were enrolled from September 2016 to March 2017 at the Spedali Civili Ophthalmic Center (Spedali Civili di Brescia) in Brescia, Italy. Patients were randomized (online statistical computing web programming www.graphpad.com/quickcalc) at a 1:1 ratio into the following two groups: monotherapy with IVR injections (ranibizumab monotherapy [RM] group; n=29) and IVR injections plus off-label topical 0.5\% indomethacin eye drops (ranibizumab plus indomethacin [RI] group; n=29). One patient per group was excluded because they moved to another eye clinic after the initial ranibizumab-loading phase.

**Inclusion and exclusion criteria**

Inclusion criteria were as follows: 1) provision of written informed consent, 2) age >40 years, and 3) presence of treatment-naive neovascular AMD, which was defined as neovascularization, fluid, or hemorrhage underneath the fovea. The presence of new active CNV was confirmed by the evidence of leakage on fluorescein angiography and fluid on spectral domain optical coherence tomography (OCT) (Spectralis OCT; Heidelberg Engineering, Heidelberg, Germany). The diagnosis was made by one investigator (AR) and thereafter confirmed by the second investigator (EG).

Exclusion criteria were as follows: 1) any prior intravitreal or retinal laser treatment, 2) pathological myopia (>7 D), 3) any concomitant eye disease, and 4) corneal epithelial defects or any condition that would affect the cornea.

**Study treatments**

Participants in both groups received an initial loading phase of 3 monthly IVR. Retreatment criteria for further injections performed by a masked examiner were as follows: 1) any intraretinal or subretinal fluid on OCT, 2) new or persistent hemorrhage, and 3) decrease in visual acuity. In the case of the absence of fluid on OCT or visual acuity drop, a fluorescein leakage of >25\% of the lesion circumference or the expansion of CNV was needed for retreatment in suspicious cases.

Patients in the RI group were given a bottle of eye drops of 0.5\% indomethacin ophthalmic solution for self-administration, which was provided free of charge by Alfa Intes Srl (Casoria, Italy). Over the 12-month study period, indomethacin was administered at a dose of one drop in the study eye TID. To verify and increase their compliance, patients were asked to bring their used bottles of indomethacin during each visit.

**Clinical evaluation**

The following examinations were performed at each visit by a blinded examiner: 1) measurement of best-corrected visual acuity (BCVA) as measured by the Early Treatment Diabetic Retinopathy study letter score, 2) fully dilated slit-lamp ophthalmic examination, 3) OCT (Spectral Domain Spectralis OCT) measurement of CRT, and 4) evaluation of adverse ocular events. Fundus photography (fluorescein and indocyanine green angiography) was performed at baseline, 3 months, and 12 months and at any monthly visit between 4 and 11 months on the basis of retreatment criteria.

**Endpoints and outcome measures**

Outcome measures were as follows: 1) mean change in study eye visual acuity; 2) mean change in CRT; 3) mean number of IVR injections over the 6-month period; and 4) adverse ocular events at 12 months.

**Statistical analyses**

To determine if there was a significant difference in changes in BCVA and CRT, repeated measures of analysis of variance
with Greenhouse–Geisser and Bonferroni corrections were performed. Independent samples t-tests were conducted to determine the statistical significance of differences in visual acuity, CRT, and number of injections between the RM and RI groups. All statistical analyses were performed using the SPSS software V.20 (IBM Corporation, Armonk, NY, USA). P-values < 0.05 were considered significant, and all values are reported as mean ± standard deviation.

**Results**

Baseline demographic and clinical data of the participants are shown in Table 1. In general, the baseline demographics between the two groups were well matched, with no significant differences between the groups. All 58 patients completed the study.

**Visual acuity**

The mean BCVA score improved significantly in both treatment groups, with the largest mean change observed in the RI group from baseline (Figure 1).

The mean change in BCVA from baseline to 12 months was −0.12±0.04 LogMAR in the RM group (P=0.04) and −0.20±0.04 LogMAR in the RI group (P=0.001), with the degree of change being significantly different between the two groups (P=0.04).

At 12 months, the mean BCVA in the RI group (0.35±0.14 LogMAR) was significantly higher than that in the RM group (0.45±0.18 LogMAR; P=0.04). Table 2 shows the changes in visual acuity at key time points.

**Table 1 Baseline patient characteristics**

<table>
<thead>
<tr>
<th>Sex, n (%)</th>
<th>RM group (n=29)</th>
<th>RI group (n=29)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Male</td>
<td>13 (44.8)</td>
<td>12 (41.4)</td>
</tr>
<tr>
<td>Female</td>
<td>16 (55.2)</td>
<td>17 (58.6)</td>
</tr>
<tr>
<td>Age (years)</td>
<td>75.1±8.9</td>
<td>75.0±7.1</td>
</tr>
<tr>
<td>Range</td>
<td>59–87</td>
<td>58–83</td>
</tr>
<tr>
<td>Visual acuity</td>
<td>0.56±0.25</td>
<td>0.56±0.29</td>
</tr>
<tr>
<td>Mean ± SD (LogMAR)</td>
<td>0.1–1</td>
<td>0.1–1</td>
</tr>
<tr>
<td>CRT (µm)</td>
<td>Mean ± SD</td>
<td></td>
</tr>
<tr>
<td>417±81.9</td>
<td>422±73.6</td>
<td></td>
</tr>
<tr>
<td>Lesion composition, n (%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Classic/predominantly classic</td>
<td>13 (44.8)</td>
<td>12 (41.4)</td>
</tr>
<tr>
<td>Minimally classic/occult</td>
<td>16 (55.2)</td>
<td>17 (58.6)</td>
</tr>
</tbody>
</table>

**Notes:** RM group: monotherapy with monthly 0.5 mg intravitreal ranibizumab injections; RI group: monthly 0.5 mg intravitreal ranibizumab injections with topical indomethacin 0.5% eye drops three times a day.

**Abbreviations:** CRT, central retinal thickness; RI, ranibizumab plus indomethacin; RM, ranibizumab monotherapy.

**Figure 1** Change in visual acuity in both groups over 12 months of treatment.

Notes: The red line indicates best-corrected visual acuity in patients treated with monthly 0.5 mg intravitreal ranibizumab and 0.5% indomethacin eye drops three times a day, and the blue line indicates best-corrected visual acuity in patients treated with monthly 0.5 mg intravitreal ranibizumab monotherapy. Mean values with standard error bars are shown for each time point.

**CRT**

A constant and significant reduction in mean CRT was observed in the RM group (−101±12.6 µm; P<0.0001) and in the RI group (−134±14.8 µm; P<0.0001) from baseline to 12 months. Figure 2 shows the changes in CRT in both groups over the 12-month treatment period. At 12 months, the mean CRT in the RM group (316±41.2 µm) was significantly higher than that in the RI group (287±31.5 µm; P=0.004). Changes in CRT over the 12-month treatment period are shown in Table 2.

**Number of injections**

The mean number of required IVR injections was 7.38±0.78 in the RM group and 6.34±0.67 in the RI group (P<0.001).

**Safety and adverse events**

No serious adverse events were observed during the study period. There was no significant difference in the number of adverse ocular events described by either group. Mild burning/stinging was reported more frequently in the RI group, while pain was similar between groups. Adverse ocular events are reported in Table 3. No issues regarding compliance with the indomethacin therapy were recorded for any patient.

**Discussion**

The results of this prospective pilot study suggest that the topical administration of indomethacin, when used in combination with ranibizumab, may produce further improvements in BCVA, retinal thickness, and the number of required IVR injections.
Ocular adverse events in the study eyes during the 12-month study period

Table 3

<table>
<thead>
<tr>
<th>Ocular adverse events</th>
<th>RM group, n (%)</th>
<th>RI group, n (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Abnormal sensation</td>
<td>9 (31)</td>
<td>12 (41)</td>
</tr>
<tr>
<td>Burning/stinging</td>
<td>14 (48)</td>
<td>16 (55)</td>
</tr>
<tr>
<td>Eye pain</td>
<td>4 (14)</td>
<td>4 (14)</td>
</tr>
<tr>
<td>Itchy eye</td>
<td>6 (21)</td>
<td>8 (28)</td>
</tr>
<tr>
<td>Headache</td>
<td>4 (14)</td>
<td>3 (10)</td>
</tr>
<tr>
<td>Sore eyelid</td>
<td>4 (14)</td>
<td>4 (14)</td>
</tr>
<tr>
<td>Retinal hemorrhage</td>
<td>1 (3)</td>
<td>0 (0)</td>
</tr>
<tr>
<td>Foreign body sensation</td>
<td>10 (34)</td>
<td>16 (55)</td>
</tr>
<tr>
<td>Conjunctivitis</td>
<td>6 (21)</td>
<td>4 (14)</td>
</tr>
<tr>
<td>Dry eye</td>
<td>7 (24)</td>
<td>9 (31)</td>
</tr>
<tr>
<td>Eye strain</td>
<td>8 (28)</td>
<td>9 (31)</td>
</tr>
<tr>
<td>Light sensitivity</td>
<td>12 (41)</td>
<td>11 (38)</td>
</tr>
<tr>
<td>Floaters</td>
<td>15 (50)</td>
<td>14 (48)</td>
</tr>
</tbody>
</table>

Notes: RM group: monotherapy with monthly 0.5 mg intravitreal ranibizumab injections; RI group: monthly 0.5 mg intravitreal ranibizumab injections with topical indomethacin 0.5% eye drops three times a day. *P<0.05 compared with baseline value for the same group. Abbreviations: RI, ranibizumab plus indomethacin; RM, ranibizumab monotherapy.
be $1,078 per injection)\textsuperscript{17} intravitreal therapy, which can be associated with serious adverse events such as endophthalmitis, retinal tears, and retinal detachment.

No serious adverse events were reported in our study. The results show that the treatments were well tolerated in all groups, with a safety profile comparable to that observed in previous studies. Furthermore, compliance with eye drop use was very high and the incidence of reported adverse events was mild to moderate.

Our study has a few limitations: first, we included a limited number of patients in a pilot study design. Second, the evaluation of adverse events was limited to eye and headache symptoms. Third, we only assessed indomethacin TID. Further studies with a larger sample size, various dosing regimens, and more in-depth follow-up are warranted to validate the results of this pilot study, especially considering the substantial visual decline during the late stages of neovascular AMD.\textsuperscript{18}

**Conclusion**

The results of this pilot study suggest that topical 0.5% indomethacin administered TID with IVR has an additive effect on CRT reduction in CNV. As previously reported, further larger studies are needed to confirm these data, especially evaluating the long-term efficacy of NSAIDs because AMD is a chronic disease with an inflammatory component. In particular, careful attention should be paid to the corneal complications associated with long-term use of topical NSAIDs.

**Acknowledgment**

This study was presented in the AAO 2016 Meeting; Chicago, IL, USA. October 15, 2016 – October 18, 2016.

**Disclosure**

The authors report no conflicts of interest in this work.

**References**