A COMPARISON OF HIGH SPATIAL RESOLUTION IMAGES FOR FINE SCALE VEGETATION MAPPING

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ABSTRACT

A Comparison of High Spatial Resolution Images for Fine Scale Vegetation Mapping

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Recent advances in airborne and spaceborne sensors have made high spatial (≤1m/pixel) and spectral resolution images (e.g. IKONOS, SPOT 5, Quickbird 2) widely available, raising questions regarding their utility for floristic identification and classification. Additionally, the use of object-oriented software to perform automated classification and mapping has increased throughout the past 20 years. Studies assessing the utility of these image and software options frequently center on large, homogeneous sites and do not address these applications to small, heterogenous areas typical of the Pacific Northwest. In this study, a high-density sampling grid was used (approximately 9.0 % sample), followed by agglomerative cluster analysis and ordination, to identify all vegetation alliances and associations on a 148-ha study site in Maple Creek, California. Supervised classification using object-oriented software was performed on three images of various high spatial resolutions (0.15 m 4-band aerial photo, 0.60 m 4-band satellite image, and 1 m 3-band satellite image). The resulting classifications were compared with the reference vegetation map (derived from plot and image data) to assess accuracy. Results show differences in classification accuracy between the 3 images with the 0.60m Quickbird image producing the highest overall accuracy (69%); followed by the 0.15m aerial photo (48%); and the 1m NAIP image (37%) when assessed at the alliance level.

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1. Introduction

The use of remote sensing technology in vegetation classification and mapping has increased over the past 30 years (Environmental Systems Research Institute and The Nature Conservancy, 1994; Jensen, 2000; Greenberg et al., 2006). Remote classification reduces the need for field sampling and alleviates access constraints, making this technology a valuable tool for resource managers (Mullerova, 2004; Leckie et al., 2005; Johansen et al., 2007). Historically, remotely-based vegetation classification was performed via manual polygon mapping with air photos (Anderson et al., 1976; Hall, 2003; Sandmann and Lertzman, 2003) as these images offered distinctly higher spatial resolution or smaller pixel sizes (Benson and MacKenzie, 1995; Lillesand et al., 2004) than satellite imagery. Recent advances in spaceborne sensors, however, have made high spatial and spectral resolution satellite images (e.g. IKONOS, SPOT 5, Quickbird 2) widely available, raising questions regarding their utility for floristic identification and classification (Carleer and Wolff, 2004; Johansen et al., 2007). Additionally, vegetation mapping methods have expanded with the development of automated classification procedures which provide image enhancement and processing techniques not available with manual mapping practices (Lillesand et al., 2004).

Higher spectral resolution, or larger number of bands in a given sensor (Lefsky and Cohen, 2003), provides opportunities for discerning vegetation characteristics unobservable in natural color (Lefsky and Cohen, 2003; Lillesand et al., 2004). The very near infrared band, in particular, allows for the development of vegetation indices such as

a normalized difference (NDVI) to assist in live canopy detection (Asner et al., 2003). Higher band numbers, however, may increase scene 'noise' and image variance (Lillesand et al., 2004).

Although higher spatial resolution imagery (≤1 m) allows for clearer visualization of ground features (Lefsky and Cohen, 2003; Wulder et al., 2004) an increase in pixel number raises the internal variability within homogeneous land cover units, causing difficulties in class discernment (Carleer and Wolff, 2004). Intra-class variability issues are particularly problematic for per-pixel classifiers which rely solely on spectral signature values (Cushnie, 1987; Woodcock and Strahler, 1987; Lefsky and Cohen, 2003). Per-pixel software limitations, such as 'salt and pepper' effects where individual pixels are classified differently from their neighbors (Yu et al., 2006), have led many interpreters to believe these classifiers are not ideal for large heterogeneous units such as vegetation classes (Song et al., 2005; Yu et al., 2006). Exploration into spectral mixture analysis (Foody and Mathur, 2006) and object-oriented software, however, have shown promise in tolerating certain levels of variability (Yu et al., 2006) previously found to be problematic for per-pixel classifiers.

Consequently, object-oriented approaches have become popular for vegetation classification particularly when using higher-resolution images (Hay et al., 2005; Chubey et al., 2006). Object-based classifiers use aggregated groups of pixels, or image objects, to train the program to identify discrete entities normally recognizable to the human eye (Hay et al., 2005). In general, these programs group spatially adjacent pixels into spectrally homogeneous objects to then be used as minimum classification units

(Yu et al., 2006). Feature Analyst 4.1 (Visual Learning Systems, 2006) is a popular object-oriented commercial software package in use today that utilizes an artificial neural network classifier (Ripley, 1996) to consider spatial attributes, such as spatial association and image texture, when performing automated feature extraction (Vanderzanden and Morrison, 2003). Object-oriented classifications do not, however, always reach the commonly recommended 80 - 85% accuracy standard (Environmental Systems Research Institute and The Nature Conservancy, 1994; Congalton and Green, 1999; Foody, 2002; USDA, 2002) especially when analyzing fine scale, floristically-based categories (e.g. alliance or association). Thus, automated software users often broaden their class scales to reach this accuracy standard, resulting in coarser, homogeneous units that may not adequately reflect the heterogeneity typical of many second-growth forests (Spies et al., 1994; Jiang et al., 2004).

Furthermore, because access and sampling issues are magnified in larger areas, studies assessing the utility of remotely-driven classification frequently center on extensive sites such as entire national forests or parks (Jiang et al., 2004; Greenberg et al., 2006; Yu et al., 2006), or large private holdings (Spies et al., 1994). Limited ground sampling in these large areas for training and validation purposes (Environmental Systems Research Institute and The Nature Conservancy, 1994; Jiang et al., 2004; Leckie et al., 2005) generally results in low-confidence reference data (Foody, 2002). In this study, however, the small scale of the site afforded a unique opportunity for comprehensive ground coverage and generation of high-confidence reference information.

In addition to image resolution and classification method considerations, image type (air photo vs. satellite) may play a role in accuracy (Lefsky and Cohen, 2003). Satellite images avoid many of the problems associated with aerial photography such as join lines in mosaicked scenes, non-standardized flight orientations, or difficulty in acquiring multidate data sets (Lefsky and Cohen, 2003; Chubey et al., 2006). Spaceborne platforms can be limited, nonetheless, by inflexibility of satellite orbit schedules, as well as increased atmospheric effects due to greater sensor-to-ground distance (Lefsky and Cohen, 2003). Perhaps one of the chief issues regarding image type is the associated cost incurred with each of these options. When dealing with a small site, the price per acre of imagery may be increased as there are frequently minimum area requirements for scene purchases. Acquiring timely aerial photographs can also be prohibitively expensive due to the cost of chartering flights. Commercial satellite imagery ranges in price from \$1 – 22/km² (Yildirim and Seker, 2004) however government subsidized programs such as Landsat and NAIP (National Agricultural Imagery Program) offer free imagery.

There are numerous decisions land managers need to make when selecting appropriate methods for classifying vegetation such as: type of image (air photo vs. satellite), image resolution (spatial and spectral), and mapping method (automated vs. manual). These decisions need to be made in the context of the manager's objectives (e.g. habitat mapping vs. timber inventory). Research assessing these image and software options will provide land managers with information regarding the utility of using high-resolution imagery coupled with object-oriented classification software when attempting to classify vegetation. The specific objectives of this study were to (1)

classify and map vegetation on a small, heterogeneous forested landscape, (2) compare the accuracy of a visual and field-based classification method with an object oriented classification method (Feature Analyst), and (3) compare the accuracy of classifying vegetation using three high spatial resolution (0.15, 0.60, and 1.0 m) digital images.

2. Materials and methods

2.1. Study area

This study was conducted on the L.W. Schatz Demonstration Tree Farm located in Maple Creek, California (Fig. 1). The 148-hectare site extends from 40°46'49" N to 40°45'56" N latitude and 123°52'21" W to 123°51'32" W longitude (T 5N, R 3E, Section 32). It ranges in elevation from 140 to 430 m, and is underlain by the Franciscan Formation, a subduction complex consisting of accreted fragments of oceanic crust and forearc sediments (Aalto and Harper, 1989).

Originally consisting of old-growth *Pseudotsuga menziesii* forest, the land was logged in the early 1950's (Schatz 2007, personal communication). It has since experienced a mixture of natural recovery and management resulting in a heterogeneous landscape mosaic typical of many northwestern forests today (Halpern and Spies, 1995).

Current vegetation includes more than 150 species (Appendix A) and is dominated by a *P. menziesii* and mixed hardwood overstory with an understory of abundant evergreen shrubs and ferns. In addition to the forested areas, the study site contains upland prairie and a transmission right-of-way where both native and non-native perennial grasses dominate.

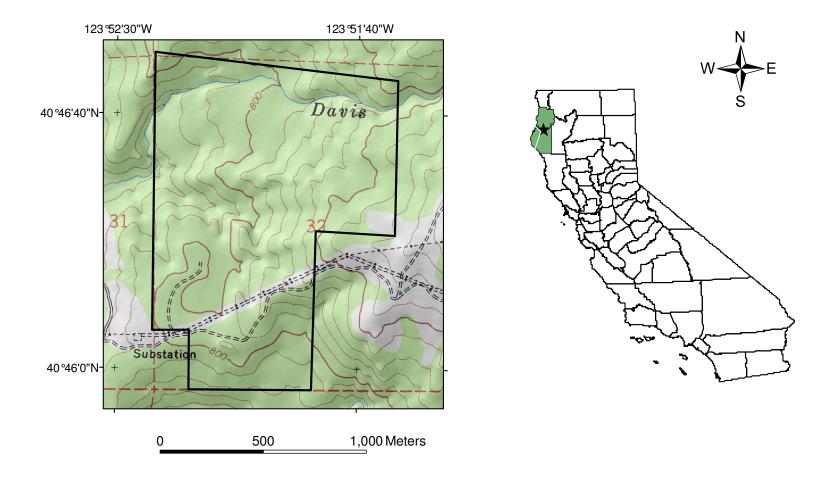


Figure 1. Location and extent of study area (outlined in black) consisting of most of the W ½ of Section 32 (T 5N, R3E), 17 miles east of Eureka, California.

2.2. Field sampling of vegetation

A high-density sampling grid scheme of 2 plots per hectare was chosen to achieve sufficient and balanced coverage of vegetation types (Cooper et al., 2006). A circular plot shape was employed for all non-riparian plots. Rectangular plots were used in riparian corridors to characterize floral composition accurately (California Native Plant Society, 2004). The maximum number of plots based on 2/ha was 296. However, the number sampled was 274 after eliminating plots overlapping the property boundary and those occurring on landslides and overly steep terrain. Two hundred and forty-eight plots were on upland, tree-dominated terrain (0.05 ha, 12.6 m radius); 8 plots were in riparian zones (0.05 ha, various dimensions); and 18 plots were in shrub- or herb-dominated areas (0.02 ha, 8.0 m radius). The plots covered approximately 9% of the tree farm.

Field data were collected from June-August, 2006 using a modified rapid assessment protocol (California Native Plant Society, 2004). Vegetation was sampled using relevé plots and modified Braun-Blanquet cover abundance scaling (Table 1; Braun-Blanquet, 1932; Lee, 2004). Ocular estimates of plant cover by species (within plot or outside of plot but providing cover within the plot) were recorded for all strata, as were average height (m) and total percentage cover by stratum (Mueller-Dombois and Ellenberg, 1974). Abiotic information recorded included: elevation (m), topographic position/landform, percent slope, aspect, and soil type (Colwell, et al., 1960).

Table 1. Cover abundance scale used in ocular estimates (Lee, 2004).

Cover Range (%)
0.001 - 0.01
0.01 - 0.1
0.1 - 1
1 - 5
5 - 15
15 - 25
25 - 50
50 - 75
75 - 100

2.3. Classification and Ordination

Vegetation data were analyzed using several multivariate approaches. Species with less than 1% total cover on the site were removed *a priori* to prevent outlier effects (McCune and Grace, 2002). Riparian plots were included in the overall analysis due to similarities in dominant vegetation with non-riparian units. Data were grouped into possible plant associations using a hierarchical clustering algorithm (Euclidean distance, Ward's linkage method) contained in PC_ORD (McCune and Grace, 2002). This method merges individual plots into groups based on species similarity. Resultant groups were then pared down through indicator analysis. Data were further analyzed via comparisons of species abundance and constancy within and between groups. A Nonmetric Multidimensional Scaling ordination was performed to further reduce the data set and graphically depict ecological relationships among plots.

The naming convention in *A Manual of California Vegetation* (Sawyer and Keeler-Wolf, 1995) was used when assigning plots to alliance and associations. Dominant species were defined as those having $\geq 50\%$ relative cover and frequency across all plots within a vegetation type.

2.4. Image Acquisition and Processing

Three images were acquired for analysis (Fig. 2). One multispectral, 4-band (0.45 $-0.90 \, \mu m$) airborne image with 0.15 m spatial resolution was acquired on June 22, 2006 for use in reference map creation and automated feature extraction. A multispectral,

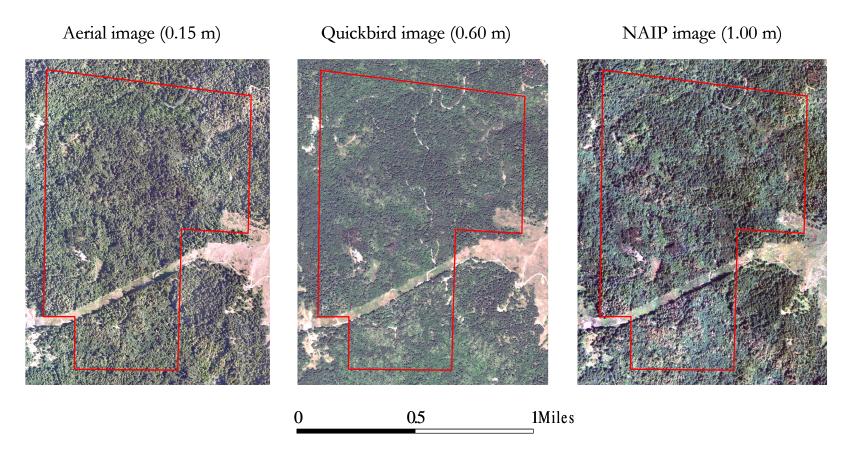


Figure 2. Digital images (and associated pixel resolution) used in analysis. Aerial image acquired on June 22, 2006, Quickbird image acquired July 28, 2006, NAIP image acquired June 15, 2005. The outline of the study site appears in red.

4-band $(0.45-0.90~\mu m)$ QuickBird satellite image of 0.6m spatial resolution acquired July 28, 2006 by Digital Globe, and a natural color 3-band $(0.45-0.69~\mu m)$ NAIP 1 m spatial resolution aerial image acquired June 15, 2005 by U.S. Department of Agriculture Farm Services Agency were used in automated feature extraction only.

The QuickBird image was orthorectified in ArcMap using a 10 m digital elevation model and rational polynomial coefficients (Barbarella et al., 2004). All other images were georeferenced and orthorectified by the provider. Image radiances were not atmospherically corrected (Lillesand et al., 2004) as times series analysis of consecutive image data was not required for this study and detailed information on the atmospheric conditions at the time of overpass was not available.

2.5. Vegetation Mapping

Vegetation polygons were visually interpreted and digitized in ArcMap using the 0.15 m resolution aerial image and labeled to alliance and association. The minimum mapping unit (MMU) was 1000 m² (approximately 2 tree-dominated ground plots) and was chosen in order to retain detail while capturing stand-level characteristics (Stohlgren et al., 1997). Vegetation polygons were laid over the remaining two images and adjusted to account for shifts in images due to error, orthorectification or georeferencing limitations. Polygons not containing at least one ground plot were verified in the field.

2.6. Automated Feature Extraction

Supervised classification of imagery was performed via automated feature extraction using Feature Analyst (Visual Learning Systems 2006) software. Training

polygons were randomly selected from the reference map using Spatial Analyst in ArcGIS 9.1 to a standard of 5 or fewer polygons (approximately 40%) per vegetation type. All vegetation classes were included in association-based classification. Vegetation classes covering < 1% of the study area or those with only one polygon were masked in the alliance-based classification. Selected input parameters included preaggregation to 500 pixels and a representation pattern appropriate for stand-level classes (Fig. 3; Vanderzanden and Morrison, 2003). All vegetation data were used in map validation.

One classification using all available spectral bands was executed with the association reference map. Several classifications were executed with the alliance reference map. Band selection methods for alliance map analyses included: use of all available spectral bands, removal of the VNIR band in the Quickbird and aerial images, application of an NDVI to the Quickbird and aerial images, and application of only the spectral bands exhibiting the best average separability as determined by a transformed divergence statistic (Swain and Davis, 1978; ERDAS IMAGINE, 2005). Overall, the greater the transformed divergence, the greater the statistical distance between training classes and the higher the probability of correct classification (Lillesand et al., 2004). In general, if a result is greater than 1,900 then the classes can be separated; between 1,700 and 1,900 the separation is fairly good. Below 1,700 the separation is poor (Jensen, 2000).

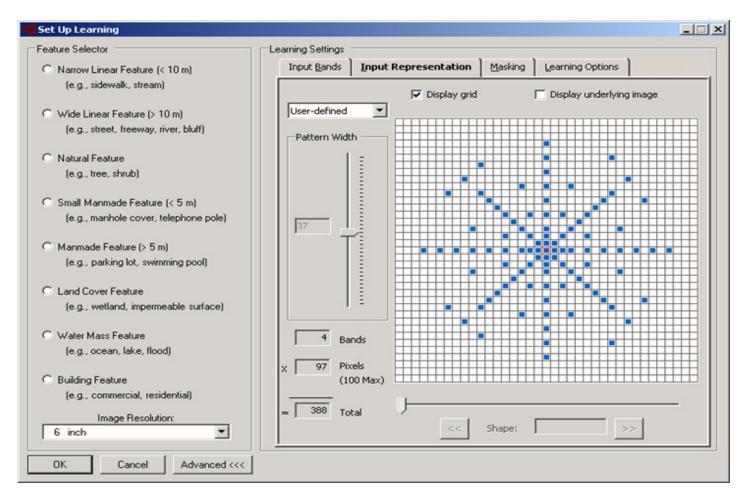


Figure 3. Feature Analyst input pattern representation. Pixels shown in blue represent those included in analysis.

To keep spatial resolution constant and test for an effect from the infrared band, the Quickbird and aerial images were resampled to 1m, and all images were run with all available spectral bands.

2.7. Accuracy Assessment

Correct vs. incorrectly classified polygons were identified in the output maps using zonal statistics in Spatial Analyst (Environmental Systems Research Institute, 2006). If the majority of pixels in the output polygon matched the reference polygon then the entire polygon was identified as correct. A majority assessment rule was chosen to account for the inherent heterogeneity within vegetation classes (Sawyer and Keeler-Wolf, 1995) and because more traditional (per-pixel) assessments of classification accuracy are often inappropriate for use in these systems (Yu et al., 2006).

Map accuracies are presented in the form of an 'error matrix' which includes measures of producer's, user's, and overall accuracy (Story and Congalton, 1986; Foody, 2002). Producer's accuracy, or error of omission, indicates the probability of a reference sample being correctly classified by the software. User's accuracy, or error of commission, indicates the probability that a pixel classified on the map represents that category on the ground (Congalton and Green, 1999). Kappa coefficients, K_{hat}, were calculated for overall map accuracy to compensate for random chance agreement (Rosenfield and Fitzpatrick-Lins, 1986).

3. Results

Twenty-five vegetation associations (Table 2; Fig. 4) and 13 vegetation alliances were classified (Table 3; Fig. 5) and mapped. *Sequoia sempervirens* plantations were included as classes in the association map as they were subcanopy dominant. *Pinus* sp. plantations were included as classes in the alliances as they were canopy dominant. The study site is primarily dominated by *P. menziesii - Abies grandis, Alnus rubra,* and *P. menziesii - Lithocarpus densiflorus* classes. Species richness per alliance varied from a low of four in a *Baccharis pilularis* alliance to a high of 21 in a *Fraxinus latifolia* riparian alliance (Table 3).

The output association maps (Fig. 6) had very low overall accuracies across all images. The Quickbird output had the highest overall accuracy (14%), followed by the NAIP (11%), and the air photo (3%). When using only 3 bands per image (R, G, B) the Quickbird output accuracy decreased to 3% while the air photo output accuracy increased to 7%.

Output maps of alliances (Fig. 7) had much higher overall accuracies across image types (Table 4) than association output maps. The Quickbird output was most accurate (69%, K_{hat} = 0.43), followed by air photo (48%, K_{hat} = 0.33), and lastly NAIP (37%, K_{hat} = 0.28) when all available bands were included in the analysis (Table 4, column a). The air photo and Quickbird alliance maps decreased in overall accuracy after removal of the infrared spectral band (Table 4, column b). Analyses using bands with the highest spectral separation had mixed results. The Quickbird and air photo accuracies decreased while the NAIP accuracy increased by 4% (Table 4, column c).

Table 2. Relative cover, frequency, and species richness of associations.

Associations	Relative	Number	Mean species	Standard
	cover (%)	of	richness	deviation of
		polygons		species richness
Umbellularia californica - Alnus rubra	12.6	16	16	5
Lithocarpus densiflorus	12.1	19	13	4
Umbellularia californica - Pseudotsuga menziesii	10.3	21	20	4
Pseudotsuga menziesii - Abies grandis	8.4	11	22	5
Lithocarpus densiflorus - Alnus rubra - Umbellularia californica	7.7	7	15	5
Pseudotsuga menzieseii/Rubus ursinus	6.9	11	19	7
Pseudotsuga menziesii/Polystichum munitum	6.5	9	20	3
Abies grandis - Pseudotsuga menziesii	6.3	15	20	4
Pseudotsuga menziesii - Lithocarpus densiflorus	6.1	12	21	6
Lithocarpus densiflorus - Umbellularia californica - Pseudotsuga				
menziesii	3.9	10	14	4
Alnus rubra	3.7	11	18	4
Transmission right-of-way	2.5	3	13	3
Tsuga heterophylla	2.1	1	9	3
Pseudotsuga menzieseii/Ceanothus thyrsiflorus	2.1	4	14	1
Lithocarpus densiflorus - Sequoia sempervirens	2.0	4	15	5
Pinus ponderosa	1.2	2	21	0
Introduced perennial grasslands	1.2	3	14	1
Pseudotsuga menziesii - Sequoia sempervirens	0.9	2	21	7
Acer macrophyllum	0.9	5	23	6
Fraxinus latifolia	0.7	5	21	2
Salix sp.	0.7	4	22	11
Ceanothus thyrsiflorus	0.6	3	18	2
Pinus radiata	0.3	1	24	0
Baccharis pilularis	0.1	2	4	0
Quercus garryana	0.1	1	17	0

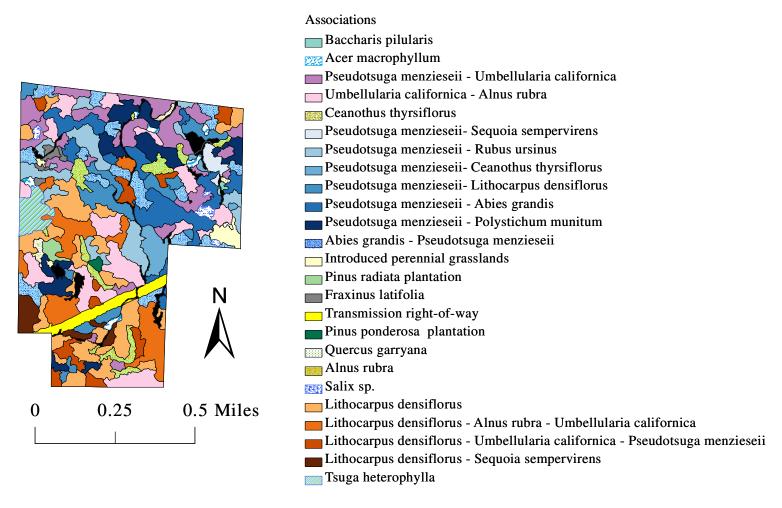


Figure 4. Thematic map showing 25 vegetative associations on study site. Areas in black were masked as non-vegetation during analysis.

Table 3. Relative cover, frequency, and species richness of alliances (*indicates vegetation classes used in automated feature extraction).

Alliances	Relative	Number of	Mean species	Standard
Amances	cover (%)	polygons	richness	deviation of
	COVCI (70)	polygons	Heimess	species
				richness
Pseudotsuga menziesii - Abies grandis*	28.1	26	21	4
Alnus rubra - Umbellularia californica - Lithocarpus densiflorus*	24.7	19	16	4
Lithocarpus densiflorus - Umbellularia californica - Pseudotsuga menziesii*	19.7	16	14	5
Pseudotsuga menziesii - Umbellularia californica*	13.0	19	20	4
Pseudotsuga menziesii*	6.9	14	19	6
Introduced perennial grassland*	3.6	6	13	2
Pine plantation*	1.0	2	21	4
Fraxinus latifolia	0.8	5	21	2
Acer macrophyllum	0.7	6	23	6
Salix sp.	0.7	4	22	11
Ceanothus thyrsiflorus	0.6	3	18	2
Baccharis pilularis	0.1	2	4	0
Quercus garryana	0.1	1	17	0

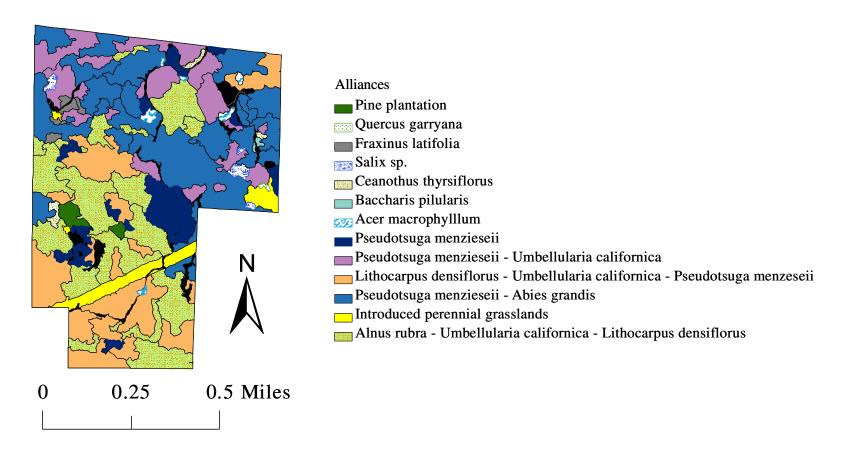


Figure 5. Thematic map showing seven vegetative alliances used in classification. Areas in black were masked as non-vegetation during analysis.

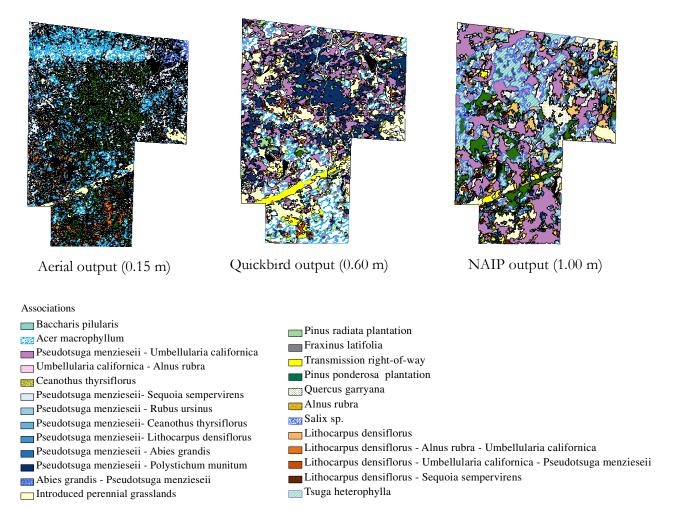


Figure 6. Feature analyst output association maps by image type. Areas in black were masked as non-vegetation during analysis.

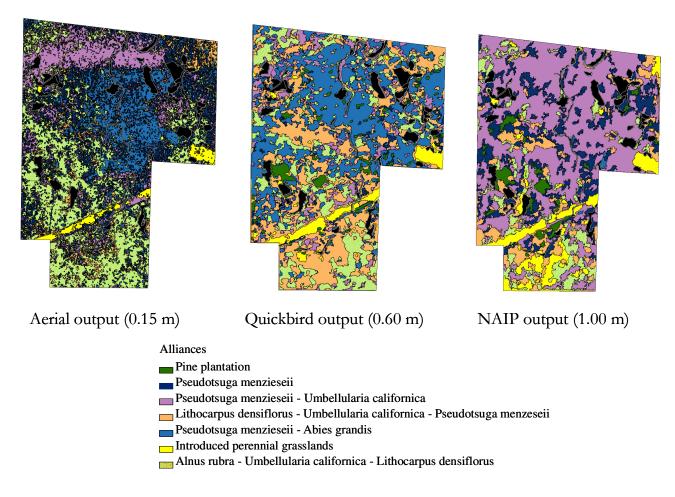


Figure 7. Feature analyst output alliance maps by image type. 7 vegetative alliances used in classification are shown. Areas in black were masked as non-vegetation during analysis.

Table 4. Overall alliance output map accuracies by image type and analysis. Band numbers used in analysis in parentheses.

		VNIR removed		
Image Spatial Resolution	All available bands a.	(3-band) b.	Select bands c.	NDVI d.
NAIP (1.00m)	0.37 (1,2,3)	n/a	0.41 (2,3)	n/a
Quickbird (0.60m)	0.69 (1,2,3,4)	0.60 (1,2,3)	0.47 (2,3)	0.54 (1,2,3,4,5)
Air photo (0.15 m)	0.48 (1,2,3,4)	0.27 (1,2,3)	0.33 (1,3,4)	0.37 (1,2,3,4,5)

The addition of the commonly used NDVI also did not improve overall accuracy (Table 4, column d). Overall accuracies decreased for the Quickbird and aerial images when resampled to 1m, implying a positive influence on accuracy from the VNIR band (Fig. 8). The highest overall accuracies for the Quickbird and aerial output maps were produced through the inclusion of all 4 bands and no further manipulation. The highest overall accuracy for the NAIP output map (41%) was produced through the removal of band 1.

The error matrices for each image (Table 5) reveal that the pine plantation and introduced perennial grassland classes generally had the highest user and producer accuracies. The pine plantations were misclassified, however, as *A. rubra* in the aerial output map. There was little consistency with other class accuracies or relative cover values (Table 6) across image types. The *A. rubra* dominant class had high user and producer accuracies but was confused with the *P. menziesii – Umbellularia californica*. The *P. menziesii – U. californica* and *P. menziesii – L. densiflorus* classes performed differently across images, however the *P. menziesii – L. densiflorus* class had relatively high user accuracies suggesting that, when identified, these classes were labeled appropriately. The *P. menziesii* class had the lowest producer accuracies (5-36%) and was frequently confused with the mixed *P. menziesii – A. grandis* class.

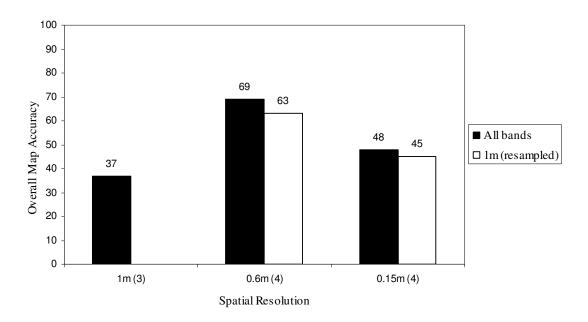


Figure 8. Overall accuracies of original and images resampled to 1 m. Number of spectral bands in parentheses.

Table 5. Error matrices for alliance map classification results by image type. Values represent area (m²) classified. Vegetation classes for the L.W. Schatz study site include: (1) Alnus rubra - Umbellularia californica - Lithocarpus densiflorus (A-Uc-Ld); (2) Pseudotsuga menziesii (Pm); (3) Pseudotsuga menziesii - Umbellularia californica (P -Uc); (4) Pseudotsuga menziesii - Abies grandis (Pm-Ag); (5) Lithocarpus densiflorus - Umbellularia californica - Pseudotsuga menziesii (Ld-Uc-Pm); (6) Introduced perennial grasslands (IPG); and (7) Pine plantation (P).

Classified data	Refer	ence data						
Class	A-Uc-Ld	Pm	Pm-Uc	Pm-Ag	Ld -UcPm	IPG	P	User (%)
(1) Classification e						11 0		(70)
(1) Classification c	TIOI IIIIIIIX IO	i the acria	i piloto (o.i	is iii, i ouii	u)		1112	
A-Uc-Ld	294645	21856	66020	49650	182548	0	6	47
Pm	0	4511	0	30454	0	0	0	13
Pm-Uc	6606	12884	74008	145180	0	0	0	31
Pm-Ag	43939	59141	37805	192590	52480	0	3750	49
Ld-Uc-Pm	0	0	0	2	43069	0	0	100
IPG	0	0	0	0	0	54014	0	100
P	0	0	0	0	0	0	0	0
Producer (%)	85	5	42	46	15	100	0	
Overall (%)		-						48
K _{hat}								0.33
(2) Classification e	error matrix fo	r the Quic	kbird imag	e (0.60 m. 4	l-band)			
A-Uc-Ld	248618	0	47270	2	0	0	0	82
Pm	0	5707	0	0	0	0	0	100
Pm-Uc	0	0	12261	14645	0	0	0	46
Pm-Ag	68084	29170	68856	348958	0	595	0	69
Ld-Uc-Pm	28489	61969	49446	54271	278098	0	0	59
IPG	0	0	0	0	0	53419	0	100
							1487	
P	0	1545	0	0	0	0	6	90
Producer (%)	70	6	7	85	100	99	100	
Overall (%)								69
K _{hat}								0.43
(3) Classification e	error matrix fo	r the NAII	P image (1.	0 m, 3-band	d)			
A-Uc-Ld	68111	3821	0	0	0	0	0	95
Pm	658	35886	3777	48625	15961	0	0	34
Pm-Uc	275181	50215	174056	369251	40984	0	0	19
Pm-Ag	0	0	0	0	0	0	0	
Ld-Uc-Pm	0	6927	0	0	173858	982	0	96
IPG	1240	0	0	0	47295	52429	0	52
							1487	
P	0	1544	0	0	0	603	6	87
Producer (%)	20	36	98	0	63	97	100	
Overall (%)								37
K _{hat}								0.28

Table 6. Relative cover values of alliances for Feature Analyst output maps by image type.

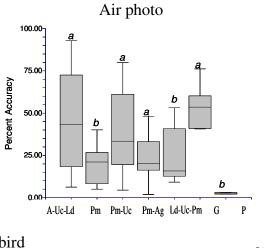
Alliance	Aerial image	Quickbird image	NAIP image	Reference map
Pseudotsuga menziesii - Abies grandis	24.0	33.1	0.3	28.1
Alnus rubra - Umbellularia californica - Lithocarpus densiflorus	30.4	17.1	9.0	23.7
Lithocarpus densiflorus - Umbellularia californica - Pseudotsuga menziesii Pseudotsuga menziesii - Umbellularia	6.9	31.0	11.8	18.7
californica	20.8	9.4	50.8	12
Pseudotsuga menziesii	14.5	1.8	16.9	6.6
Introduced perennial grassland	3.3	4.3	8.1	3.6
Pine plantation	0.2	3.3	3.2	1

When the *P. menziesii* and *P. menziesii* – *A. grandis* classes were combined into a "mixed conifer" class the overall accuracies of the air photo, Quickbird, and NAIP images increased to 54%, 72%, 41%, respectively.

Differences in vegetation class accuracies were evaluated using a non-parametric Kruskal-Wallis ANOVA procedure, followed by Kruskal-Wallis Z and Bonferroni correction. Significant differences (p <0.05) between the accuracies of vegetation types for each image were detected (Figure 9).

3.1. Sources of Error

Spectral separation of classes was calculated for all three images using the transformed divergence statistic (Swain and Davis, 1978; ERDAS IMAGINE, 2005). Results showed that all images did not have ideal spectral separation implying spectral overlap between classes. The Quickbird data had the best statistical separation between categories. All images had similar minimum values. In general, the classes with the highest producer and user accuracies had the highest spectral separation (Appendix B). The introduced perennial grass class was the most spectrally distinct with the highest transform divergence values across all images (air photo = 995, Quickbird = 1432, NAIP = 503 average TD). Classes that were confused with each other typically had transformed divergence values <100 but were not always those with the highest spectral overlap. For example, in the aerial image the pine plantation class was confused with the *Alnus rubra - Umbellularia californica - Lithocarpus densiflorus* class (TD = 227) and not the *Pseudotsuga menziesii* class (TD = 17).



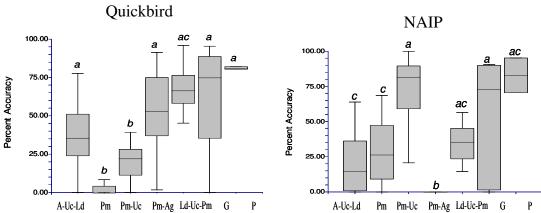


Figure 9. ANOVA test for differences in vegetation class accuracy across image types. Different letters above the bars indicate significant differences in Tukey post-hoc comparisons between the classes. Vegetation codes are defined in Table 5.

Increasing internal variability in vegetation classes is correlated with decreasing classification accuracy (Cushnie, 1987; Cochrane, 2000). Higher intra-class variability lessens the separation between classes, making thematic distinction difficult. The average standard deviations of the digital numbers within vegetation types revealed differences across images. The air photo has the most intra-class variability (31.1) which may be due to the high number of pixels. It did not, however, have the lowest accuracies when all 4 bands were used, implying an important contribution from the VNIR band. The Quickbird image had the least amount of intra-class variance (11.2) and the highest accuracies.

Although factors such as polygon size, total vegetation coverage, and number of training polygons have been shown to influence accuracy (Foody et al., 1995; Foody and Arora, 1997; Staufer and Fischer, 1997; Ellis and Wang, 2006; Foody and Mathur, 2006), logistic and linear regressions of these variables did not result in significant findings with the exception of the NAIP image where producer accuracy increased as class coverage decreased (p = 0.03). This relationship is exemplified in the pine plantation high accuracies despite low cover (1%) and low number of training polygons (2).

4. Discussion

In all cases, the Quickbird image produced the highest levels of overall accuracy when compared to the air photo and the NAIP image. Image processing techniques, such as resampling, inclusion of an NDVI, selection of bands with the highest transformed divergence values, or use of natural color bands (R, G, B) alone, did not alter this outcome. These techniques did, however, generally lower the overall and class accuracies of the Quickbird and air photo. The highest overall accuracies for these two images were produced using all available bands. The highest accuracy for the NAIP image, however, resulted from the removal of the blue band (450 – 520 nm). Blue bands can add unwanted noise to an image oftentimes due to atmospheric scattering (Kimes et al., 2006). The NDVI did not produce higher accuracies since this index generally compensates for terrain effects such as surface slope, aspect, or elevation, (Lillesand et al., 2004) and these factors were not significantly different between vegetation types. Additionally, the addition of the NDVI may have also added unwanted noise through the inclusion of redundant reflectance values.

What accounts for these differences in image performance? Generally speaking it results from the software's ability to discriminate between classes. Spectral separation and internal variability of classes were predominant factors influencing class discernment (Cushnie, 1987; Cochrane, 2000). An increase in intra-class variability causes a reduction of statistical separability between classes (Cushnie, 1987; Yu et al., 2006).

The Quickbird image had the highest overall transformed divergence values, the lowest intra-class variability, and the highest accuracies. This pattern was also found with the pine plantation and introduced perennial grass vegetation types. High accuracies occurred in these classes despite their low percentage of vegetative cover, as well as small number of training polygons (Foody, 2002; Chubey et al., 2006) suggesting that spectral separation supersedes these other variables in effect.

Another mechanism behind image performance relates to spatial resolution. It can be difficult to assess the appropriate threshold for spatial resolution. Too fine a resolution may increase the internal variability within homogeneous land cover units too much (Woodcock and Strahler, 1987; Aplin et al., 1997; Greenberg et al., 2006; Yu et al., 2006). At too coarse a resolution a number of vegetation types may be combined within one pixel, resulting in a spectrally noisy signal and yielding poor classification results (Cushnie, 1987; Yu et al., 2006; Johansen et al., 2007). The results of the 3-band input trials imply that a 0.6 m spatial scale may be best suited for alliance-level classification, followed by 1 m and finally 0.15 m. The higher resolution aerial photo may aid in visual interpretation, but it is not a 'best fit' when using automated classifiers.

Accuracies less than 80% may be due to land management history and to the early seral stages in this landscape (Jiang et al., 2004; Lu, 2005; Rapp et al., 2005). My study site was a heterogeneous mosaic of young forest, plantation, and remnant old-growth trees typical of post-logged coniferous forests in the Pacific Northwest today (Spies et al., 1994; Jiang et al., 2004). Due to fragmentation of the landscape and resultant growth patterns, vegetation classes were rarely composed of solely one or two dominant species,

such as a pine or grass, but were instead a mix of conifer and hardwood species of varying ages and sizes. For example, some classes defined by the Federal Geographic Data Committee (1997) have differences of 10% tree cover in some of their alliances which would not consistently produce a significant change in spectral signal (Greenberg et al., 2006). Dissimilarities in structural attributes of forest stands may have a greater effect on reflectance characteristics than tree species composition (Lefsky and Cohen, 2003). Additionally, evergreen forest types have been shown to lack unique spectral reflectance characteristics due to limited phenological differences (Lillesand et al., 2004). In my study the most ecologically mixed classes tended to be the most confused. Although some interpreters emphasize the importance of spectrally 'pure' classes, there may be more inherent variability within one tree crown than between species or classes (Leckie et al., 1992). Thus, prioritizing class homogeneity may produce ecologically meaningless polygons.

Categorical scale is another important factor in classification (Marceau et al., 1994; Ju and Gopal, 2005; Rapp et al., 2005). The map of associations had extremely low accuracies likely due to large number of input classes and the presence of subcanopy vegetation (Greenberg et al., 2006). To date, LIDAR data has been most effective in directly measuring three-dimensional distributions of plant canopy and sub-canopy (Lefsky and Cohen, 2003) thus a laser-altimetry tool may be more suited for this level of detail. Other studies (Greenberg et al., 2006; Yu et al., 2006) have generally shown low accuracies when attempting to classify alliances because of the influence of sub-canopy vegetation, small sample sizes, and species co-dominance.

Additional issues affecting accuracy not directly explored in my study include positional error of GPS and imagery (Foody, 2002), temporal differences in image acquisition (Lefsky and Cohen, 2003), join lines in mosaicked aerial photos (Lefsky and Cohen, 2003), as well as accuracy assessment method (Gopal and Woodcock, 1994; Stehman, 1997; Foody, 2002; Liu et al., 2007).

Overall accuracies were somewhat low with fine scale classifications, and only one image approached the commonly used 80% accuracy threshold (Environmental Systems Research Institute and The Nature Conservancy, 1994). The software produced the highest accuracies when utilizing all available bands, thus removing the need for image enhancement or other interpreter-based processing techniques (Hay et al., 2005).

The specific algorithms employed by Feature Analyst software included a type of neural network classifier that considers spatial context, or image 'texture', in addition to the brightness values of the pixels (Visual Learning Systems, 2006). Neural network classifiers have been successful in supervised classification of community data owing in part to their non-parametric nature which is appropriate for non-normally distributed data (Černá and Chytrý, 2005), as well as their ability to learn by example and generalize (Foody and Arora, 1997). Although artificial neural networks have given accurate class predictions compared to other supervised methods they are also accused of having a 'black-box' approach (Černá and Chytrý, 2005) which hides the underlying process. Foody and Arora (1997) found that neural network classifiers performed well on small training sets, however this software could not resolve classes in the alliance map that were represented by only 1 polygon, regardless of size. Therefore, types that are 'rarer'

such as *Tsuga heterophylla*, could not be identified through this process, and, it can be difficult to have enough polygons for training on such a small scale.

Additionally, the input representation pattern employed was limited in pixel number (100 per band). Therefore, less area was covered with each pass in the higher resolution images. This resulted in certain features of distinct shape, such as the electrical right-of-way, being misclassified in the association map. Thus, processing time would be extended by the necessity of running several passes or employing different input patterns to extract various shapes.

5. Conclusions

Object-oriented software may not be the ideal method to achieve high accuracy for alliance-level classification and mapping. Automated feature extraction has been shown to perform better in more homogeneous landscapes or at coarser categorical scales. To date, remote sensing is not as accurate and precise in measuring vegetation that an investigator utilizing manual and field-based mapping techniques can achieve (Greenberg et al., 2006). Thus, choosing this method of classification should be weighed against considerations such as cost and time limitations.

Spatial and spectral resolutions are important parameters to consider when choosing imagery for vegetation mapping. It is wise to use imagery at a spatial resolution that is appropriate for both the features being classified (Woodcock and Strahler, 1987) and the method of classification. In short, higher spatial resolution does not always produce the highest map accuracies with automated methods, and lower spatial resolution is not ideal for manual mapping. Additionally, the processing time associated with each image type should be considered as larger file sizes may lead to cumbersome hardware demands.

More work is needed to explore the utility of very high spatial resolution imagery in the field of vegetation classification and mapping. As satellites continue to offer finer pixel grains and additional spectral bands, they are fast becoming a viable choice over aerial imagery which is frequently cost-intensive. Additionally, more studies classifying second-growth heterogeneous vegetation to fine categorical scales are needed to

adequately assess the application of object-oriented classification procedures on sites reflecting the changing landscape.

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Appendix A. Vascular plant species found in study location. Nomenclature follows the USDA PLANTS Database (USDA, 2007).

Family	Species	Common name		
Aceraceae	Acer macrophyllum Pursh	bigleaf maple		
Aceraceae	Acer circinatum Pursh vine maple			
Anacardiaceae	Toxicodendron diversilobum (Torr. & Gray) Greene	Pacific poison-oak		
Apiaceae	Daucus carota L.	wild carrot		
Apiaceae	Heracleum maximum Bartr.	cow parsnip		
Apiaceae	Osmorhiza chilensis Hook. & Arn.	mountain sweet cicely		
Apiaceae	Sanicula crassicaulis Poepp. ex DC.	Pacific blacksnakeroot		
Araliaceae	Aralia californica S. Wats.	elk clover		
Aristolochiaceae	Asarum caudatum Lindl.	wild ginger		
Asteraceae	Achillea millefolium L.	common yarrow		
Asteraceae	Adenocaulon bicolor Hook.	American trailplant		
Asteraceae	Baccharis pilularis DC.	coyotebrush		
Asteraceae	Cirsium vulgare (Savi) Ten.	bull thistle		
Asteraceae	Erechtites minima (Poir.) DC.	coastal burnweed		
Asteraceae	Gnaphalium sp. L.	cudweed		
Asteraceae	Hieracium albiflorum Hook.	white hawkweed		
Asteraceae	Leucanthemum vulgare Lam.	oxeye daisy		
Asteraceae	Madia elegans D. Don ex Lindl.	common madia		
Asteraceae	Petasites palmatus (Ait.) Gray	coltsfoot		
Asteraceae	Taraxacum officinale G.H. Weber ex Wiggers	dandelion		
Berberidaceae	Achlys californica Fukuda & Baker	California deer-foot		
Berberidaceae	Mahonia repens (Lindl.) G. Don	Oregon grape		
Berberidaceae	Vancouveria hexandra (Hook.) Morr. & Dcne.	white insideout flower		
Betulaceae	Alnus rubra Bong.	red alder		
Betulaceae	Corylus cornuta var. californica (A. DC.) Sharp	California hazelnut		
Blechnaceae	Blechnum spicant (L.) Sm.	deer fern		
Boraginaceae	Plagiobothrys sp. Fisch. & C.A. Mey.	popcorn flower		
Caprifoliaceae	Lonicera hispidula (Lindl.) Dougl. ex Torr. & Gray	honeysuckle		
Caprifoliaceae	Sambucus racemosa L.	red elderberry		
Caprifoliaceae	Symphoricarpos albus (L.) Blake	common snowberry		
Caryophyllaceae	Stellaria media (L.) Vill.	common chickweed		
Celastraceae	Euonymus occidentale Nutt. ex Torr. var. occidentale	western burning bush		
Clusiaceae	Hypericum perforatum L.	common St. Johnswort		
Cornaceae	Cornus sericea L.	American dogwood		
Cucurbitaceae	Marah oreganus (Torr. ex S. Wats.) T.J. Howell	coastal manroot		
Cupressaceae	Sequoia sempervirens (Lamb. ex D. Don) Endl.	coast redwood		
Cupressaceae	Sequoiadendron giganteum (Lindl.) Buchh.	giant sequoia		
Cyperaceae	Carex deweyana Schwein.	Dewey's sedge		
Cyperaceae	Carex obnupta Bailey	slough sedge		
Cyperaceae	Carex sp. L.	sedge		
Cyperaceae	Cyperus eragrostis Lam.	tall flatsedge		
Dennstaedtiaceae	Pteridium aquilinum (L.) Kuhn	bracken fern		

Appendix A. Vascular plant species found in study location. Nomenclature follows yhe USDA PLANTS Database (USDA, 2007; continued).

Family	Species	Common name		
Dipsacaceae	Dipsacus fullonum L.	Fuller's teasel		
Dryopteridaceae	Athyrium filix-femina (L.) Roth	lady fern		
Dryopteridaceae	Polystichum munitum (Kaulfuss)	sword fern		
Equisetaceae	Equisetum arvense L.	common horsetail		
Equisetaceae	Equisetum hyemale L. var. affine (Engelm.) A.A. Eat.	giant scouring rush		
Ericaceae	Arbutus menziesii Pursh	Pacific madrone		
Ericaceae	Arctostaphylos columbiana Piper	hairy manzanita		
Ericaceae	Gaultheria shallon Pursh	salal		
Ericaceae	Vaccinium ovatum Pursh	evergreen huckleberry		
Ericaceae	Vaccinium parvifolium Sm.	red huckleberry		
Fabaceae	Lathyrus sp. L.	sweet-pea		
Fabaceae	Lotus corniculatus L.	broadleaf birdsfoot trefoil,		
Fabaceae	Lupinus sp. L.	lupine		
Fabaceae	Melilotus alba (L.) Lam.	yellow sweetclover		
Fabaceae	Trifolium sp. L.	clover		
Fabaceae	Vicia sativa L.	spring vetch		
Fagaceae	Chrysolepis sempervirens Kellogg Hjelmqvist	bush chinquapin		
Fagaceae	Lithocarpus densiflorus (Hook. & Arn.) Rehd.	tanoak		
Fagaceae	Quercus garryana Dougl. ex Hook.	Oregon white oak		
Gentianaceae	Centaurium muehlenbergii (Griseb.) W. Wight ex Piper	Muhlenberg's centaury		
Geraniaceae	Geranium dissectum L.	cut-leaved geranium		
Grossulariaceae	Ribes bracteosum Dougl. ex Hook.	stink currant		
Grossulariaceae	Ribes menziesii Pursh	canyon gooseberry		
Grossulariaceae	Ribes sanguineum Pursh	red-flowering currant		
Hydrangeaceae	Whipplea modesta Torr.	yerba de selva		
Hydrophyllaceae	Hydrophyllum tenuipes Heller	Pacific waterleaf		
Hydrophyllaceae	Nemophila pedunculata Dougl. ex Benth.	littlefoot nemophila		
Hydrophyllaceae	Phacelia sp. Juss	phacelia		
Iridaceae	Iris douglasiana Herbert	Douglas iris		
Iridaceae	Sisyrinchium bellum S. Wats.	blue-eyed-grass		
Juncaceae	Juncus patens E. Mey.	spreading rush		
Juncaceae	Juncus sp. L.	rush		
Lamiaceae	Mentha arvensis L.	wild mint		
Lamiaceae	Prunella vulgaris L.	self heal		
Lamiaceae	Satureja douglasii (Benth.) Kuntze	yerba buena		
Lamiaceae	Stachys rigida Nutt. ex Benth. var. rigida	rigid hedge-nettle		
Lauraceae	Umbellularia californica (Hook. & Arn.) Nutt.	California bay		
Liliaceae	Chlorogalum sp. Kunth	soap plant		
Liliaceae	Clintonia andrewsiana Torr.	bead lily		
Liliaceae	Dichelostemma capitatum (Benth.) Wood	bluedicks		
Liliaceae	Dichelostemma ida-maia (Wood) Greene	firecracker flower		
Liliaceae	Disporum hookeri Torr.	Hooker's fairy bells		
Liliaceae	Lilium kelloggii Purdy	Kellog's Lily		
Liliaceae	Lilium sp. L.	lily		
Liliaceae	Scoliopus bigelovii Torr.	fetid adder's tongue		

Appendix A. Vascular plant species found in study location. Nomenclature follows the USDA PLANTS Database (USDA, 2007; continued).

Family	Species	Common name		
Liliaceae	Maianthemum racemossum (L.) Link	false lily of the valley		
Liliaceae	Maianthemum stellatum (L.) Link	starry false lily of the valley		
Liliaceae	Trillium ovatum Pursh	coast trillium		
Linaceae	Linum bienne P. Mill.	pale flax		
Oleaceae	Fraxinus latifolia Benth.	Oregon ash		
Onagraceae	Circaea alpina L.	enchanter's nightshade		
Onagraceae	Epilobium sp. L.	fireweed		
Orchidaceae	Corallorrhiza striata Lindl.	striped coral root		
Oxalidaceae	Oxalis oregana Nutt.	redwood sorrel		
Papaveraceae	Dicentra formosa (Haw.) Walp.	Pacific bleeding heart		
.	Philadelphus lewisii Pursh ssp. californicus (Benth.)			
Philadelphaceae	Munz	wild mock orange		
Pinaceae	Abies grandis (Dougl. ex D. Don) Lindl.	grand fir		
Pinaceae	Pinus ponderosa P.& C. Lawson	ponderosa pine		
Pinaceae	Pinus radiata D. Don	Monterey pine		
Pinaceae	Pinus sylvestris L.	Scot's pine		
Pinaceae	Pseudotsuga menziesii (Mirbel) Franco	Douglas-fir		
Pinaceae	Tsuga heterophylla (Raf.) Sarg.	western hemlock		
Plantaginaceae	Plantago lanceolata L.	narrow-leaved plantain		
Poaceae	Agrostis stolonifera L.	creeping bentgrass		
Poaceae	Anthoxanthum odoratum L.	sweet vernal grass		
Poaceae	Avena sp. L.	oats		
Poaceae	Briza minor L.	little rattlesnake grass		
Poaceae	Bromus hordeaceus L.	soft brome		
Poaceae	Cortaderia jubata (Lem.) Stapf	pampas grass		
Poaceae	Cynosurus echinatus L.	bristly dogtail grass		
Poaceae	Dactylis glomerata L.	orchard grass		
Poaceae	Deschampsia cespitosa (L.) Beauv.	tufted hair-grass		
Poaceae	Elymus glaucus Buckl.	blue wildrye		
Poaceae	Hierochloe occidentalis Buckl.	California sweetgrass		
Poaceae	Holcus lanatus L.	velvet grass		
Poaceae	Hordeum sp. L.	barley		
	Leymus ×vancouverensis (Vasey) Pilger (pro sp.)			
Poaceae	$[mollis \times triticoides]$	wildrye		
Poaceae	Lolium perenne L.	English ryegrass		
Poaceae	Phalaris sp. L.	canarygrass		
Poaceae	Phleum pratense L.	common timothy		
Polemoniaceae	Navarretia squarrosa (Eschsch.) Hook & Arn.	skunkweed		
Polygonaceae	Rumex acetosella L.	common sheep sorrel		
Polygonaceae	Rumex crispus L.	yellow dock		
Polypodeaceae	Adiantum pedatum L.	northern maidenhair fern		
Portulaceae	Claytonia perfoliata Donn ex Willd.	miner's lettuce		
Portulaceae	Claytonia sibirica L.	Siberian candyflower		
Primulaceae	Trientalis borealis Raf. ssp. latifolia (Hook.) Hultén	broadleaf starflower		

Appendix A. Vascular plant species found in study location. Nomenclature follows the USDA PLANTS Database (USDA, 2007; continued).

Family	Species	Common name		
	Pentagramma triangularis (Kaulfuss) Yatskievych,			
Pteridaceae	Windham & Wollenweber	goldenback fern		
Pyrolaceae	Pyrola picta Sm.	white-veined wintergreen,		
Ranunculaceae	Ranunculus californicus	buttercup		
Rhamnaceae	Ceanothus thyrsiflorus Eschsch.	blue blossom		
Rhamnaceae	Rhamnus californica Eschsch.	coffeeberry		
Rhamnaceae	Rhamnus purshiana (DC.) Cooper	cascara buckthorn		
Rosaceae	Fragaria vesca L.	woodland strawberry		
Rosaceae	Heteromeles arbutifolia (Lindl.) M. Roemer	toyon		
Rosaceae	Holodiscus discolor (Pursh) Maxim.	oceanspray		
Rosaceae	Oemleria cerasiformis (Hook & Arn.) J.W. Landon	oso berry		
Rosaceae	Prunus emarginata (Hook.) Walp.	bitter cherry		
Rosaceae	Rosa californica Cham. & Schldl.	California rose		
Rosaceae	Rosa gymnocarpa Nutt.	wood rose		
Rosaceae	Rubus discolor Focke	himalayan blackberry		
Rosaceae	Rubus parviflorus Nutt.	thimbleberry		
Rosaceae	Rubus spectabilis Pursh	salmonberry		
Rosaceae	Rubus ursinus Cham. & Schlecht.	California blackberry		
Rubiaceae	Galium aparine L.	common bedstraw		
Salicaceae	Salix hookeriana Barratt ex Hook.	dune willow		
Salicaceae	Salix scouleriana Barratt ex Hook.	Scouler's willow		
Salicaceae	Salix sitchensis Sanson ex Bong.	Sitka willow		
Salicaceae	Salix sp. L.	willow		
Saxifragaceae	Tolmiea menziesii (Pursh) Torr. & Gray	pig-a-back plant		
Scrophulariaceae	Scrophularia californica Cham. & Schlecht.	California bee-plant		
Scrophulariaceae	Veronica americana Schwein. ex Benth.	water speedwell		
Urticaceae	Urtica dioica L.	stinging nettle		
Violeaceae	Viola sp. L.	violet		

Appendix B. Transformed divergence values for vegetation classes by image type. Vegetation codes are defined in Table 5.

Class	A-Uc-Ld	Pm	Pm-Uc	Pm-Ag	Ld -UcPm	IPG	P
(1) Spectral	separation for t	he aerial ph	oto (0.15 m, 4-	band)			
A-Uc-Ld	0						
Pm	133	0					
Pm-Uc	120	83	0				
Pm-Ag	134	23	33	0			
Ld-Uc-Pm	149	136	102	97	0		
IPG	1261	1029	991	999	698	0	
P	227	176	350	258	288	1112	0
(2) Spectral	separation for t	he Quickbir	d image (0.61	m, 4-band)			
A-Uc-Ld	0						
Pm	224	0					
Pm-Uc	59	112	0				
Pm-Ag	160	27	69	0			
Ld-Uc-Pm	177	99	71	113	0		
IPG	1606	1345	1483	1530	1196	0	
P	934	395	740	493	632	1419	0
(3) Spectral	separation for t	he NAIP im	age (1.0 m, 3-l	band)			
A-Uc-Ld	0						
Pm	119	0					
Pm-Uc	40	56	0				
Pm-Ag	122	23	31	0			
Ld-Uc-Pm	224	198	297	299	0		
IPG	489	548	650	695	137	0	
P	644	310	589	452	386	741	0