Multispectral Pattern Recognition Reveals a Diversity of Clinical Signs in Intermediate Age-Related Macular Degeneration

Angelica Ly,1,2 Lisa Nivison-Smith,1,2 Nagi Assaad,1,3,4 and Michael Kalloniatis1,2

1Centre for Eye Health, University of New South Wales, Sydney, New South Wales, Australia
2School of Optometry and Vision Science, Faculty of Science, University of New South Wales, Sydney, New South Wales, Australia
3Department of Ophthalmology, Prince of Wales Hospital, Randwick, New South Wales, Australia
4Department of Ophthalmology, Sutherland Hospital, Taren Point, New South Wales, Australia

Correspondence: Michael Kalloniatis, Centre for Eye Health, School of Optometry and Vision Science, Faculty of Science, University of New South Wales, Sydney, New South Wales, Australia
m.kalloniatis@unsw.edu.au.
Submitted: September 29, 2017 Accepted: March 6, 2018
Citation: Ly A, Nivison-Smith L, Assaad N, Kalloniatis M. Multispectral pattern recognition reveals a diversity of clinical signs in intermediate age-related macular degeneration. Invest Ophthalmol Vis Sci. 2018;59:1790-1799. https://doi.org/10.1167/iovs.17-25076

Purpose. To develop a proof-of-concept, computational method for the quantification and classification of fundus images in intermediate age-related macular degeneration (AMD).

Methods. Multispectral, unsupervised pattern recognition was applied to 184 fundus images from 10 normal and 36 intermediate AMD eyes. The imaging results of preprocessed, grayscale images from three modalities (infrared, green, and fundus autofluorescence) were automatically classified into various clusters sharing a common spectral signature, using a k-means clustering algorithm. Class separability was calculated by using transformed divergence (D_T). The classification results for large drusen, pigmentary abnormalities, and areas unaffected by AMD were compared against three expert observers for concordance, and to calculate sensitivity and specificity.

Results. Multispectral, unsupervised pattern recognition successfully identified a finite number of AMD-specific, statistically separable signatures in eyes with intermediate AMD. By using a correct classification criterion of >83% for identical clusters and a total of 1693 expert annotations, the sensitivity and specificity of multispectral pattern recognition for the detection of AMD lesions was 74% and 98%, respectively. Large drusen and pigmentary abnormalities were correctly classified in 75% and 68% of instances, respectively.

Conclusions. We describe herein a novel approach for the classification of multispectral images in intermediate AMD. Automated classification of intermediate AMD, using multispectral pattern recognition, has moderate sensitivity and high specificity, when compared against clinical experts. The methods described may have a future role in AMD screening or monitoring.

Keywords: macular disease, image analysis, multimodal, ocular imaging, drusen

Age-related macular degeneration (AMD) is defined as a degenerative disease of the retinal pigment epithelium and is a leading cause of visual impairment worldwide.1 It is subdivided clinically into three stages—early, intermediate, and late.2 Early AMD features small to medium-sized drusen, while in intermediate AMD the drusen are large (>125 μm) and may be accompanied by hyper- or hypopigmentary abnormalities. Late AMD represents the most advanced form associated with significant visual morbidity and presents as two subtypes—geographic atrophy or neovascular AMD.

Accurate detection and classification of AMD is fundamental to optimal management and mitigation of future vision loss in select patients. For instance, earlier identification of patients with intermediate AMD may enable earlier adoption of preventative strategies such as nutritional supplementation or cessation of smoking.3 Routine assessment by primary care providers, however, may result in the misclassification of 25% of eyes with early or moderate AMD as normal.4 Several factors contribute to the difficulty and a lack of uniformity in AMD assessment.5 The condition is phenotypically heterogeneous. Evaluation of clinical markers for determining AMD stage and risk of progression (drusen size, pigmentary abnormalities), using funduscopy and/or retinal photography, may also vary between clinicians.6 Other disease-specific prognostic biomarkers that may better identify an individual’s overall risk of progression may also not be routinely considered in a clinical setting. For example, drusen regression represents an ominous sign of AMD progression associated with geographic atrophy7–9 but is typically laborious to identify, necessitating time and expertise-intensive alignment and inspection of retinal photographs taken on different occasions, if available. Drusen subtypes have also been identified, and this subtyping may be applied to better identify an individual’s overall risk of progression.10

The accurate assessment of AMD is further complicated by the advent of multiple advanced imaging modalities that are now widely available for the clinical evaluation and differential diagnosis of macular structure (optical coherence tomography; infrared imaging; monochromatic retinal photography—blue-, red-, or green-filtered images; and fundus autofluorescence). These technologies enable earlier and more accurate disease detection by providing detailed phenotype data with high...
Multispectral Pattern Recognition in AMD

However, each imaging modality is also beset with a range of specific advantages and limitations. For example, infrared imaging holds the highest sensitivity for reticcular pseudodrusen, yet may not provide the same prognostic insight as fundus autofluorescence. A computational approach integrating the findings from multiple modalities might potentially assist in identifying robust markers of disease state with greater consistency. This could potentially solve the problem of identifying areas of drusen regression by emphasizing change within a certain signature. Several digital image classification studies have already explored various automated and semiautomated techniques for AMD screening. However, relatively few have considered using a pixel-based pattern recognition approach to integrate results of current commercially available modalities in intermediate AMD. Thus, the aim of this study was to develop and apply a computational approach (multispectral pattern recognition) to statistically classify the spectral characteristics of normal eyes and lesions in AMD, and to subsequently appraise its clinical utility against expert grading.

**METHODS**

This was a retrospective, diagnostic study of 36 eyes of 20 patients with bilateral intermediate AMD who attended the Centre for Eye Health for a macular assessment in 2015 (Supplementary Fig. S1). The Centre for Eye Health is an intermediate-tier eye-care establishment that provides imaging and visual system diagnostic services, operating primarily as a service to eye-health practitioners in commercial practice so that they can optimally manage their patients. Intermediate AMD was defined according to the Beckman initiative for macular research clinical classification scheme, and included any eye with at least one large druse (≥125-μm diameter or exceeding the average width of a retinal vein at the disc margin) within 3 mm of the macular center or pigmentary abnormalities associated with medium drusen in patients aged 55 years or older. Ten normal eyes (five patients) were also analyzed. Eyes with incomplete or poor image quality, pseudophakia, or macular disease other than intermediate AMD were excluded. Patient written consent was obtained in accordance with the Declaration of Helsinki and approved by a Biomedical Human Research Ethics Advisory Panel of the University of New South Wales, Sydney, Australia.

Methods used for the pattern recognition analysis of images are summarized in Figure 1. Each classified image series, from each eye, included readily available ocular imaging modalities: a reference color fundus photograph, and 8-bit grayscale fundus autofluorescence (FAF). Grayscale IR815nm, and Green532nm; and infrared (IR815nm, Spectralis Heidelberg Retina Angiograph 2; Heidelberg Engineering, Heidelberg, Germany) were also registered and masked. No further manipulations were performed on the color photographs. The three preprocessed IR815nm, Green532nm, and FAF images were assigned to the red, green, and blue channel, respectively, and preliminarily assessed qualitatively in rgb triplet format by using the merge channels function in ImageJ (Fig. 1B, panel 2).

**Multispectral Pattern Recognition**

Multispectral digital analysis uses the spectral data from each pixel in a series of images generated by using narrow bands of light at various imaging wavelengths. Different wavelengths penetrate the fundus to different depths and hence different modalities contain unique information regarding lesion characteristics and location. Pixel values (ranging from 0 to 255 for an 8-bit image) from each unit in an image are categorized into common spectral theme classes and the results outputted by using a thematic pseudocolor map or classified image. The clinical meaning of these classes may then be determined a posteriori.

Grayscale, preprocessed IR815nm, Green532nm, and FAF images from each eye were automatically classified by using unsupervised K-means cluster analysis and commercially available satellite imaging software (http://www.pigeomatics.com/; provided in the public domain by PCI Remote Sensing, Markham, Ontario, Canada) with the following parameters: 16 maximum classes, 16 maximum iterations, and a 0.01 minimum threshold (Fig. 1B, panel 3). The principles of cluster analysis have been described elsewhere. Although other clustering algorithms and parameters were explored, we describe herein the best combination as judged by the authors, using simultaneous comparison against the reference fundus photograph (Supplementary Fig. S2). Class separability was calculated by using transformed divergence (DT; KCLUS algorithm, PCI Remote Sensing), as previously described. Briefly, DT describes a method of calculating the separability or statistical difference (correct classification) between spectral signatures. It derives from measuring the difference in pixel values between all pairs from a predefined number of groups. DT measures are presented as a value between zero and two: zero indicates extremely poor classification accuracy or overlap between signatures of two classes, while two indicates complete separation between the two classes (analogous to 100% classification accuracy).

Pattern recognition theory was originally developed for satellite image analysis and encompasses clustering analysis and tests for cluster separability. The cluster separability indices include DT, which is effectively similar to separability indices reported by univariate signal detection theory used in psychology, but DT is able to report separability in N-dimensional space. A larger separability value equates to better classification accuracy, and the statistic is resistant to relatively large deviations from normality. Increasing the number of data channels only increases the value of the statistic if the additional channel contributes further separability.

The Final Image Series for Each Eye

Each application of cluster analysis resulted in a classified image, or pseudocolor map, whereby a hue represents a unique class or spectral signature. The two classes with the lowest separability were merged and the process repeated until a minimum DT cutoff value between all classes was met. We investigated the diagnostic utility of multispectral,
FIGURE 1. Schematic showing the data extraction, processing, qualitative and quantitative analyses used in the study. (A) Input imaging results from each eye included a reference color fundus photograph, infrared, green, and fundus autofluorescence scanning laser ophthalmoscopy image. Note the numerous differences in image alignment, magnification, and size. (B) Following preprocessing, the imaging results were merged and reviewed qualitatively as an rgb triplet image then quantitatively by using pattern recognition analysis. (C) Final image series from one eye. Each color in the classified image represents a class or spectral signature of retinal disease. Three accuracy levels or endpoints for pattern recognition classification were considered ($D_1 > 1.5$, $D_2 > 1.7$, $D_3 > 1.9$). Percentages refer to the probability of correct classification, derived by Swain and King.32 (D) The sensitivity and specificity of the method was evaluated by using a ground truth determined by expert grading. CFP, color fundus photograph; IR, infrared; FAF, fundus autofluorescence.
unsupervised pattern recognition by using three endpoint
criteria—D_T equal to 1.5, 1.7, or 1.9, which translates to a
>83%, >90%, and >98% probability of correct classification,
respectively (Fig. 1C). The final output is a statistically
robust, classified image derived from multispectral pattern
recognition for each eye.

**Ground Truth: Evaluation of Diagnostic Accuracy
Against Expert Grading**

To evaluate the diagnostic accuracy of the classified images, a
ground truth was defined by using two clinical experts (highly
trained optometrists with over 15 years of combined optometric
experience, with 8 years providing ocular imaging
diagnostic services in an optometry-ophthalmology collabora-
tive care setting). The graders manually annotated 36 color
fundus photographs for all visible (1) large drusen, (2)
hyperpigmentary abnormalities, and (3) hypopigmentary
abnormalities (up to a maximum of 20 instances each), using
the Cell Counter plugin in ImageJ. Graders also annotated an
additional 20 locations each indicating blood vessels and
background. The graders were not advised of the diagnosis of
any of the study eyes but instructed to use the Beckman
initiative definitions of the lesions. Drusen grading circles C0
(63-μm diameter) and C1 (125-μm diameter) were appended to
each image for comparative purposes. Disagreements between
the two graders were resolved by a third grader (AL). Ten
percent of all annotations were also confirmed by a medical
retinal specialist (NA). Optical coherence tomography findings
were not considered during the grading; however, select
images are included in this article for illustrative purposes.

For the classified images, each class was manually catego-
rized as either an AMD lesion (coincident predominantly with
large drusen, hyper- or hypopigmentary abnormalities) or
background class (blood vessels or background). In the
classified images, eyes without an identifiable AMD lesion
class were considered poorly classified and excluded from the
sensitivity analysis (5/36, 14%; case example shown in Fig. 2C
and further described in the Results), resulting in a total of
1693 annotations across 31 eyes with intermediate AMD.
Classified images and the annotated color fundus reference
photograph for the 31 eyes were then compared for
concordance and agreement. True positives of the lesions
(which for simplicity we describe as sensitivity) and true
negatives (specificity) were calculated (Fig. 1D).

**RESULTS**

**Pattern Recognition Applied to Normal Eyes**

In the normal group, there were three females, three
Caucasians, and two Asians aged between 60 and 66 years
(Table 1). Macular images from normal, healthy eyes were
characterized by relatively low spectral diversity, that is, greater
uniformity in the grayscale input images (Fig. 2A). The
rgb images consequently displayed a minimal range of colors or
phenotype diversity. Comparatively, eyes with AMD showed a
AMD with limited agreement between the reference fundus photo-
graph and classified images. In eye (C), the classified image lacks any
identifiable AMD lesion classes (recognizable large drusen, hyper-
or hypopigmentary abnormalities). Note that a stricter D_T criterion was
associated with poorer discrimination of AMD lesions. In eyes (A) and
(B), the central artifact in the IR815nm image was identified as a
unique signature in the classified images. The lack of changes in the
normal eye provides strong evidence that the algorithm is indeed
identifying AMD-specific lesions.
more diverse appearance in the classified images (Fig. 2B). Imaging artifacts were identified as a unique signature (Figs. 2A, 2B).

Pattern Recognition Applied to AMD Eyes

AMD participants were predominantly female (13/20, 65%), Caucasian (17/20, 85%), with ages ranging between 56 and 82 years. Three patients were Asian (3/20, 15%). In the multispectral pattern recognition classified images (also described as pseudocolor maps), each separable spectral signature was identified by a distinct color. Well-classified AMD images showed notable colocalization with the distribution and position of drusen and other AMD lesions, while poorly classified eyes did not (i.e., Fig. 2B versus Fig. 2C). The latter typically also displayed a lower inherent signal to noise ratio (ratio of brightness to darkness of AMD lesions despite preprocessing) in the input images, suggesting the result might be a limitation of the input modalities. Incidentally, the rgb triplet images were also a useful means of validating the accuracy of image registration. Poor registration typically appeared as kerf\textsuperscript{35} surrounding blood vessels in the rgb image and where visible, prompted realignment.

Across the AMD dataset, the number of unique spectral signatures (colors in the pseudocolor maps) in intermediate AMD ranged between 2 and 11 (Table 2). Classification accuracy for the AMD annotations ranged between 79% and 100% for each eye and depended on the underlying appearance of AMD lesions in the composite imaging. A stricter, that is, higher DT endpoint, was associated with poorer discrimination of lesions. Consequently, the DT\textsubscript{1.5} criterion (>83% probability of correct classification) displayed the greatest diagnostic utility. In that classification set, 1543 of 1695 annotations were correctly classified (91% agreement). Two annotations could not be classified owing to convolution by imaging artifacts. Large drusen, hyperpigmentary abnormalities, and hypopigmentary abnormalities were concordantly discriminated in 75%, 68%, and 75% of all annotated instances, respectively, yielding an overall sensitivity of 74% for AMD lesions. By comparison, classification accuracy of blood vessels and background was greater at 99% and 96%, respectively, consistent with a specificity of 98% (Table 1). Without excluding the poorly classified eyes (n = 5), 1743/1932 annotations were correctly classified, resulting in an overall sensitivity and specificity of 68% and 98%, respectively. An additional analysis was conducted to determine if the number of samples from each eye affected overall proportions used to calculate sensitivity, such as to skew the result. There was no statistically significant correlation (P = 0.49) between the percentage of correct classification and the number of AMD lesion annotations on an eye level.

Lesion-Specific Observations

This section describes additional observations attained by using empirical, successive comparison of the grayscale images, rgb images, pseudocolor maps, and the reference fundus photographs. Each color (class) of the pseudocolor map representing a separable spectral signature corresponded to a specific anatomic structure, such as the fundus background, blood vessels, or a distinct type of AMD lesion (drusen, hyper- or hypopigmentary abnormalities; Fig. 3). Using this approach with a DT criterion of 1.5, a signature corresponding to large drusen was separable in 31 of 35 eyes (89%). Hyperpigmentary abnormalities and hypopigmentary abnormalities were identified in 4 of 10, and 1 of 3 eyes (33%), respectively.

Large drusen in each input, grayscale image typically appeared brighter than the background signal, especially in the Green\textsubscript{532nm} and FAF images (Fig. 4), and consequently appeared green or blue-green in the rgb images. Interestingly, the drusen often colocalized with more than one theme class in one eye, including changes within or encircling the drusen (intra- or perilesional signatures in 18 eyes; Fig. 4B). Occasionally, several classes represented by various pseudocolors were discovered across drusen (both within and between units in the rgb and classified images), reminiscent of drusen diversity (Fig. 4C). Before pattern recognition, these drusen appeared similar in the fundus images. The three different drusen-related signatures in Figure 4C correlated with differences in the internal reflectivity of the drusen and in the

---

**Table 1. Baseline Demographic and Descriptive Data of the Normal and Intermediate AMD Cohort**

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Control Patients</th>
<th>Intermediate AMD</th>
</tr>
</thead>
<tbody>
<tr>
<td>No. of subjects</td>
<td>5</td>
<td>20</td>
</tr>
<tr>
<td>No. of eyes, n (%)</td>
<td>10</td>
<td>50</td>
</tr>
<tr>
<td>Right</td>
<td>5 (50)</td>
<td>17 (47)</td>
</tr>
<tr>
<td>Left</td>
<td>5 (50)</td>
<td>19 (53)</td>
</tr>
<tr>
<td>Age, y</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Median (IQR)</td>
<td>61 (6)</td>
<td>73 (6)</td>
</tr>
<tr>
<td>Range</td>
<td>60–66</td>
<td>56–82</td>
</tr>
<tr>
<td>Sex, n (%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Female</td>
<td>3 (60)</td>
<td>13 (65)</td>
</tr>
<tr>
<td>Male</td>
<td>2 (40)</td>
<td>7 (35)</td>
</tr>
<tr>
<td>Ethnicity, n (%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Caucasian</td>
<td>3 (60)</td>
<td>17 (85)</td>
</tr>
<tr>
<td>Asian</td>
<td>2 (40)</td>
<td>3 (15)</td>
</tr>
</tbody>
</table>

---

**Table 2. Classification Accuracy of Multispectral Pattern Recognition Using the Total Dataset From 36 Eyes With Intermediate AMD and Three Transformed Divergence (DT) Endpoints**

<table>
<thead>
<tr>
<th>Diagnostic Index</th>
<th>DT\textsubscript{1.5} Endpoint Correct Classification &gt;83%</th>
<th>DT\textsubscript{1.7} Endpoint Correct Classification &gt;90%</th>
<th>DT\textsubscript{1.9} Endpoint Correct Classification &gt;98%</th>
</tr>
</thead>
<tbody>
<tr>
<td>Agreement</td>
<td>1543/1695 (91%)</td>
<td>1509/1695 (89%)</td>
<td>1491/1695 (88%)</td>
</tr>
<tr>
<td>Disagreement</td>
<td>1481/1695 (9%)</td>
<td>1821/1695 (11%)</td>
<td>2001/1695 (12%)</td>
</tr>
<tr>
<td>Unclassified owing to artifacts</td>
<td>2/1695 (&lt;1%)</td>
<td>2/1695 (&lt;1%)</td>
<td>2/1695 (&lt;1%)</td>
</tr>
<tr>
<td>Sensitivity</td>
<td>74%</td>
<td>71%</td>
<td>62%</td>
</tr>
<tr>
<td>Specificity</td>
<td>98%</td>
<td>96%</td>
<td>98%</td>
</tr>
<tr>
<td>Minimum No. of classes(^*)</td>
<td>5</td>
<td>3</td>
<td>2</td>
</tr>
<tr>
<td>Maximum No. of classes(^*)</td>
<td>11</td>
<td>8</td>
<td>6</td>
</tr>
<tr>
<td>Average No. of classes(^*)</td>
<td>7</td>
<td>5</td>
<td>4</td>
</tr>
</tbody>
</table>

\(^*\) The number of classes is denoted by the number of colors in each pseudocolor map or classified image. For further details, please refer to the text under the subsection entitled “Lesion Specific Observations.”
integrity of the inner segment ellipsoid zone overlying the drusen.

If separable, pigmentary abnormalities were typically assigned to a single class (Fig. 5). Hyperpigmentary abnormalities appeared bright in IR815nm and FAF imaging though dark (decreased reflectivity, increased absorption) in the Green532nm images, rendering them bright pink in the rgb images (Figs. 5A, 5B). Hypopigmentary abnormalities appeared least commonly of all the AMD lesions (in three eyes only, as based on expert annotation) and where present, appeared blue-green in rgb (Fig. 5C). A lack of separability occurred twice and may relate to convolution by underlying drusen.

DISCUSSION

Despite a range of widely available ocular imaging technologies, the diagnosis of AMD and its subtypes is still complex, incongruous, and difficult. Signs of AMD, such as drusen or pigmentary abnormalities, may be hard to identify owing in part to low contrast, interindividual variations, and
red saturation of the fundus. This study showcased the high specificity of a multispectral, unsupervised pattern recognition approach in classifying intermediate AMD, using multimodal imaging. Specifically, we found AMD lesions have specific spectral signatures that can be displayed on a single multispectral pseudocolor map to simplify multimodal image interpretation.

Applications for Multispectral Pattern Recognition

The strength of this study lies in its capacity to objectively and quantitatively integrate the unique findings of multiple imaging modalities. In particular, this method appeared to capitalize on the advantages of scanning laser ophthalmoscopy—namely, high contrast, averaging, and depth discrimination (axial optical sectioning). Evaluating the finite number of colors in the output pseudocolor map is simpler than assessing multiple images and therefore is amenable to interpretation by nonexperts, which might shorten the time and resources required for grading. Input imaging results were also well-tolerated en face modalities that can be acquired undilated.

The steps that allowed useful classification of retinal architecture relied upon (1) accurate image registration, (2) preprocessing, including pseudobackground correction and the contrast enhancement steps, (3) appropriate masking of nonmacular areas, and (4) stepwise, down merging in the unsupervised classification approach. Unprocessed images were widely prone to spurious classifications confounded by nonmacular signatures, physiological variations in visibility of the choroidal vasculature, differences in fundus pigmentation (Supplementary Fig. S1), and background intensity gradient differences across the image (dependent on the acquisition conditions). By beginning with a large predefined number of classes and merging down, the final classified image also appeared more clinically meaningful than the image generated by prespecifying a fixed smaller number of classes (Supplementary Fig. S2).

Interpretation of the Spectral Signatures

Our study confirmed that tissue types in the retina have an identifiable spectral signature. In combination with spatial context, these signatures may aid in the detection and diagnosis of disease, including sub- or preclinical features (those preceding or not meeting standard clinical criteria for disease, e.g., dynamic changes in rhodopsin regeneration) or for tracking disease progression (Supplementary Fig. S3). Light delivered to the ocular tissues undergoes fluorescence, scattering, and absorption, primarily by lipofuscin, hemoglobin, melanin, and water. Macular pigment and choroidal vasculature in our series were never identified as separate spectral signatures, likely owing to the background correction step in preprocessing. An additional finding was that drusen were more often accurately discriminated than

![Image series of three eyes with large drusen in intermediate AMD suitably classified by using pattern recognition and a transformed divergence criterion of 1.5, or >85% probability of correct classification. (A) Separable large drusen appear as a single red-coded signature in the theme map. Note that large drusen sometimes colocalized with a range of spectral signatures. (B) An example where an individual druse appears to be encircled by a different perilesional spectral signature (transformed divergence separability = 1.77). (C) Drusen that appear similar in the fundus image (arrowheads) are separated into a number of drusen-related unique signatures and also encircled by a perilesional signature, similar to the eye in (B). (D) The three different drusen-related signatures of case (C) may reflect drusen diversity. Using optical coherence tomography, the drusen classes (blue, orange, and pink) differed in internal reflectivity and the overlying appearance of the inner segment ellipsoid zone. For the blue-coded druse, the overlying ellipsoid zone is minimally disturbed. The orange druse appears to abut the ellipsoid zone, while the pink druse causes a more frank discontinuity.](http://arvojournals.org/)
pigmentary abnormalities. This may relate to difficulties in grading and the characteristics of these different lesions in the component imaging: drusen appear bright on all modalities with a reflectance peak approximating 560 nm, while pigmentary abnormalities appear dark (due to absorption) in green light. Previous studies have also shown that melanin best absorbs light between 450- and 700-nm wavelength, though 450- to 600-nm range light may also be absorbed by other ocular components.

As illustrated in Figure 4, our method also appeared to discriminate subtypes of drusen, which may relate to the integrity of the overlying outer retinal layers and may therefore correlate with risk of atrophy. Our observations of unique perilesional spectral signatures may represent a limitation in image registration, that is, residual areas of kerf. Alternatively, it may suggest prescient extension of disease. For example, separable signatures lying just within the margins of drusen might reflect the indistinct border characteristics or mechanical distortion of the retina by soft drusen. Thus, we chose not to remove these signatures in the final analysis.

Study Limitations

As with all imaging, the methodology introduced in the foregoing sections should be viewed as adjunctive rather than a replacement of traditional imaging and funduscopy. This article reports on a proof-of-concept study and will benefit from future work using a replication sample of larger size. The three modalities (structural images) included were chosen for their ready commercial availability. The sensitivity of our technique might also be improved by varying the number and/or type of input channels (spectral bands) into the algorithm, such as en face images from dye studies, near-infrared FAF, scanning laser tomography, or a dedicated multispectral imaging device.

This study evaluated the appearance and spectral signature separability of typical intermediate AMD lesions (those that are hyperreflective in the input imaging) in Asian and Caucasian eyes only, and did not consider spatial information or lesions of other stages (reticular pseudodrusen, geographic atrophy, signs of neovascular AMD) or ethnicities. The “ground truth” was established via consensus opinion of clinical experts. Although those graders were masked to the diagnosis of the eyes, they were instructed to annotate each eye for lesions in accordance with the Beckman initiative for macular research clinical classification scale. Consequently, some bias was inevitable.

Finally, the application of the described technique to diseases or stages other than intermediate AMD was beyond the scope of this study. Late AMD was not evaluated. The present work described a case control diagnostic accuracy study, and the reported sensitivity and specificity values may indeed be lower in the presence of other confounding retinal diseases. The effect of lens status, for example, pseudophakia, may also introduce other variables, which will require clarification before more widespread adoption of the method. Further research designed as a cohort study applying the technique to patients with these potential confounders is currently underway.

![Figure 5](http://arvojournals.org/)

**Figure 5.** Case examples of three eyes with hyper- or hypopigmentary changes in intermediate AMD (arrowheads). (A) An eye with separable hyperpigmentary changes that appeared bright with a high pixel value on infrared and autofluorescence imaging, pink in the rgb triplet image, and a separate color, orange, in the theme map. (B) An instance of hyperpigmentary changes apparently convoluted by the underlying drusen changes. Although the changes appear similar to (A) in the input imaging, the final classified image shows the same signature/color as the background drusen. (C) Hypopigmentary changes marked by the arrowhead are separable, appearing red in the final classified image. As in Figure 3, additional high-resolution optical coherence tomography line scans have been provided for illustrative purposes.
CONCLUSIONS

This work represents a novel, proof-of-concept study demonstrating that multispectral, unsupervised pattern recognition can be used to classify fundus images in intermediate AMD and has applications in improving our understanding of phenotypes in AMD and identifying disease progression. This form of image analysis was able to accurately identify AMD lesions with high specificity and could ultimately be incorporated into a fully automated diagnostic tool or clinical decision-making support system suitable for AMD and other diseases, in non-specialized environments and telemedicine to facilitate the early and timely detection of eye disease.

Acknowledgments

The authors thank Erica Fletcher and Laura Downie for technical advice relating to pattern recognition, Robert Marc and Rebecca Pfeiffer for technical advice and access to the University of Utah image registration program (ir-tweak), Lindsay Moore and Elizabeth Wong for performing the AMD grading, and Tyson Xu for technical support.

Supported in part by an Australian Government Research Training Program scholarship and a National Health and Medical Research Council (NHMRC) Grant (No. 1033224). Guide Dogs NSW/ACT is a partner in the NHMRC grant and also provided a supplementary PhD scholarship for AL and support for LN-S.

Disclosure: A. Ly, P. L. Nivison-Smith, P. N. Assaad, None; M. Kalloniatis, P.

References


