Visual outcomes of age-related macular degeneration patients undergoing intravitreal ranibizumab monotherapy in an urban population

Khadijah Basheer
Evelyn Mensah
Tina Khanam
Neda Minakaran
Ophthalmology Department, Central Middlesex Hospital, London, UK

Aim: To compare the visual outcomes of an urban population with age-related macular degeneration (AMD) undergoing ranibizumab monotherapy to the results from major clinical trials.

Procedures: Prospective data was collected from 164 wet AMD patients receiving intravitreal ranibizumab. Visual acuities were obtained with the Early Treatment Diabetic Retinopathy Study chart. All patients underwent a loading phase of three monthly treatments of ranibizumab. Patients were monitored monthly using a retreatment criterion. Treatment was further individualized by sequentially lengthening follow-up intervals when stable.

Results: At 12 and 24 months, respectively, the percentage of eyes that maintained vision was 91% and 88.6%. We found that 20.3% of eyes had improved vision at 12 months and 20% at 24 months. At 12 months, 8.3% of eyes’ vision worsened and 12% worsened at 24 months.

Conclusion: Individualized ranibizumab monotherapy is effective in preserving vision in wet AMD and follows the same trends as the pivotal trials.

Keywords: visual acuity, comparison of age-related macular degeneration treatment trials, choroidal neovascular membrane

Introduction
Age-related macular degeneration (AMD) is the commonest cause of severe visual impairment in older people in the developed world.1,2 The two late phenotypes of AMD are geographic atrophy and wet AMD; together, these are responsible for two-thirds of registrations of visual impairment or blindness in the United Kingdom.2 Vision loss in wet AMD is caused by formation of a choroidal neovascular membrane (CNVM) responding to vascular endothelial growth factor (VEGF). It is thought that wet AMD accounts for only 10%–15% of all AMD cases but is responsible for over 80% of AMD-related severe visual impairment or visual loss.3

The minimally classic/occult trial of the anti-VEGF antibody ranibizumab in the treatment of neovascular AMD (MARINA) and anti-VEGF antibody for the treatment of predominantly classic choroidal neovascularization in AMD (ANCHOR) trials were large multicenter double-blind studies which established that continuous treatment with intravitreal ranibizumab was an effective treatment for wet AMD.4,5 Following this, comparison of age-related macular degeneration treatment trials (CATT) proved that treatment with ranibizumab on an as-required basis following the initial three monthly loading dose was equally as effective as continuous monthly treatments.6,7

The CATT trial retreatment criteria was based on signs of CNVM activity: fluid on optical coherence tomography (OCT), new or persistent hemorrhage, decreased visual acuity as compared with the previous examination, dye leakage, or increased lesion size on fundus fluorescein angiography.6
Following these trials, ranibizumab intravitreal injections have been employed as first-line treatment for wet AMD. Since this change in the treatment of wet AMD, little has been published in the UK regarding real population visual outcomes. We wanted to analyze our inner-city AMD patient cohort and compare their visual outcomes following ranibizumab monotherapy to the major clinical trials.

Aims
To compare the results from an inner-city population to the results achieved in the major clinical trials described in the introduction.

Methods
From December 2007 to January 2012, prospective data were collected from 164 patients who underwent intravitreal ranibizumab treatment for wet AMD and entered into an in-house, password-protected spreadsheet. Baseline and follow-up visual acuity measurements were obtained with the Early Treatment Diabetic Retinopathy Study (ETDRS) chart. Patient consent to publish their demographics and visual outcome data was gained when the decision to proceed with treatment was made. The study was approved by the ethics committee at London North West Healthcare NHS Trust.

The patients who were included in the study were aged 50 years or older; had visual acuity between 0.00 and 2.30 logMAR; and had the presence in the study eye of previously untreated active CNVM as demonstrated by hemorrhage or leakage on FFA and OCT.

Patients who had received previous treatment, transferred to another trust, or who subsequently had another diagnosis were excluded from our analysis, as were patients who had no more than the three treatments in the loading phase. After applying these exclusions, 27 patients were removed from the study and the data for 123 eyes of 106 patients were analysed. None of the patients refused treatment. Data were collected for age, sex, baseline visual acuity, and monthly visual acuities over 2 years and for the mean number of treatments per year. This study did not analyze the disease progression of different AMD types; we solely wanted to report the visual outcome data of a real-life population undergoing ranibizumab monotherapy.

All patients underwent a loading phase of three monthly treatments of 0.5 mg in 0.05 mL of intravitreal ranibizumab. Following this, patients were monitored monthly and the following retreatment criteria were used to establish whether further treatment was required:
- Loss of letters with fluid on OCT
- Any fluid on OCT – intraretinal fluid, subretinal fluid
- Increased size of pigment epithelial detachment
- New retinal hemorrhage
- New CNVM.

In addition, if FFA revealed active CNVM, treatment was repeated each month for three months. Treatment was further individualized by sequentially lengthening follow-up intervals when a period of stability had been established.

Each intravitreal ranibizumab injection was performed in theatre under aseptic conditions and chloramphenicol 0.5% eye drops were administered four times a day for 5 days following the treatment. During and post-procedure, the patients were monitored for topical and general adverse events.

Similar to both the ANCHOR and MARINA trials, efficacy analyses were performed on an intention-to-treat basis with the use of a last-observation-carried-forward method for missing data.4,5

Results
Demographics
Similar to the ANCHOR trial but in contrast to the MARINA and CATT trials, our results had a majority of male patients (59%). Our population with a mean age of 83 years was much older than patients included in the other trials. Our largest age group was between 80 and 89 years compared to 75–84 years in the ANCHOR and MARINA trials and 70–79 years in the CATT trial. All the trials had a Caucasian majority of between 97% and 98%; we also demonstrate a Caucasian majority but of only 62% with 38% of patients from other ethnicities. Demographics are summarized in Table 1.

Visual outcomes
We compared our 1-year and 2-year visual outcomes to those found from the ANCHOR, MARINA, and CATT trials (Figures 1 and 2). We used their same criteria for defining the

| Table 1 Comparison of present study demographics and mean baseline vision to pivotal trials |
|---------------------------------|-----------------|-----------------|-----------------|-----------------|
| Demographics                    | Present study   | CATT trial      | ANCHOR trial    | MARINA trial    |
| Age                             | 83              | 79              | 76              | 77              |
| Mean (years)                    | 80–89           | 70–79           | 75–84           | 75–84           |
| Largest group (years)           | Male (%)        | 51              | 40              | 54              | 37              |
| Sex                             | Female (%)      | 49              | 59              | 46              | 63              |
| Ethnicity                       | Caucasian (%)   | 62              | 98              | 97              | 97              |
| Other (%)                       | 38              | 3               | 4               | 3               |
| Mean baseline vision (ETDRS letters) | 35             |                 |                 |                 |

Abbreviation: ETDRS, Early Treatment Diabetic Retinopathy Study.
**Year 1 visual outcomes**

![Graph comparing the visual outcomes after 1 year from this study to the CATT, ANCHOR, and MARINA trials.](#)

**Abbreviation:** ETDRS, Early Treatment Diabetic Retinopathy Study.

**Year 2 visual outcomes**

![Graph comparing the visual outcomes after 2 years from this study to the CATT, ANCHOR, and MARINA trials.](#)

**Abbreviation:** ETDRS, Early Treatment Diabetic Retinopathy Study.
three visual outcomes. The primary outcome measure was the percentage of eyes that maintained vision, meaning the percentage of eyes that lost less than 15 letters from baseline.4–6 Secondary outcome measures included the percentage of eyes that improved vision (meaning those that gained 15 letters or more from baseline) and the mean change in visual acuity.4–6 All 123 eyes underwent at least 12 months of monitoring and 61 eyes (50% of the initial study population) had visual outcome data for 24 months.

At 12 months, the percentage of eyes that maintained vision in our study was 91% compared to 96% in the CATT trial, 96.4% in the ANCHOR trial, and 94.6% in the MARINA trial. We found that 20.3% of eyes had improved vision compared to 25% in the CATT trial, 40.3% in the ANCHOR trial, and 33.8% in the MARINA trial. Finally, 8.1% of our study eyes’ vision worsened compared to 4% in the CATT trial, 3.6% in the ANCHOR trial, and 5.4% in the MARINA trial (Table 2).

At 24 months, the percentage of eyes that maintained vision in our study was 91% compared to 96% in the CATT trial, 96.4% in the ANCHOR trial, and 94.6% in the MARINA trial.4–6 We found that 20.3% of eyes had improved vision compared to 25% in the CATT trial, 40.3% in the ANCHOR trial, and 33.8% in the MARINA trial. Finally, 8.1% of our study eyes’ vision worsened compared to 4% in the CATT trial, 3.6% in the ANCHOR trial, and 5.4% in the MARINA trial (Table 2).

When comparing the mean visual progress over 2 years, our results did not show the initial steep gain of letters that was shown in the trials (Figure 3). Instead, the mean change in vision over 2 years was only between zero and three ETDRS letters. We also found that the change in mean visual acuity was minimal ranging between 0.58 and 0.70 logMAR over 2 years (Figure 4). The mean number of injections given per eye was ten over 2 years; six in year 1 (range three to 12) and four in year 2 (range one to 12), compared to 12.6 in the CATT trial (Table 2).

On investigating this further we found that from the group whose vision worsened by 15 letters or more, 73% of these eyes had severe vision loss (30 letters or more; range 16–97). Although this is a relatively small percentage of the total number of eyes (9%), this significant loss of letters may have been enough to skew the mean visual acuity and mean change in vision, thus hiding any initial dramatic increase in vision.

### Discussion

The pivotal continuous and individualized clinical trials have changed how wet AMD is treated today. The trials were either continuous, where monthly injections of ranibizumab were given over 2 years, or individualized where, after the three monthly loading doses, the patient was evaluated monthly but ranibizumab therapy was only given according to set criteria relating to the activity of wet AMD in that individual. The ANCHOR and MARINA trials were the two continuous trials that recruited 423 and 716 patients, respectively. They both observed that 90% of patients maintained vision, and improvement in vision occurred in 41% of patients in the ANCHOR trial and 35% in the MARINA trial. The mean change in visual acuity from baseline was 10.7 letters in the ANCHOR trial and 6.6 letters in the MARINA trial. The recently published CATT trial was significantly larger with 1,208 patients. They compared visual outcomes of eyes that underwent treatment with ranibizumab to those that had
bevacizumab and also the visual outcomes of patients who underwent continuous treatment versus individualized, as-needed treatment. The arm of the trial that observed individualized treatment with ranibizumab showed maintenance of vision in 93% of patients, improvement of vision in 33% of patients, and a mean change in visual acuity of 6.7 letters over 2 years.

The results from our study follow a similar pattern to the pivotal trials, where the initial findings from year 1 were largely maintained throughout year 2. At 2 years, the percentage of eyes that maintained vision was 88.6%, compared to 93% in the CATT trial and 90% in the ANCHOR and MARINA trials. However, the number of eyes that improved vision was approximately a third less than the pivotal trials; 19.5% compared to 33% in the CATT trial, 41% in the ANCHOR trial, and 35% in the MARINA trial. The number of eyes whose vision worsened by losing more than 15 letters was greater than in all the trials; 11.5% compared to 7% in the CATT trial, and 4% and 6% in the ANCHOR.
Table 3 Comparison of present study results to ten other studies of real life aMD populations treated with ranibizumab monotherapy around the world

<table>
<thead>
<tr>
<th></th>
<th>Present study UK</th>
<th>Bandukwala et al11 Canada</th>
<th>Cohen et al10 France</th>
<th>Dadgostar et al12 USA</th>
<th>Feng et al17 China</th>
<th>Malgorzata and Stankiewicz3 Poland</th>
<th>Hjelmqvist et al14 Sweden</th>
<th>Kang et al15 Korea</th>
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<td>Male (%)</td>
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<td>60</td>
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<td>90</td>
<td>83.8</td>
<td>84.9</td>
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<td>gain of more than</td>
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<td>change (ETDRS letters)</td>
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<td>5.2</td>
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<td>2</td>
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<td>10.9</td>
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Abbreviations: aMD, age-related macular degeneration; eTDRS, Early Treatment Diabetic Retinopathy Study; N/A, not applicable; P, prospective; R, retrospective.

Conclusion

This prospective study of individualized ranibizumab monotherapy over 24 months shows that individualized therapy in an inner-city population is effective in preserving and gaining vision compared to other single centers. Currently, there is no data published from the United Kingdom. However, ten other papers have been published by other centers (see Table 3 and Figure 3), and patients who failed to attend further appointments (n=4). Patients with poor visual outcomes often had treatment with ranibizumab monotherapy over 24 months showed that individualized therapy is effective in preserving vision in an inner-city population is effective in preserving vision.

The authors report no conflicts of interest in this work.

Disclosure

MARINA trials; respectively. The lack of dramatic visual improvement over 2 years, was also observed in the mean visual acuity over 2 years, which remained fairly static at 0.6 logMAR (range -0.1 to 2.0). As mentioned previously, this may be due to the fact that 9% of eyes did particularly poorly, and the average visual outcomes. Our results do follow the same trend as the pivotal trials with low patient numbers. Other reasons included: patients who failed to attend further appointments (n=4). Patients with poor visual outcomes often had treatment with ranibizumab monotherapy over 24 months showed that individualized therapy is effective in preserving vision in an inner-city population is effective in preserving vision.

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References


