Newly diagnosed exudative age-related macular degeneration treated with pegaptanib sodium monotherapy in US community-based practices: medical chart review study

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Abstract

Background: Studies have shown that early detection and treatment of neovascular age-related macular degeneration (NV-AMD) can delay vision loss and blindness. The objective of this study was to evaluate the efficacy/safety of intravitreal pegaptanib sodium monotherapy in treatment-naïve subjects with newly diagnosed NV-AMD and to gain insight into characteristics of lesions treated in community-based practices.

Methods: From seven private US practices, charts were retrospectively reviewed on 73 subjects with previously untreated subfoveal choroidal NV-AMD treated with their first dose of pegaptanib monotherapy on/after 4/1/2005 through 6/5/2006, receiving ≥4 treatments at 6-week intervals over 21 weeks. Primary endpoint: mean visual acuity (VA) change from baseline to month 6.

Results: 75% of lesions were occult, and 82% were subfoveal. From baseline to month 6, mean VA change was -0.68 lines; 58% and 16% gained ≥0 and ≥3 lines of VA, and 70% were responders (<3 lines lost). In 35 subjects with early disease, 80% were responders with a mean gain of 0.46 lines.

Conclusion: Pegaptanib is effective in real-world patients with treatment-naïve NV-AMD in uncontrolled community-based retina practices.

Background

Age-related macular degeneration (AMD) is a chronic, progressive disease that results in a loss of central vision and significant functional impairment. It is the leading cause of blindness in Western developed countries [1,2]. Neovascular AMD (NV-AMD) represents 10 to 15% of all AMD cases but accounts for 90% of AMD-related severe vision loss [3]. Choroidal neovascularization (CNV) causes disruption of the structure and function of the retinal pigment epithelium and the retinal photoreceptors. Prevalence of late forms of AMD (defined as the presence of NV-AMD or geographic atrophy [4]) increases exponentially with age [5].

Rapid vision loss is a key characteristic of NV-AMD, such that the proportion of untreated patients who develop severe vision loss (> 6 lines) can reach up to 42% in 3 years of follow-up [6]. Additionally, patients with CNV in one eye have an estimated 43% probability of progression to NV-AMD in the fellow eye within 5 years [7]. Patients with visual impairment from AMD lose independence, suffer from fall-related injuries, experience high levels of depression and anxiety, develop annoying visual hallucinations, and require low vision aids [8-12]. Direct medical and non-medical costs and the cost of progression to blindness all contribute to the economic burden of AMD both to the patient and to society. Estimated annual societal costs of all NV-AMD patients in Canada, France, Germany, Spain, and the United Kingdom are substantial, ranging from €671 to €3278 million [13].

Studies show that the early detection and treatment of AMD can delay vision loss and blindness and thus significantly reduce the economic burden of the disease [14,15]. The primary purpose of early treatment ideally
would be to defer progression or to promote visual improvement. Prior to the development of anti-vascular endothelial growth factor (VEGF) inhibitors, NV-AMD patients were treated with laser photocoagulation and photodynamic therapy (PDT) with verteporfin (Visudyne®). PDT had limited use as it was only approved to treat the predominantly classic type lesion, representing approximately 20% of NV-AMD patients [16] and it merely slowed vision loss [17].

The quest for alternative treatment options in NV-AMD has been precipitated by the increasing prevalence of the disease and by the associated side effects and unsatisfactory outcomes with approved therapies. Anti-VEGF treatments, such as pegaptanib sodium (Macugen®), ranibizumab (Lucentis®) and bevacizumab (Avastin®), are the first pharmacological treatments to address an underlying pathological factor of the CNV of NV-AMD and to address disease progression without healthy ocular tissue destruction. Pegaptanib sodium, a selective anti-VEGF therapy approved for the treatment of all subtypes of NV-AMD, was introduced into the US market in January 2005. Results from phase II/III pivotal clinical trials [18,19] showed that pegaptanib was effective in patients with subfoveal NV-AMD regardless of lesion subtype (i.e. predominantly classic, minimally classic, or occult). Approximately 70% of patients treated with 0.3 mg pegaptanib had stabilised or improved vision (lost < 15 letters [< 3 lines] compared to baseline) at 54 weeks compared to 55% of patients receiving standard-of-care treatment. Additionally, pegaptanib was well tolerated, with the majority of adverse events being ocular in nature and transient.

Currently available efficacy and safety data for pegaptanib are from clinical trials, which may not accurately reflect pegaptanib’s real-world use and potential visual outcomes. Further, there is a need to understand which disease characteristics define earlier lesions in order to identify patients who may have a better response to anti-VEGF therapy (i.e. pegaptanib). We performed a retrospective chart review study in newly diagnosed NV-AMD patients initially treated with 0.3 mg pegaptanib in the US to evaluate actual clinical experience with intravitreal pegaptanib monotherapy and to explore the characteristics of lesions in patients in whom a better response to pegaptanib monotherapy was observed.

Methods
Study design
This retrospective medical chart review study included 73 newly diagnosed NV-AMD subjects recruited from seven retina specialist offices/clinics in different geographic regions of the US who were treated with pegaptanib monotherapy. Subjects were required to have at least one eye (study eye) that was newly diagnosed with NV-AMD and that was previously untreated for this condition prior to pegaptanib therapy. Subjects had best-corrected Snellen visual acuity (VA) of 20/40 to 20/200 in the study eye when pegaptanib therapy was initiated and were free from any other ocular pathology that would impair VA. Subjects must have received pegaptanib monotherapy for a minimum of four treatments at 6-week intervals over a 21-week period in the study eye, with initial therapy on or after 1 April 2005 through 5 June 2006. All subjects were ≥50 years of age and were excluded if they had participated in an investigational drug study within the study period. Site study staff identified potential charts for review according to study inclusion and exclusion criteria and contacted subjects of the potential charts to obtain written informed consent prior to data abstraction. No medical interventions or invasive procedures were required by the study protocol.

This study was conducted according to the tenets of the Declaration of Helsinki. The study protocol and subject informed consent document were approved by the Allendale Investigational Review Board, a central human investigation review board. This research was compliant with Health Insurance Portability and Accountability Act policies and procedures.

Data abstracted
Abstracted chart data included subjects’ demographic characteristics, NV-AMD diagnosis date, VA at the time of NV-AMD diagnosis, comorbid medical conditions, baseline clinical characteristics of the study eye (including VA, angiographic subtype, lesion size, lesion characteristics and lesion location based on fluorescein angiography) prior to pegaptanib monotherapy initiation, date of injection visits, VA assessment at each injection visit and ocular adverse events.

Endpoints and statistical analysis
The primary endpoint was mean change in best-corrected VA from baseline to month 6. Secondary endpoints included proportions of subjects who gained ≥0, 1, 2 and 3 lines of VA and proportion of those who lost <3 lines, 3 to <6 lines and 6 or more lines in best-corrected VA from baseline to month 6. Safety analysis was performed based on a pre-specified list of adverse events common with intravitreal injections.

We calculated summary statistics (means and standard deviations [SD] for continuous variables and frequency distributions for categorical variables) to describe sample demographic and clinical characteristics at baseline and clinical characteristics of study eyes at each treatment visit. Mean change in VA from baseline was calculated both in logarithm of the minimum angle of resolution (logMAR) and in line units of VA at each treatment.
visit and at the month 6 visit minus VA at the baseline visit.

We compared mean change in VA (logMAR; t-test or Wilcoxon nonparametric test) and proportions of subjects who gained or lost VA (Fisher’s exact test) from baseline to month 6 between those classified as having early lesions and those not having early lesions. Two definitions of early disease (early lesion) from the VISION study [20] were used for the assessment. Definition No.1 defined early disease as a lesion size of < 2 disc areas, baseline VA in the study eye of ≥ 20/80 (≥ 54 Early Treatment of Diabetic Retinopathy Study [ETDRS] letters), and absence of scarring or atrophy within the lesion. Definition No.2 defined early disease as occult with no classic CNV, absence of lipid, and better VA at baseline in the fellow eye (i.e. worse VA at baseline in the study eye). Further, we evaluated subject baseline characteristics and VA change from baseline to month 6 across study sites to assess whether any differences in treatment patterns, subject characteristics or outcomes were observed among sites using t-test, Wilcoxon test, or Fisher’s exact test, as appropriate.

Statistical significance was evaluated at the 0.05 level with no adjustments for multiple comparisons. All analyses were performed using PC-SAS version 9.1 (SAS Institute, Cary, NC, USA).

Results
Baseline demographics and clinical characteristics
Data were collected between August and November 2006 from 73 NV-AMD subjects’ medical charts. Subject demographic characteristics and comorbid conditions are summarised in Table 1. The median age of the subjects was 79 years (range, 58–92 years). A majority of the subjects were female (62%) and had at least one comorbid disease (86%).

At baseline, the mean (SD) VA in the study eye was 0.62 (0.24) logMAR, equivalent to approximately 20/80 Snellen, and the mean time from NV-AMD diagnosis to baseline (first treatment) was 2.4 months (Table 2). A majority of subjects had occult lesions (75%); lesion size < 4 disc areas (66%); subfoveal lesion locations (82%). Few subjects had pigment epithelial detachment, retinal angiomatous proliferation, cystoid macular oedema (CME), fibrosis or geographic atrophy. Approximately one-third (34%) had presence of blood reported.

VA change from baseline
Overall sample
On average, subjects’ VA remained stable through the fourth pegaptanib treatment. There was a slight decline from baseline to the month 6 visit (mean change: 0.07 logMAR [-0.68 lines]; Table 3), with 58% of subjects gaining ≥ 0 lines, 16% gaining ≥ 3 lines, 12% losing > 0 to < 3 lines, and 11% losing ≥ 6 lines. In all, 70% of subjects lost < 3 lines of VA.

Evaluation of mean VA change from baseline to month 6 by baseline NV-AMD characteristics showed that only angiographic subtype was significantly associated with VA change: subjects with occult lesions had improvement in mean VA (-0.01 logMAR [0.09 lines]) while those with predominantly classic and minimally classic lesions had a decline in mean VA (0.27 and 0.34 logMAR, [-2.64 and -3.50 lines], respectively; overall p-value = 0.0065 from one-way analysis of variance).

By early disease definitions
Eighteen subjects met early disease definition No.1 (lesion size < 2 disk areas, baseline VA of the study eye ≥ 20/80 and absence of scarring or atrophy), and 35 subjects met early disease definition No. 2 (occult with

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**Table 1 Baseline demographic characteristics and comorbid conditions.**

| Characteristic                                      | Study sample |
|-----------------------------------------------------|--------------|
| Age (years)                                         | 73           |
| Mean (SD)                                           | 78.3 (7.0)   |
| Median (range)                                      | 79 (58 – 92) |
| Gender, n (%)                                       |              |
| Female                                              | 45 (61.6)    |
| Male                                                | 27 (37.0)    |
| Missing                                             | 1 (1.4)      |
| Ethnicity, n (%)                                    |              |
| White, non-Hispanic                                 | 26 (35.6)    |
| Missing                                             | 47 (64.4)    |
| Reported comorbid disease, n (%)                    | 63 (86.3)    |
| Comorbid disease, n (%)                             |              |
| Diabetes                                            | 5 (6.8)      |
| Cancer                                              | 13 (17.8)    |
| Asthma                                              | 4 (5.5)      |
| Chronic obstructive pulmonary disease               | 2 (2.7)      |
| Arthritis and rheumatism                            | 22 (30.1)    |
| Headache (migraine, cluster)                        | 1 (1.4)      |
| Chronic neck or back pain                           | 1 (1.4)      |
| Heart disease                                       | 12 (16.4)    |
| Stroke                                              | 2 (2.7)      |
| Sleep disturbance                                   | 1 (1.4)      |
| Comorbid disease categories, n (%)                  |              |
| Ocular*                                             | 31 (42.5)    |
| Other                                               | 55 (75.3)    |
| Comorbid diseases†                                   |              |
| N                                                   | 63           |
| Mean (SD)                                           | 3.1 (1.5)    |
| Median (range)                                      | 3 (1 – 7)    |

* Includes diabetic retinopathy, glaucoma, congenital degeneration of the retina, ocular tumour, cataract, and low vision due to other reasons.
† Among subjects with comorbid conditions.
SD = standard deviation.
no classic CNV, absence of lipid, and better VA in the fellow eye at baseline). Subjects with missing data who could not be classified with regard to early disease definitions were excluded from the analyses reported below.

Subjects meeting early disease definition No.1 showed a decline in mean VA at all treatment visits, and a significant difference in VA change was observed at the third visit relative to those who did not meet the definition (0.08 versus -0.06 logMAR [-0.89 versus 0.61 lines]; \( p = 0.0283 \); Table 4). Subjects meeting early disease definition No.2 showed improvement in mean VA from baseline at all treatment visits (\( p < 0.05 \) for the third, fourth, and month 6 visits difference between those who met the definition and those who did not meet the definition). Additionally, 26% of subjects meeting early disease definition No.2 had \( \geq 3 \)-line gains from baseline to month 6 compared with only 6% among subjects meeting early disease definition No.1 (Figure 1). Using early disease definition No.2, 80% of patients were responders (lost < 3 lines of VA). Mean change in VA from baseline to month 6 was a loss of 0.16 logMAR (-1.61 lines) for definition No.1 and an improvement of -0.05 logMAR (0.46 lines) for definition No.2, with the latter group showing more favourable outcomes compared to the entire sample (0.07 logMAR [-0.68 lines]).

Evaluation of baseline subject clinical characteristics showed that subjects meeting early disease definition

| Table 2 Baseline clinical characteristics of study eyes. |
|--------------------------------------------------------|
| Characteristic                                         | Study eye |
| Best-corrected VA (logMAR)                             |           |
| N                                                      | 73        |
| Mean (SD)                                             | 0.62 (0.24) |
| Median (range)                                        | 0.54 (-0.18 – 1.00) |
| Time from NV-AMD diagnosis to baseline, months         |           |
| N                                                      | 73        |
| Mean (SD)                                             | 2.4 (5.9) |
| Median (range)                                        | 0 (0 – 31) |
| Angiographic subtype, n (%)                            |           |
| Minimally classic                                     | 4 (5.5)  |
| Occult                                                | 55 (75.3) |
| Predominantly classic                                 | 11 (15.1) |
| Missing                                               | 3 (4.1)  |
| Lesion size (disc area), n (%)                         |           |
| < 2                                                   | 26 (35.6) |
| ≥ 2 and < 4                                           | 22 (30.1) |
| ≥ 4                                                   | 11 (15.1) |
| Missing                                               | 14 (19.2) |
| Lesion location, n (%)                                 |           |
| Subfoveal                                             | 60 (82.2) |
| Extrafoveal                                           | 1 (1.4)  |
| Juxtafoveal                                           | 6 (8.2)  |
| Missing                                               | 6 (8.2)  |
| Pigment epithelial detachment, n (%)                   |           |
| Absent                                                | 46 (63.0) |
| Present                                               | 16 (21.9) |
| Missing                                               | 11 (15.1) |
| Retinal angiomatic proliferation, n (%)                |           |
| Absent                                                | 58 (79.5) |
| Present                                               | 1 (1.4)  |
| Missing                                               | 14 (19.2) |
| Cystoid macular oedema, n (%)                          |           |
| Absent                                                | 52 (71.2) |
| Present                                               | 7 (9.6)  |
| Missing                                               | 14 (19.2) |
| Fibrosis, n (%)                                        |           |
| Absent                                                | 57 (78.1) |
| Present                                               | 3 (4.1)  |
| Missing                                               | 13 (17.8) |
| Geographic atrophy, n (%)                              |           |
| Absent                                                | 54 (74.0) |
| Present                                               | 10 (13.7) |
| Missing                                               | 9 (12.3)  |
| Presence of blood, n (%)                               |           |
| Yes                                                   | 25 (34.2) |
| No                                                    | 43 (58.9) |
| Missing                                               | 5 (6.8)  |

Table 2: Baseline clinical characteristics of study eyes. (Continued)

| Estimated percentage of lesion with blood, n (%)       |           |
| ≤ 50%                                                 | 12 (16.4) |
| > 50%                                                 | 3 (4.1)   |
| Missing                                               | 58 (79.5) |

NV-AMD = neovascular age-related macular degeneration; SD = standard deviation; VA = visual acuity.

### Table 3 Visual acuity (VA; logMAR*) in the study eye by treatment.

| Treatment visit | Mean VA | Mean VA change from baseline |
|-----------------|---------|-----------------------------|
|                 | n Mean (SD) | n Mean (SD) |
| Second treatment| 73 0.60 (0.32) | 73 -0.02 (0.25) (↑) |
| Third treatment | 73 0.59 (0.28) | 73 -0.03 (0.23) (↑) |
| Fourth treatment| 73 0.63 (0.35) | 73 0.01 (0.29) (↑) |
| Month 6 visit   | 73 0.69 (0.39) | 73 0.07 (0.33) (↑) |

* ↑ improved; ↓ declined. Negative logMAR changes indicate improvement; positive logMAR changes indicate declination.

LogMAR = logarithm of the minimum angle of resolution; SD = standard deviation.
No.1 had a mean of 1.1 (median 0.0) months from NV-AMD diagnosis to baseline assessment while those who did not meet the definition for early disease had a mean of 3.1 (median 0.0) months, a difference that did not reach statistical significance in either the mean or the median. Even though more subjects meeting definition No.1 had CME (17% versus 10%, respectively) and pigment epithelial detachment (28% versus 20%, respectively) and fewer were reported to have the presence of blood (22% versus 34%, respectively) compared to those not meeting the definition, none of the differences reached statistical significance (Table 5). Subjects meeting early disease definition No.2 had a mean of 2.5 (median 0.0) months from NV-AMD diagnosis to baseline assessment while those who did not meet this definition had a mean of 0.9 (median 0.0) months (p-values not significant). Even though fewer subjects meeting early disease definition No.2 had CME (3% versus 19%, respectively) or presence of blood (26% versus 48%, respectively) and more subjects had pigment epithelial detachment (31% versus 7%, respectively), only the difference for pigment epithelial detachment reached statistical significance (Table 5).

By study sites
When mean VA change from baseline to month 6 was evaluated by study sites, one site (No.7) showed a mean VA gain of -0.04 logMAR (0.50 lines) compared to a mean loss of VA ranging from 0.02-0.30 logMAR (-0.17 to -3.00 lines) for all other sites. Figure 2 shows proportions of subjects with VA change by site. Demographic and baseline clinical characteristics of the subjects were compared between site No.7 and all other sites to explore any subject or practice characteristics that might lead to substantially more subjects treated at site No.7 maintaining or improving their VA from baseline to month 6 compared to other sites (83% versus 49%, respectively; p = 0.0134; Table 6). There was no statistically significant difference observed in subjects’ baseline demographic or clinical characteristics between the sites. However, the mean time from NV-AMD diagnosis to

| Early disease definition | Met definition | Did not meet definition | p-value |
|--------------------------|---------------|-------------------------|---------|
| Definition No.1<sup>1</sup> |               |                         |         |
| Baseline to second treatment visit | 18 | 0.04 (0.33) | 41 | -0.06 (0.21) | 0.1702 |
| Baseline to third treatment visit | 18 | 0.08 (0.29) | 41 | -0.06 (0.20) | 0.0283 |
| Baseline to fourth treatment visit | 18 | 0.12 (0.35) | 41 | -0.04 (0.26) | 0.0597 |
| Baseline to month 6 visit | 18 | 0.16 (0.39) | 41 | 0.03 (0.31) | 0.1799 |
| Definition No.2<sup>2</sup> |               |                         |         |
| Baseline to second treatment visit | 35 | -0.06 (0.18) | 31 | 0.03 (0.32) | 0.1627 |
| Baseline to third treatment visit | 35 | -0.09 (0.18) | 31 | 0.03 (0.28) | 0.0339 |
| Baseline to fourth treatment visit | 35 | -0.08 (0.22) | 31 | 0.09 (0.34) | 0.0155 |
| Baseline to month 6 visit | 35 | -0.05 (0.28) | 31 | 0.13 (0.34) | 0.0235 |

<sup>1</sup>Negative logMAR changes indicate improvement; positive logMAR changes indicate declination.
<sup>2</sup>Early disease definition No.1: lesion size < 2 disk area, baseline VA study eye ≥ 20/80 Snellen, no scar or atrophy.
<sup>3</sup>Early disease definition No.2: occult and better VA at baseline in fellow eye (worse in study eye).

LogMAR = logarithm of the minimum angle of resolution; SD = standard deviation.
initiation of treatment was much shorter for site No.7 subjects compared to that of the other sites overall (0.33 versus 3.13 months, respectively; p = 0.0808), and substantially more subjects at site No.7 did not have geographic atrophy (94% versus 67%, respectively; p = 0.0545).

**Ocular safety**

Pegaptanib appears to be safe for use (Table 7). There was one report of endophthalmitis and two reports of geographic atrophy (occurrence rates of 0.003 and 0.006, respectively, in 326 injections), but the history of these subjects is not known.

**Discussion**

The efficacy-related findings of this real-world observational study are similar to those previously reported in pegaptanib clinical trials. In the current study, the mean change in VA from baseline to month 6 was a decline of -0.68 lines (0.07 logMAR), and 70% of subjects lost fewer than three lines. The VISION trials [18] showed a decline of approximately 8 letters, and 70% of subjects lost fewer than three lines of VA after treatment over a period of 54 weeks.

There is no universally accepted definition of early CNV secondary to AMD. The early lesion definitions used in this analysis were based on those used in the exploratory analysis of the VISION study results [20], matching as closely as possible those definitions using the data we had available. Subjects in our study who met early definition No.2 had significantly greater improvement in VA from baseline to month 6 compared to the decline in VA in those who did not meet the definition (-0.05 versus 0.13 logMAR [0.46 versus -1.29 lines], respectively; p = 0.0235). This result was likely driven by the high proportion of patients with occult lesions in the study while few subjects with predominantly classic or minimally classic lesions were included. This suggests that clinicians when selecting out earlier lesions are defining these lesions with occult characteristics. Our current study found that, on average, subjects with occult lesions treated with pegaptanib had an improvement in VA from baseline to month 6 while subjects with predominantly classic or minimally classic lesions had a decline in VA (-0.01 versus 0.27 versus 0.34 logMAR [0.09 versus -2.64 versus -3.50 lines], respectively; p = 0.0065).

In this study, we observed a significant difference in clinical efficacy in subjects across study sites. In particular, 83% of subjects treated at site No.7 showed either an improvement or maintenance of VA from baseline to month 6, with a mean improvement of -0.04 logMAR (0.50 lines), while subjects from all other sites had a mean decline of 0.11 logMAR (-1.07 lines), with only 49% of subjects showing either an improvement or maintenance of VA (p = 0.0134). The earlier treatment of subjects at site No.7 might have accounted for the better results - the mean time from NV-AMD diagnosis to treatment was 0.3 months for site No.7 compared to 3.1 months for all other sites (p = 0.0808). Whether or not anti-VEGF efficacy can be related to the duration of existing CNV disease has not been clearly defined. In a recent study by Boyer et al. [21], efficacy of ranibizumab did not appear to differ across the different quartiles of duration of NV-AMD, which is the opposite of what we found in this study with pegaptanib administration.

Although large randomised, prospective studies have confirmed benefit with treatment, the question remains as to which type of CNV lesions responds best to treatment. Different characteristics were identified in previous studies with no consensus. For example, the TAP [Treatment of Age-related Macular Degeneration with Photodynamic Therapy] and VIP [Verteporfin in Photodynamic Therapy] Study Groups [22] concluded that lesion size was a strong predictor of VA outcome based upon observed PDT treatment and outcome while the VISION trials [20] found that anti-VEGF treatment of early lesions (defined on the basis of the combined characteristics of size, lipid, scarring and time of presentation) had a positive impact on VA outcome. A different conclusion was drawn from the MARINA study [21] subanalysis of subjects with minimally classic and occult lesions treated with ranibizumab in which neither lesion size nor duration of NV-AMD was found to have a direct relationship with VA outcomes. Our study also attempted to evaluate which of the subjects' baseline angiographic characteristics had an impact on VA outcome and found that those who met the early disease definition using the criteria of occult with no classic CNV, absence of lipid, and better VA at baseline in the fellow eye had better VA outcomes. There is no agreed upon and accepted early lesion definition at the present time.

This study intended to evaluate newly diagnosed NV-AMD patients treated with pegaptanib monotherapy. Even though we did not provide participating clinicians with a definition of ‘newly diagnosed,’ we did not find differences in angiographic characteristics across patients' time since diagnosis. However, we did observe a relationship between time since diagnosis and primary study outcomes of VA change from baseline to month 6 with shorter time being associated with better outcomes. The finding is consistent with the oncology literature that shows that younger tumours that are earlier in their angiogenic process of new blood vessel development appear to be more susceptible to certain types of anti-VEGF therapy [23,24].
There are several limitations to this study. The study included only those practices and patients who were willing to participate. Two sites had a total of four patients who met study inclusion criteria and who were willing to participate, with one patient having maintained VA and the other three having lost more than three lines from baseline to month 6. We cannot be certain that these four individuals are representative of the universe of NV-AMD patients treated in these two practices. Conversely, one does not know how representative patients from site No.7 are to the NV-AMD population either. The sample size of the study limited our ability to perform multivariate analyses that might better support interpreting study findings. In addition, although the proportion of patients losing VA from baseline to month 6 appears to correlate with the presence of classic lesions, this is most likely biased by the fact that our sample included few subjects with classic lesions. Further, the study did not collect optical coherence tomography data for each visit, which might have provided a more accurate summary of NV-AMD characteristics, enabling us to evaluate the relationship between VA change and disease characteristics. Finally, the data were collected for 6 months only; it is not known if the

| Characteristic                                      | Met definition | Did not meet definition | p-value | Met definition | Did not meet definition | p-value |
|-----------------------------------------------------|----------------|-------------------------|---------|----------------|-------------------------|---------|
| Best-corrected VA (logMAR), mean (SD)               | 0.43 (0.10)    | 0.68 (0.27)             | 0.0010  | 0.62 (0.24)    | 0.64 (0.27)             | 0.7536  |
| Time from NV-AMD diagnosis to baseline (months), mean (SD) | 1.11 (3.12)    | 3.12 (6.94)             | 0.2456  | 2.49 (5.43)    | 0.90 (2.77)             | 0.1488  |
| Angiographic subtype, n (%)                         |                |                         | 0.6817  |                |                         | <0.0001 |
| Minimally classic                                   | 1 (5.6)        | 3 (7.3)                 | 0 (0.0) | 4 (12.9)       |                         |         |
| Occult                                              | 13 (72.2)      | 33 (80.5)               | 35 (100.0) | 16 (51.6)       |                         |         |
| Predominantly classic                               | 4 (22.2)       | 5 (12.2)                | 0 (0.0) | 11 (35.5)      |                         |         |
| Missing                                             | 0 (0.0)        | 0 (0.0)                 | 0 (0.0) | 0 (0.0)        |                         |         |
| Lesion location, n (%)                              |                |                         | 0.1345  |                |                         | 1.0000  |
| Subfoveal                                           | 14 (77.8)      | 38 (92.7)               | 30 (85.7) | 28 (90.3)       |                         |         |
| Extralevel                                           | 1 (5.6)        | 0 (0.0)                 | 0 (0.0) | 0 (0.0)        |                         |         |
| Juxtapfoveal                                        | 2 (11.1)       | 3 (7.3)                 | 3 (8.6) | 2 (6.5)        |                         |         |
| Missing                                             | 1 (5.6)        | 0 (0.0)                 | 2 (5.7) | 1 (3.2)        |                         |         |
| Pigment epithelial detachment, n (%)                |                |                         | 0.5091  |                |                         | 0.0202  |
| Present                                             | 5 (27.8)       | 8 (19.5)                | 11 (31.4) | 2 (6.5)        |                         |         |
| Absent                                              | 13 (72.2)      | 33 (80.5)               | 19 (54.3) | 26 (83.9)       |                         |         |
| Missing                                             | 0 (0.0)        | 0 (0.0)                 | 5 (14.3) | 3 (9.7)        |                         |         |
| Cystoid macular oedema, n (%)                       |                |                         | 0.6639  |                |                         | 0.0582  |
| Present                                             | 3 (16.7)       | 4 (9.8)                 | 1 (2.9) | 6 (19.4)       |                         |         |
| Absent                                              | 15 (83.3)      | 37 (90.2)               | 26 (74.3) | 22 (71.0)       |                         |         |
| Missing                                             | 0 (0.0)        | 0 (0.0)                 | 8 (22.9) | 3 (9.7)        |                         |         |
| Fibrosis n (%)                                      |                |                         | 1.0000  |                |                         | 0.1585  |
| Present                                             | 0 (0.0)        | 2 (4.9)                 | 1 (2.9) | 2 (6.5)        |                         |         |
| Absent                                              | 18 (100.0)     | 39 (95.1)               | 26 (74.3) | 27 (87.1)       |                         |         |
| Missing                                             | 0 (0.0)        | 0 (0.0)                 | 8 (22.9) | 2 (6.5)        |                         |         |
| Geographic atrophy, n (%)                           |                |                         | 0.1637  |                |                         | 0.2125  |
| Present                                             | 0 (0.0)        | 6 (14.6)                | 4 (11.4) | 5 (16.1)       |                         |         |
| Absent                                              | 18 (100.0)     | 35 (85.4)               | 25 (71.4) | 25 (80.6)       |                         |         |
| Missing                                             | 0 (0.0)        | 0 (0.0)                 | 6 (17.1) | 1 (3.2)        |                         |         |
| Presence of blood, n (%)                            |                |                         | 0.5404  |                |                         | 0.0806  |
| Yes                                                 | 4 (22.2)       | 14 (34.1)               | 9 (25.7) | 15 (48.4)       |                         |         |
| No                                                  | 14 (77.8)      | 27 (65.9)               | 24 (68.6) | 16 (51.6)       |                         |         |
| Missing                                             | 0 (0.0)        | 0 (0.0)                 | 2 (5.7)  | 0 (0.0)        |                         |         |

* Early disease definition No.1: lesion size < 2 disk area, baseline VA study eye ≥ 20/80 Snellen, no scar or atrophy within the lesion.
†Early disease definition No.2: occult and better VA at baseline in fellow eye (worse in study eye).

LogMAR = logarithm of the minimum angle of resolution; NV-AMD = neovascular age-related macular degeneration; SD = standard deviation; VA = visual acuity.

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results would have been different over a longer treatment period.

**Conclusion**

The efficacy of pegaptanib in community-based practices appears to confirm findings from the VISION trials and a published analysis of patients with earlier disease. There appears to be a trend for patients with earlier lesions to respond more favourably to pegaptanib. Due to our small sample size, there was significant variability of outcomes by site and by patient. Still, shorter time from NV-AMD diagnosis to initiation of pegaptanib treatment appears to be associated with better treatment outcomes and enhanced clinical benefits. It is a common theme across the medical and scientific literature that earlier intervention, prior to permanent damage, is more likely to achieve a beneficial outcome. Other large NV-AMD clinical studies [21,22] have attempted to define and interpret outcome based on lesion characteristics. In this study, though, other disease characteristics did not seem useful in identifying *a priori* responders to treatment. Further research is warranted to fully understand and determine NV-AMD disease characteristics that help predict outcomes. Their results will enable NV-AMD therapy to be targeted to provide the greatest benefit to both patients and society.

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**Table 6 Comparison of baseline characteristics and clinical efficacy between site No.7 versus other sites.**

| Characteristic                                      | Site No.7 | Other sites | p-value |
|-----------------------------------------------------|-----------|-------------|---------|
| Baseline best-corrected VA (logMAR)                 |           |             |         |
| Mean (SD)                                           | 0.61 (0.26) | 0.62 (0.24) | 0.8429  |
| Time from NV-AMD diagnosis to baseline (months)     |           |             |         |
| N                                                   | 18        | 55          |         |
| Mean (SD)                                           | 0.33 (0.77) | 3.13 (6.65) | 0.0808  |
| Median (range)                                      | 0.0 (0–3) | 0.0 (0–31)  |         |
| Angiographic subtype, n (%)                         |           |             | 0.1937  |
| Minimally classic                                  | 0 (0.0)   | 4 (7.3)     |         |
| Occult                                             | 12 (66.7) | 43 (78.2)   |         |
| Predominantly classic                              | 5 (27.8)  | 6 (10.9)    |         |
| Missing                                            | 1 (5.6)   | 2 (3.6)     |         |
| Lesion size (disc area), n (%)                     |           |             | 0.1376  |
| <2                                                 | 9 (50.0)  | 17 (30.9)   |         |
| ≥ 2 and < 4                                        | 7 (38.9)  | 15 (27.3)   |         |
| ≥ 4                                                | 1 (5.6)   | 10 (18.2)   |         |
| Missing                                            | 1 (5.6)   | 13 (23.6)   |         |
| Geographic atrophy, n (%)                          |           |             | 0.0545  |
| Present                                            | 0 (0.0)   | 10 (18.2)   |         |
| Absent                                             | 17 (94.4) | 37 (67.3)   |         |
| Missing                                            | 1 (5.6)   | 8 (14.5)    |         |
| Presence of blood, n (%)                           |           |             | 0.4277  |
| Present                                            | 4 (22.2)  | 21 (38.2)   |         |
| Absent                                             | 13 (72.2) | 30 (54.5)   |         |
| Missing                                            | 1 (5.6)   | 4 (7.3)     |         |
| Met early disease definition No.1*                 | 5 (29.4)  | 13 (31.0)   | 1.0000  |
| Met early disease definition No.2†                 | 5 (29.4)  | 30 (61.2)   | 0.0281  |
| VA, baseline to month 6 (logMAR)                    |           |             |         |
| Mean (SD)                                           | -0.04 (0.22) | 0.11 (0.36) | 0.0970  |
| VA change, baseline to month 6                      |           |             |         |
| Gained ≥ 0 lines                                    | 15 (83.3) | 27 (49.1)   | 0.0134  |
| Gained ≥ 3 lines                                    | 3 (16.7)  | 9 (16.4)    | 1.0000  |
| Lost < 3 lines                                      | 1 (5.6)   | 8 (14.5)    | 0.4365  |
| Lost ≥ 6 lines                                      | 0 (0.0)   | 8 (14.5)    | 0.1871  |

* Early disease definition No.1: lesion size <2 disk area, baseline VA study eye ≥ 20/80 Snellen, no scar or atrophy.
† Early disease definition No.2: occult and better VA at baseline in fellow eye (worse in study eye).

LogMAR = logarithm of the minimum angle of resolution; NV-AMD = neovascular age-related macular degeneration; SD = standard deviation; VA = visual acuity.
Table 7 Ocular adverse events over a 6-month period from pegaptanib treatment initiation.

| Adverse event                | Number of subjects with occurrence | Total number of occurrences | Rate of occurrence by number of injections* |
|------------------------------|------------------------------------|-----------------------------|--------------------------------------------|
| Endophthalmitis              | 1†                                 | 1†                          | 0.003                                      |
| Retinal detachment           | 0                                  | 0                           | 0                                          |
| Increased intraocular pressure| 0                                  | 0                           | 0                                          |
| Retinal tear                 | 0                                  | 0                           | 0                                          |
| Traumatic cataract           | 0                                  | 0                           | 0                                          |
| Vitreous haemorrhage         | 0                                  | 0                           | 0                                          |
| Geographic atrophy           | 2†                                 | 2†                          | 0.006                                      |
| Other                        | 0                                  | 0                           | 0                                          |

* Based on 326 total injections received by study subjects.
† Unclear history.

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