1 Title

- 2 Short-interval fires and vegetation change in southern California
- 3

4 Short running title

- 5 Short-interval fires and vegetation change
- 6

7 Authors

- 8 Stephanie M. Lucero¹, Nathan C. Emery², Carla M. D'Antonio^{1,3}
- ⁹ ¹Department of Ecology, Evolution, and Marine Biology, University of California, Santa
- 10 Barbara, Santa Barbara, CA, USA.
- ² Department of Plant Biology, Michigan State University, East Lansing, MI, USA.
- 12 ³Environmental Studies Program, University of California, Santa Barbara, Santa Barbara, CA,
- 13 USA.
- 14

15 Correspondence

16 Stephanie M. Lucero, Department of Ecology, Evolution, and Marine Biology, University of

17 California, Santa Barbara, Santa Barbara, CA, USA.

- 18 Email: <u>sama@ucsb.edu</u>
- 19
- 20 Funding

21 The California Energy Commission (500-10-045, CMD) and the University of California, Santa

- 22 Barbara provided funding.
- 23

24 Abstract

25 Questions

26	In southern California, shortened fire return intervals may contribute to a decrease in
27	native chaparral shrub presence and an increase in non-native annual grass presence. To test the
28	hypothesis that short-fire return intervals promote a loss in shrub cover, we examined the
29	contribution of single short-interval fires and abiotic conditions on the change of shrub cover
30	within Ventura and Los Angeles counties. Through evaluating pre- and post-fire historical aerial
31	images, we answered the following questions, 1) How has vegetation type cover changed after
32	repeat fires? and 2) What landscape variables contribute the most to the observed change?
33	
34	Location
35	Ventura County and Los Angeles County, California, USA.
36	
37	Methods
38	We assessed the impact of a single short-interval fire by comparing vegetation recovery in
39	adjacent once- and twice-burned fire burn polygons (long- and short-interval respectively). Pixel
40	plots were examined within each polygon and vegetation cover was classified to vegetation type.
41	We determined the best predictor of vegetation type cover with a linear mixed effects model
42	comparison using Akaike Information Criterion.
43	
44	Results
45	Pre-fire and post-fire community type cover was highly correlated. Burn interval was the best
46	predictor of tree cover change (lower cover in twice-burned pixel plots). Aspect was the best

48	was the best predictor of chaparral cover change (positive correlation) and sage scrub cover
49	change (negative correlation). Conversion of chaparral to sage scrub cover was more likely to
50	occur than conversion of chaparral to annual grass cover.
51	
52	Conclusions
53	Our study did not find extensive evidence of a decrease in chaparral shrub cover due to a single
54	short-interval fire. Instead, post-fire cover was highly correlated with pre-fire cover. Chaparral
55	recovery, however, was dynamic suggesting that stand recovery may be strongly influenced by
56	local scale conditions and processes.
57	
58	Keywords
59	southern California, chaparral, short-interval fire, vegetation change, historical aerial
60	photographs, georectification, linear mixed-effects model, type conversion, sage scrub, grass
61	
62	
63	
64	
65	
66	
67	
68	
69	
70	
71	

72 Introduction

73 Fire-adapted ecosystems around the world are experiencing significant changes to the historical 74 fire regime due to human influences (Piñol et al., 1998, Gillett et al., 2004, Syphard et al., 2007b, 75 Nowacki and Abrams, 2008). The interval time between fire events (fire return interval), is one 76 of the most direct way humans can alter a fire regime. Fire suppression has lengthened the 77 interval time between fires in the northern Rockies (Barrett and Arno, 1982), in western 78 Washington (Everett et al., 2000), and in the Sierra Nevada mountains (McKelvey et al., 1996) 79 leading to with fire-adapted tree species being replaced by fire-sensitive, shade-tolerant species. 80 A shorter fire interval can also put fire-adapted ecosystems at risk. In southern California, a 81 shorter fire return interval is considered one of the leading drivers of native shrubs being 82 replaced by non-native annual grasses (Keeley, 2000, Haidinger and Keeley, 1993, Zedler et al., 83 1983). 84 In southern California chaparral ecosystems, the historical fire return interval ranges from 85 30-90 years (Keeley et al., 2004, Van de Water and Safford, 2011) and in some locations may be 86 as long as 150 years (Syphard et al., 2006). In this Mediterranean-type climate region, fires are 87 typically crown fires (Hanes, 1971) occurring in the summer and fall (Beyers and Wakeman, 88 2000). Post fire, these stands generally return to pre-fire canopy cover within the first decade 89 (Hope et al., 2007, Peterson and Stow, 2003) and to pre-fire stand structure within the second 90 decade following fire (Hanes, 1971, Schlesinger and Gill, 1978).

In contrast, the current fire return interval can be much shorter due to an expansion of the
wildland-urban-interface (Syphard et al., 2007a) and increased year-round anthropogenic
ignitions caused by increasing human populations (Keeley and Fotheringham, 2001). Climate
change is also expected to shorten mean fire return intervals as temperatures in southern
California become warmer and precipitation more variable, thus increasing the risk of ignition

96 (Krawchuk and Moritz, 2012, Polade et al., 2017). Furthermore, with the introduction of non-97 native annual grasses, chaparral communities could be driven toward a new successional 98 trajectory leading to the extirpation of native shrub species and the expansion of non-native 99 annual grasses (Brooks et al., 2004, Keeley and Brennan, 2012, D'Antonio and Vitousek, 1992). 100 A shortened fire interval has been recognized as one of the main causes of chaparral 101 conversion to non-native annual grasses (Keeley, 2000, Syphard et al., 2019b). This is because 102 many chaparral species require five-to-ten years to reach reproductive maturity (Zammit and 103 Zedler, 1993) and at least 15 years to establish a robust seedbank (Syphard et al., 2018b, Park et 104 al., 2018, Keeley, 2000). This is especially true of obligate seeding species, which are not fire 105 resistant and germinate from seed following fire (Keeley, 1991). Extirpation of obligate seeding 106 species and an increase in non-native species have been observed in the field when a second fire 107 occurred in short succession (Zedler et al., 1983, Haidinger and Keeley, 1993, Keeley and 108 Brennan, 2012, Jacobsen et al., 2004). 109 A single short-interval fire, however, does not always lead to chaparral conversion. 110 Previous work with remotely sensed data suggest that water availability (Park et al., 2018, 111 Syphard et al., 2019a), elevation (Meng et al., 2014), and mean annual temperature (Storey et al., 112 2021) explain more variation among chaparral stand recovery (although also see Syphard et al. 113 (2019b)). These studies utilized imagery with 30-m spatial resolution and/or datasets with a 30-m 114 scale. 115 There exists a research gap between in-person observations following a single fire event 116 and landscape-scale observations across multiple fire events. The goal of this study was to

address this limitation by using high resolution (1-meter) historical aerial imagery to observe

detailed vegetation regrowth across the landscape and across multiple historical- and short-

119 interval fire events.

120	In this study, we measured the difference in vegetation type cover following once- or
121	twice-burned fire polygons (long- and short-interval fires, respectively) using historical aerial
122	photographs ranging from 1956 and 2003 across two counties where, if chaparral conversion
123	were occurring, we should be able to detect chaparral loss and decline. Through evaluation of
124	pre- and post-fire images, we asked 1) How has vegetation type cover (i.e., chaparral, sage scrub
125	grass, tree) changed after repeat fires? and 2) What landscape variables contribute to the
126	observed changes in vegetation cover?

- 127
- 128 Methods

129 Mapping the occurrence of short-interval fires

130 Fire history data (1879-2009) were acquired from the Fire and Resources Assessment Program 131 (FRAP) database (CALFIRE, www.fire.ca.gov), reporting fires \geq 4 hectares. The FRAP database 132 was used because we required spatially explicit polygons for our analysis (although see Syphard 133 and Keeley (2016) for the limitations of using the FRAP database). The shapefile was clipped in 134 ArcMap (ArcGIS 10.1) to select for wildfires that fell within the study area of Ventura and Los 135 Angeles counties and then processed to create an Interval Wildfire Occurrence map (Figure 1), 136 each polygon having its own unique wildfire history based on original wildfire perimeters. 137 The attribute table of the Interval Wildfire Occurrence map was exported to MS Excel 138

(Microsoft 2011) and new metrics such as "Number of Fires" and "Minimum Fire Interval" were calculated (Appendix 1). Wildfire perimeters were filtered to eliminate single wildfires that were reported by multiple agencies, for example if a polygon had multiple "Fire Alarm Dates" in the same year (Jacobsen et al., 2004). The modified data table was finally joined back to the merged shapefile in ArcMap.

144 Selection of twelve paired sites in Ventura and Los Angeles Counties

Ventura and Los Angeles Counties are ideal for determining the effects of a short-interval
wildfire because the region is highly vulnerable to fire during the dry months (typically JulyOctober) when Santa Ana wind conditions promote fast spreading wildfires (Hughes and Hall,
2010). In addition, the number of short-interval wildfires is predicted to increase as the
population of southern California continues to grow (Keeley and Fotheringham, 2001, Myers and
Pitkin, 2013).

151 Study sites were selected from the Interval Wildfire Occurrence map (Figure 1). We 152 defined a "short-interval fire" as a fire return interval of five or fewer years. To quantify the 153 effects of a single short-interval fire on vegetation, we selected polygons that experienced two 154 wildfires within a five-year period that had an adjacent polygon that experienced only one wildfire within the same five-year period. All polygons had to be at least ≥ 0.5 km² to allow for 155 156 sufficient subsampling and analysis. Polygons that experienced a short-interval fire were considered "twice-burned" polygons (e.g., burned in 1962 and 1967) and adjacent polygons that 157 158 only experienced the latter fire (e.g., 1967) were considered "once-burned" polygons. Selecting 159 paired polygons that burned in the second fire, allowed for the polygons to experience the same 160 number of post-fire recovery years. Long-interval polygons had, on average, 26.6 ± 5.2 (standard 161 error mean) years of regrowth since the last wildfire and two of the twelve polygons (Site 3 and 162 Site 9) had no prior record of wildfire since the early 1900s. For two sites (Site 4 and Site 6), 163 long-interval polygons burned in the first wildfire year (instead of the second wildfire) and were analyzed in images ≥19 years post-fire. This exception was allowed assuming any difference in 164 165 vegetation cover between long-interval and short-interval polygons would be negligible after ≥ 19 166 years of regrowth (Zammit and Zedler, 1993).

167	In addition to burn history, sites were selected to represent vegetation variation within the
168	two counties and along a moisture gradient determined by their distance from the coast. Sites
169	closer to the coast are generally more mesic and sites farther inland are generally more arid
170	(Franklin, 1998). Water availability influences chaparral and sage scrub community composition
171	and extent (Mooney and Parsons, 1973, Poole and Miller, 1981). Sites were selected to represent
172	the vegetation communities within Los Padres National Forest in Ventura County and within the
173	Angeles National Forest and the Santa Monica Mountain National Recreation Area in Los
174	Angeles County.
175	
176	Selecting aerial photographs
177	Historical aerial photographs (HAPs) were acquired from the Map and Imagery Laboratory
178	(MIL) at the University of California, Santa Barbara in 2011 to 2014
179	(www.library.ucsb.edu/mil). HAPs were chosen between 1952 and 2009 for corresponding
180	wildfires spanning 1956 to 2003. Pre-fire HAPs were selected as close to before the first wildfire
181	as possible to record initial vegetation cover and post-fire HAPs were selected ≥ 6 years
182	following the second wildfire to capture maximal vegetation cover without encountering a third
183	wildfire (Appendix 1). Vegetation communities were assumed to return to pre-fire canopy cover
184	within six years following wildfire (Muller et al., 1968, Schlesinger and Gill, 1978). Seasonality
185	of images was not controlled for under the assumption that mature communities appear
186	distinguishable year-round (i.e., chaparral appears darker with a closed canopy year-round; grass
187	appears lighter and has no visible canopy structure year-round). Final HAP selection was based
188	on photograph availability and adherence to fire criteria.
189	

190 Georectifying aerial photographs

191 To compare pre- and post-fire vegetation cover on a pixel-by-pixel basis, all HAPs were 192 georectified to the same base image. Grayscale, 2009, one-meter spatial resolution, digital 193 orthophoto quarter quads (DOQQ) of Ventura or Los Angeles County, collected by the United 194 States Geological Survey, were used as the base image. Temporally stable objects such as large 195 shrubs or trees, rock outcrops, and crests and troughs of the mountainous landscape were used as 196 registration points (RPs). Dirt roads and permanent structures were also used, although these 197 more permanent features were rare in the HAPs due to the remoteness of the polygons. Because 198 the terrain of the HAPs was mountainous and highly variable, RPs were placed at a high density 199 to increase warping accuracy. Each HAP was then warped using triangulation and pixels were 200 resampled to the nearest neighbor, creating a georectified HAP with one-meter spatial resolution. 201 Georectified HAPs (gHAPs) were then mosaicked to minimize edge distortion and 202 increase spatial accuracy for vegetation analysis. Mosaicked gHAPs covered the entire long-203 interval and short-interval polygon of a site under pre-fire and post-fire conditions. Only two 204 sites (Sites 2 and 6) were not georectified across their entire long-interval polygon due to a lack 205 of available HAPs and/or their extensive size and instead an equivalent or greater area to the 206 short-interval polygon was georectified.

207 Mosaicked gHAPs were validated for their spatial accuracy by identifying 40-100 RPs 208 corresponding to the 2009 DOQQ base map. Validation RPs had a final root mean square error 209 of ten pixels (i.e., ten meters) or less.

210

211 Plot selection on north and south aspects within long- and short-interval polygons

Aspect is a large influencer of vegetation type cover (Hanes, 1971) with south facing aspects receiving more solar radiation than north facing aspects. Random points were generated within each site's pre-fire gHAP and approximately eight points in the long-interval polygon and eight

215	points in the short-interval polygon were selected for vegetation analysis. At each point, a 50 x
216	50 pixel plot (50 x 50 m) was established. Pixel plots (hereafter "plots") were distributed
217	between north and south facing aspects (north: 0.0° to 67.5° or 292.5° to 360°; south: 112.5° to
218	247.5°) to account for differences in solar irradiance (northern aspects receive less solar
219	irradiance than southern aspects). Plots were shifted if needed to ensure they did not overlap a
220	mountain ridge or valley and to fit entirely on one aspect. Aspect was verified with 30-meter
221	USGS Digital Elevation Model (DEM) data and/or visually with Google Earth (Google Earth
222	Pro 7.3.0.3832). All pre-fire plots were replicated in the post-fire mosaicked gHAP to capture
223	vegetation regrowth at the same location.
224	In total, 198 plots were analyzed with 99 plots in long-interval polygons and 99 plots in
225	short-interval polygons. One hundred and four plots had northern aspects and 94 plots had
226	southern aspects. Plots were considered independent after including site as a covariate in plot
227	level analysis and found no significant influence on plot level results.
228	Prescribed burns included in the FRAP database were reviewed and only two of the 198
229	plots overlapped with a prescribed burn. These two plots were removed from the analysis.
230	
231	Quantifying vegetation type cover within plots
232	To quantify vegetation type cover at each plot, the "dot grid" method was used (Floyd
233	and Anderson, 1982, Dublin, 1991). A ten-by-ten grid (100 points) was overlaid on each plot
234	with a spacing of five pixels (five meters) between each point. Vegetation cover was classified to
235	vegetation type: chaparral, grass, sage scrub or tree. All grass cover was assumed to be non-
236	native dominated based on the 1930's Wieslander Maps and the 2001 USDA California
237	Vegetation map. For classification consistency, all sites were examined twice to account for
238	initial training and improvement in classification over time.

239	To improve classification accuracy, solar zenith was considered to account for shadows
240	and Google Earth was referenced for vegetation type cover and seasonal changes (available
241	years: 1990-2015). The authors traveled to six of the twelve sites to confirm that site cover
242	approximated vegetation type cover observed in the HAPs (e.g., a matrix of chaparral, sage
243	scrub, and grass in the HAPs were a matrix of vegetation in the field). However, verification of
244	the HAPs was infeasible due to many of the images being decades old.
245	Percent cover (%) was quantified by tallying the number of points classified within each
246	vegetation type (100 points = 100% cover). Pre-fire vegetation cover was subtracted from post-
247	fire vegetation cover to quantify the amount of vegetation type change within each plot.
248	
249	Datasets for abiotic variables
250	Aspect was calculated from USGS digital elevation models (DEMs) with a 30 x 30 meter
251	horizontal resolution and a one-meter vertical resolution. Distance from the coast was calculated
252	in ArcMap by determining the centroid point of each polygon and measuring the shortest direct
253	distance to the coastline (Appendix 1).
254	Moisture availability following fire, influences seedling survival and thus eventual
255	vegetation type cover (Pratt et al., 2014, Venturas et al., 2016). It was calculated from PRISM
256	(http://www.prism.oregonstate.edu/explorer/) averaging the annual precipitation during the first
257	five years of regrowth following the latter wildfire.
258	
259	Statistical analysis
260	Statistical analysis was conducted using RStudio (RStudio, Inc. version 0.98.1103) and R version
261	3.3.2 (http://www.rstudio.com/) and were either run at the plot level (e.g., 99 once-burned and 99

twice-burned plots) or at the polygon level (e.g., 12 once-burned polygons and 12 twice-burned
polygons). Polygon values were calculated as the mean of their plot values.

264 As post-fire vegetation type cover at the plot level was highly correlated with pre-fire 265 vegetation type cover (Figure 2), the residuals of a linear regression for pre- and post-fire 266 vegetation types were derived to account for pre-existing plant communities. A linear mixed-267 effects model of plot data (N = 198) was run for each vegetation type with site as a random effect 268 to reduce the effects of spatial autocorrelation. The models to predict the residuals were: burn 269 interval, aspect, years after fire (when post-fire HAPs were taken), distance to coast, and the 270 five-year average rainfall post-fire. In total, five models were run for each vegetation type. The 271 Akaike Information Criterion (AIC) values of each model were compared within a vegetation 272 type. The model(s) with the lowest AIC value were further investigated for each vegetation type. 273 If the best model predictor was categorical, then an ANOVA was performed to determine 274 statistical significance and if the model predictor was continuous, a linear regression was 275 performed to determine significance.

276

277 Results

278 Trends in vegetation type cover comparing pre-fire and post-fire conditions

279 For each vegetation cover type, post-fire conditions were highly correlated with pre-fire

280 conditions (N = 198) (Figure 2). Linear regressions were significant for chaparral percent cover

281 (P < 0.001, $r^2 = 0.93$), sage scrub percent cover (P < 0.001, $r^2 = 0.91$), grass percent cover (P < 0.001, $r^2 = 0.91$), grass percent cover (P < 0.001, $r^2 = 0.91$), grass percent cover (P < 0.001, $r^2 = 0.91$), grass percent cover (P < 0.001, $r^2 = 0.91$), grass percent cover (P < 0.001, $r^2 = 0.91$), grass percent cover (P < 0.001, $r^2 = 0.91$), grass percent cover (P < 0.001, $r^2 = 0.91$), grass percent cover (P < 0.001, $r^2 = 0.91$), grass percent cover (P < 0.001, $r^2 = 0.91$), grass percent cover (P < 0.001, $r^2 = 0.91$), grass percent cover (P < 0.001, $r^2 = 0.91$), grass percent cover (P < 0.001, $r^2 = 0.91$), grass percent cover (P < 0.001, $r^2 = 0.91$), grass percent cover (P < 0.001, $r^2 = 0.91$), grass percent cover (P < 0.001, $r^2 = 0.91$), grass percent cover (P < 0.001, $r^2 = 0.91$), grass percent cover (P < 0.001, $r^2 = 0.91$), grass percent cover (P < 0.001, $r^2 = 0.91$), grass percent cover (P < 0.001, $r^2 = 0.91$), grass percent cover (P < 0.001, $r^2 = 0.91$), grass percent cover (P < 0.001, $r^2 = 0.91$), grass percent cover (P < 0.001, $r^2 = 0.91$), grass percent cover (P < 0.001, $r^2 = 0.91$), grass percent cover (P < 0.001, $r^2 = 0.91$), grass percent cover (P < 0.001, $r^2 = 0.91$), grass percent cover (P < 0.001, $r^2 = 0.91$), grass percent cover (P < 0.001, $r^2 = 0.91$), grass percent cover (P < 0.001, $r^2 = 0.91$), grass percent cover (P < 0.001, $r^2 = 0.91$), grass percent cover (P < 0.001, $r^2 = 0.91$), $r^2 = 0.91$, $r^2 = 0.91$

282 0.001, $r^2 = 0.93$), and tree percent cover (P < 0.001, $r^2 = 0.94$).

283

284 *Linear mixed-effects model*

285	When analyzing chaparral vegetation type residuals, the models with the lowest AIC values
286	included aspect and years since fire (Table 1). An ANOVA of aspect yielded a trend towards a
287	difference between north and south facing aspects ($P = 0.083$, F = 3.05) with south-facing
288	aspects showing greater decreases in cover than north-facing aspects (Figure 3, top right).
289	Change in chaparral cover was positively correlated with longer recovery time post-fire (Figure
290	3, top left, $P < 0.001$, t = 3.507). For sage scrub residuals, the models with the lowest AIC were
291	also aspect and years after fire. Plots on north-facing aspects had greater positive change in sage
292	scrub percent cover than south-facing slopes (Figure 3, middle right, $P = 0.034$, F = 4.567).
293	Change in sage scrub cover was negatively correlated with years after fire (Figure 3, middle left,
294	P = 0.005, t = -2.823). The lowest AIC value for grass residuals was years after fire, however the
295	difference in AIC value among models was very slight. A linear regression of the change in
296	percent grass and years after fire was not significant (Figure 3, bottom left, $P = 0.185$, t = -
297	1.332). For tree residuals, the best model was burn interval. Once-burned plots had more
298	negative change in tree cover than short-interval plots (Figure 4, ANOVA, $P = 0.030$, F = 4.769).
299	This pattern was still significant when the outlier site (Site 11, Plot 7: -39.97) was removed
300	(ANOVA, <i>P</i> = 0.048, F = 3.97).
201	

301

302 *Transition matrix*

303 The majority of pixels showed no change in cover (Figure 5). The majority pixels located in

304 once-burned plots returned to their pre-fire cover (chaparral: 86.3%; sage scrub: 80.4%; grass:

305 76.3%; tree: 74.0%). Pixels in twice-burned plots also often returned to their pre-fire cover

306 (chaparral: 87.1%; sage scrub: 78.4%; grass: 70.0%; tree: 65.7%).

The transition direction and the proportion of pixels that changed in cover were similar
between once- and twice-burned locations. Chaparral pixels were more likely to convert to sage

scrub than to grass (once-burned: 8.4% and 1.3% respectively; twice-burned: 10.5% and 0.7%
respectively). Sage scrub pixels were more likely to convert to chaparral than to grass (once
burned: 10.8% and 5.2% respectively; twice-burned: 14.0% and 5.5% respectively). Grass pixels
that converted were most likely to convert to sage scrub cover (once-burned: 17.0%; twiceburned: 23.8%).

314

315 Discussion

The results of this study found that a single short-interval fire, as quantified in twiceburned plots, did not lead to significant vegetation type change of chaparral, sage scrub or grass cover at the landscape scale. A short-interval fire was only a significant predictor of reduced vegetation cover for plots dominated by trees (Table 1). Instead, overall vegetation recovery trends were driven by the pre-existing cover (i.e., post-fire cover was significantly correlated with pre-fire cover) (Figure 2).

There were, however, a large number of plots that either increased or decreased in cover about the mean. Suggesting there is greater heterogeneity at the individual plot level than there is at the landscape level. Ninety-eight out of 198 chaparral plots and 91 out of 198 sage scrub plots showed an increase or decrease in woody cover following fire. This heterogeneity in canopy regrowth is consistent with previous findings (Syphard et al., 2018a, Park et al., 2018, Storey et al., 2021).

While we set out to observe if there was evidence of chaparral conversion to grass following a single short-interval fire, we found there was greater conversion from chaparral to sage scrub (10.5%) than chaparral to grass (0.6%). In comparison, sage scrub was more likely to convert to grass (5.5%). These interactions may be influenced by these communities' physical distribution. Sage scrub communities are typically found below chaparral at lower elevations,

which tend to be hotter and drier, and adjacent to urban areas, which tend to be dominated by
annual grasses. Similar results were found by Syphard et al. (2018a, 2018b, 2006) and Meng et
al. (2014).

Aspect was a significant driver of sage scrub cover (Figure 3, middle right) and a strong driver of chaparral cover (Figure 3, top right) with cover increasing on north-facing aspects and decreasing on south-facing aspects. This result was unexpected as north-facing aspects tend to have more mesic conditions and receive less solar radiation, conditions where chaparral typically outcompete sage scrub (Syphard et al., 2006). Miller et al. (1983) showed similar dry down periods (0-2 weeks) between north- and south-facing aspects, indicating soil moisture on opposing aspects may not be as different as originally expected.

There was strong evidence that chaparral cover increased on north-facing aspects and
decreased on south-facing aspects (Figure 3, top right), which is consistent with previous
research (Hanes, 1971, Keeley and Keeley, 1981) and other studies that found metrics of soil
aridity to be positively correlated with chaparral loss (Syphard et al., 2019a) and grass presence
(Park et al., 2018).

Another explanation as to why no significant difference was detected between north- and south-facing aspects could be due to the expansion of *Malosma laurina*, a tenacious facultative resprouter, on south-facing aspects, which can outcompete other species when water resources are limited (Thomas and Davis, 1989). Because *M. laurina* can reach similar heights and canopy densities as mixed chaparral stands, locations dominated by *M. laurina* would have been classified as "chaparral" in the HAPs.

While we were readily able to differentiate between vegetation cover types (e.g., grass, sage scrub, chaparral, tree), we were not able to identify vegetation to species, which meant we were unable to detect if there was a change in community composition or a change in the presence of

357 obligate seeding, obligate resprouting, or facultative resprouting shrub species. These 358 reproductive strategies are predicted to respond differentially to increased fire frequency 359 (Keeley, 1991, Syphard et al., 2006, Franklin et al., 2004) and distribution can vary by water 360 availability (Mooney and Parsons, 1973, Poole and Miller, 1981, Franklin, 1998, Meentemeyer 361 and Moody, 2002). 362 In addition, no significant difference in chaparral cover on north- or south-facing aspects 363 could be due to the method of data collection. Since values from the dot-grid method spanned 0-364 100%, plots with close to 100% chaparral cover pre-fire had little room to increase in chaparral 365 cover post-fire. Within plots that had 100% chaparral cover pre-fire, a decrease in cover or "no 366 change" was the only possible outcome. Indeed, average pre-fire chaparral cover on north-facing 367 (more mesic) aspects was $72.19 \pm 3.78\%$ compared to only $26.33 \pm 3.49\%$ on south-facing (more 368 arid) aspects. North-facing aspects had more chaparral cover to lose at the outset. 369 Calculating the relative change in vegetation cover was explored, however it inflated non-370 biologically important results (e.g., 1 pixel post-fire / 2 pixels pre-fire = decrease of 50%) and 371 hid larger community changes (e.g., 60 pixels post-fire / 100 pixels pre-fire = decrease of 40%). 372 For this reason, all changes in cover were calculated as their absolute value (e.g., 1 pixel post-fire 373 - 2 pixels pre-fire = decrease by 1).

Chaparral and sage scrub cover showed strong yet opposing trends in succession following fire. Chaparral cover increased significantly with additional years after fire whereas sage scrub cover strongly declined with additional years after fire (Figure 3, left). This supports previous studies that suggest sage scrub can be successional to chaparral given enough time without fire (McPherson and Muller, 1967, Gray, 1983). This transition in canopy cover is consistent with the assumption that chaparral stands require five to thirty years to recover after wildfire (Hanes, 1971, Hope et al., 2007, Schlesinger and Gill, 1978) and subshrubs, common in

sage scrub communities, decline as chaparral shrubs mature around them (Syphard et al., 2006).
Thus, it could be possible that with enough fire-free years, chaparral cover could expand via the
establishment of chaparral species within sage scrub stands.

Number of years post-fire and vegetation succession may also explain some of the vegetation change detected in Syphard et al. (2018a). Some locations observed to convert from shrub to grass cover may have been dominated by or included herbaceous cover in the process of recovering following fire and may not represent the mature community. The Day fire (2006), Zaca fire (2007), and La Brea fire (2009) occurred within seven years prior to the 2013 Landfire vegetation map, so it is within the window of expected recovery that some of these locations were still recovering and may eventually return to shrub cover absent of short-interval fires.

391 We hypothesized sites closer to the coast would be more mesic and sites farther from the 392 coast would be more arid, leading to less vegetation conversion near the coast and more 393 conversion inland. We did not, however, find distance to the coast to be a significant factor. This 394 could be due to vegetation trends being more strongly correlated with moisture gradients driven 395 by elevation than by distance from the coast. Indeed, Meng et al. (2014) and Syphard et al. 396 (2019b) found elevation to be a strong driver of vegetation change with more chaparral 397 conversion occurring at lower elevations. Analysis by Storey et al. (2019) included a more 398 extensive range of chaparral stands in southern California and found vegetation change was 399 greatest in the eastern portion of their study region and at higher elevations and was a result of 400 drought and fire interactions.

401 The five-year average rainfall following the latter fire was also found to not be a
402 significant driver of vegetation type change, which is in line with other studies (Storey et al.,
403 2020, Meng et al., 2014), although see Storey et al. (2021). Storey et al. (2020) found climactic
404 water deficit to be a stronger predictor of chaparral recovery compared to total precipitation and

405 water conditions preceding fire were generally more predictive of recovery than conditions 406 following fire. Finer temporal scale patterns of rainfall (e.g., light rainfall over a month vs. a one-407 day downpour, early winter vs. late winter) will also likely be important to consider in the 408 seasons leading up to fire and the first year following fire. This will especially be true of 409 locations that are already water limited (e.g., < 500 mm y⁻¹).

410 While almost all polygons included in this study experienced additional fires before and 411 after the years of analysis, multiple short-interval fires were not included in this analysis nor was 412 the entire fire history (1878-2009) of a location. Storey et al. (2021) included additional fire 413 history by comparing locations in southern California that burned once, twice, or three times 414 within 25 years. They found no difference in shrub recovery between stands that burned once or 415 twice but did find a decrease in shrub recovery when locations burned three times within 25 416 years. Investigating sites that experienced multiple short-interval fires will continue to be highly 417 valuable as will investigating the impact of variability in years between fires (Zedler, 1995) on 418 vegetation change.

Another factor that might have contributed to our results is the time frame of analysis (i.e., 1956-2003). Resilient shrub species may have already been selected for in locations that experienced multiple fires. Thus, any vegetation change that would have occurred due to a shortinterval fire may have already occurred, prior to when imagery was first available.

While this study did not find a consistent increase in grass cover following a single shortinterval fire, non-native annual grasses still pose a risk to native plant communities. When nonnative annual grasses senesce they create highly ignitable, continuous fuel across the landscape that can carry fire into chaparral stands that otherwise would not have ignited (Keeley, 2000). An expanding wildland-urban-interface (Syphard et al., 2007b) and climate change (Krawchuk and Moritz, 2012) are also expected to create additional risks for southern California's wildlands.

429

430 Conclusion

431 The aim of this study was to identify how vegetation type cover changed in response to a 432 single short-interval fire (defined as two fires within five years) and how environmental variables 433 contributed to vegetation type change across the landscape. We did not find extensive evidence 434 that short-interval fires promoted a landscape-scale loss of shrublands. Instead, we found 435 chaparral cover only slightly declined in twice-burned plots. Tree cover was the only vegetation 436 type to significantly decline following a single-short interval fire. Our data are consistent with 437 the hypothesis that chaparral conversion to other vegetation types occurs slowly and is likely 438 influenced by its abiotic environment. Aspect was the best predictor of differences in sage scrub 439 cover while the number of years post-fire was the best predictor of chaparral and sage scrub 440 stand regrowth. Post-fire and pre-fire cover, overall, were highly correlated for all vegetation 441 cover types, although variation at the pixel plot level was apparent, suggesting that vegetation 442 change occurring at the local scale may be influenced primarily by local scale processes. 443 Therefore, caution is advised when managing fire-prone vegetation types as these landscapes can 444 be highly heterogeneous and outcomes at one location may not reflect landscape level patterns. 445

446

447 Acknowledgements

We are grateful to E. Burley, S. Suresh, C. Allred, T. Madden, and M. Plummer for their
assistance processing aerial images; to S. Peterson for guidance in ENVI and ArcGIS; and to P.
Dennison, M. Moritz, and R. Oono for their advice, guidance, and feedback on earlier versions

451 this manuscript.

453	
454	Citations
455	
456	BARRETT, S. W. & ARNO, S. F. 1982. Indian Fires as an Ecological Influence in the Northern
457	Rockies. Journal of Forestry, 80, 647-651.
458	BEYERS, J. L. & WAKEMAN, C. D. 2000. Season of burn effects in southern California
459	chaparral. Second interface between ecology and land development in California. Open-
460	File Report 00-62. Sacramento, CA: US Department of the Interior, Geological Survey,
461	45-55.
462	BROOKS, M. L., D'ANTONIO, C. M., RICHARDSON, D. M., GRACE, J. B., KEELEY, J. E.,
463	DITOMASO, J. M., HOBBS, R. J., PELLANT, M. & PYKE, D. 2004. Effects of
464	invasive alien plants on fire regimes. Bioscience, 54, 677-688.
465	DUBLIN, H. T. 1991. Dynamics of the Serengeti-Mara woodlands: an historical perspective.
466	Forest & Conservation History, 35, 169-178.
467	D'ANTONIO, C. & VITOUSEK, P. M. 1992. Biological invasions by exotic grasses, the
468	grass/fire cycle, and global change. Annual Review of Ecology and Systematics, 23, 63-
469	87.
470	EVERETT, R. L., SCHELLHAAS, R., KEENUM, D., SPURBECK, D. & OHLSON, P. 2000.
471	Fire history in the ponderosa pine/Douglas-fir forests on the east slope of the Washington
472	Cascades. Forest Ecology and Management, 129, 207-225.
473	FLOYD, D. A. & ANDERSON, J. E. 1982. A new point interception frame for estimating cover
474	of vegetation. Vegetatio, 50, 185-186.
475	FRANKLIN, J. 1998. Predicting the distribution of shrub species in southern California from
476	climate and terrain-derived variables. Journal of Vegetation Science, 9, 733-748.

- 477 FRANKLIN, J., COULTER, C. L. & REY, S. J. 2004. Change over 70 years in a southern
- 478 California chaparral community related to fire history. *Journal of Vegetation Science*, 15,
 479 701-710.
- 480 GILLETT, N., WEAVER, A., ZWIERS, F. & FLANNIGAN, M. 2004. Detecting the effect of
- 481 climate change on Canadian forest fires. *Geophysical Research Letters*, 31.
- 482 GRAY, J. T. 1983. Competition for light and a dynamic boundary between chaparral and coastal
 483 sage scrub. *Madrono*, 43-49.
- 484 HAIDINGER, T. L. & KEELEY, J. E. 1993. Role of high fire frequency in destruction of mixed
 485 chaparral. *Madrono*, 141-147.
- 486 HANES, T. L. 1971. Succession after fire in chaparral of southern California. *Ecological*487 *Monographs*, 41, 27-&.
- 488 HOPE, A., TAGUE, C. & CLARK, R. 2007. Characterizing post-fire vegetation recovery of
- 489 California chaparral using TM/ETM+ time-series data. *International Journal of Remote*490 Sensing, 28, 1339-1354.
- HUGHES, M. & HALL, A. 2010. Local and synoptic mechanisms causing Southern California's
 Santa Ana winds. *Climate Dynamics*, 34, 847-857.
- 493 JACOBSEN, A., DAVIS, S. & FABRITIUS, S. 2004. Fire frequency impacts non-sprouting
- 494 chaparral shrubs in the Santa Monica Mountains of southern California. *Ecology*,
- 495 *conservation and management of mediterranean climate ecosystems. Millpress,*
- 496 *Rotterdam, Netherlands.*
- 497 KEELEY, J. E. 1991. Seed-germination and life-history syndromes in the California chaparral.
- 498 *Botanical Review*, 57, 81-116.

- 499 KEELEY, J. E. Fire and invasive species in Mediterranean-climate ecosystems of California.
- 500 Proceedings of the invasive species workshop: the role of fire in the control and spread of
- 501 invasive species. Fire conference, 2000. 81-94.
- 502 KEELEY, J. E. & BRENNAN, T. J. 2012. Fire-driven alien invasion in a fire-adapted
- 503 ecosystem. *Oecologia*, 169, 1043-1052.
- 504 KEELEY, J. E., FOTHERINGHAM, C. & MORITZ, M. A. 2004. Lessons from the october
- 505 2003. Wildfires in Southern California. *Journal of Forestry*, 102, 26-31.
- 506 KEELEY, J. E. & FOTHERINGHAM, C. J. 2001. Historic fire regime in Southern California
- 507 shrublands. *Conservation Biology*, 15, 1536-1548.
- KEELEY, J. E. & KEELEY, S. C. 1981. Post-fire regeneration of southern California chaparral.
 American Journal of Botany, 68, 524-530.
- 510 KRAWCHUK, M. & MORITZ, M. 2012. *Fire and climate change in California: changes in the*
- 511 *distribution and frequency of fire in climates of the future and recent past (1911-2099),*
- 512 California Energy Commission.
- 513 MCKELVEY, K. S., SKINNER, C. N., CHANG, C.-R., ERMAN, D. C., HUSARI, S. J.,
- 514 PARSONS, D. J., VAN WAGTENDONK, J. W. & WEATHERSPOON, C. P. An
- 515 overview of fire in the Sierra Nevada. Pages 1033-1040 in: Sierra Nevada Ecosystem
- 516 Project, Final Report to Congress, Vol. II, Assessments and Scientific Basis for
- 517 Management Options. Davis, CA: University of California, Centers for Water and
- 518 Wildland Resources. Report No. 37., 1996.
- 519 MCPHERSON, J. K. & MULLER, C. H. 1967. Light competition between Ceanothus and Salvia
- 520 shrubs. *Bulletin of the Torrey Botanical Club*, 41-55.

521	MEENTEMEYER	, R. K	K. & MOODY,	A. 2002. Distribution of p	ant life history types in
		/	,		

- 522 California chaparral: the role of topographically-determined drought severity. *Journal of*523 *Vegetation Science*, 13, 67-78.
- 524 MENG, R., DENNISON, P. E., D'ANTONIO, C. M. & MORITZ, M. A. 2014. Remote Sensing
- Analysis of Vegetation Recovery following Short-Interval Fires in Southern California
 Shrublands. *Plos One*, 9.
- 527 MILLER, P. & POOLE, D. 1983. The influence of annual precipitation, topography, and
- vegetative cover on soil moisture and summer drought in southern California. *Oecologia*,
 56, 385-391.
- MOONEY, H. A. & PARSONS, D. J. 1973. Structure and function of the California chaparral—
 an example from San Dimas. *Mediterranean type ecosystems*. Springer.
- 532 MULLER, C. H., HANAWALT, R. B. & MCPHERSON, J. K. 1968. Allelopathic control of
- herb growth in the fire cycle of California chaparral. *Bulletin of the Torrey Botanical Club*, 225-231.
- 535 MYERS, D. & PITKIN, J. 2013. The Generational Future of Los Angeles: Projections to 2030
- and Comparisons to Recent Decades. *California Demographic Futures*. USC Sol Price
 School for Public Policy.
- NOWACKI, G. J. & ABRAMS, M. D. 2008. The demise of fire and "mesophication" of forests
 in the eastern United States. *BioScience*, 58, 123-138.
- 540 PARK, I. W., HOOPER, J., FLEGAL, J. M. & JENERETTE, G. D. 2018. Impacts of climate,
- 541 disturbance and topography on distribution of herbaceous cover in Southern California
- 542 chaparral: Insights from a remote-sensing method. *Diversity and Distributions*, 24, 497-
- 543 508.

544	PETERSON, S. & STOW, D. 2003. Using multiple image endmember spectral mixture analysis
545	to study chaparral regrowth in southern California. International Journal of Remote
546	Sensing, 24, 4481-4504.
547	PIÑOL, J., TERRADAS, J. & LLORET, F. 1998. Climate warming, wildfire hazard, and
548	wildfire occurrence in coastal eastern Spain. Climatic change, 38, 345-357.
549	POLADE, S. D., GERSHUNOV, A., CAYAN, D. R., DETTINGER, M. D. & PIERCE, D. W.
550	2017. Precipitation in a warming world: Assessing projected hydro-climate changes in
551	California and other Mediterranean climate regions. Scientific reports, 7, 1-10.
552	POOLE, D. & MILLER, P. 1981. The distribution of plant water stress and vegetation
553	characteristics in southern California chaparral. American Midland Naturalist, 32-43.
554	PRATT, R. B., JACOBSEN, A. L., RAMIREZ, A. R., HELMS, A. M., TRAUGH, C. A.,
555	TOBIN, M. F., HEFFNER, M. S. & DAVIS, S. D. 2014. Mortality of resprouting
556	chaparral shrubs after a fire and during a record drought: physiological mechanisms and
557	demographic consequences. Global change biology, 20, 893-907.
558	SCHLESINGER, W. H. & GILL, D. S. 1978. Demographic studies of the chaparral shrub,
559	Ceanothus megacarpus, in the Santa Ynez Mountains, California. Ecology, 59, 1256-
560	1263.
561	STOREY, E. A., STOW, D. A., O'LEARY, J. F., DAVIS, F. W. & ROBERTS, D. A. 2021.
562	Does short-interval fire inhibit postfire recovery of chaparral across southern California?
563	Science of The Total Environment, 751, 142271.
564	STOREY, E. A., STOW, D. A., ROBERTS, D. A., O'LEARY, J. F. & DAVIS, F. W. 2019.
565	Evaluating Drought Impact on Postfire Recovery of Chaparral Across Southern
566	California. Ecosystems.

- 567 STOREY, E. A., STOW, D. A., ROBERTS, D. A., O'LEARY, J. F. & DAVIS, F. W. 2020.
- 568 Evaluating Drought Impact on Postfire Recovery of Chaparral Across Southern
 569 California. *Ecosystems*.
- 570 SYPHARD, A. D., BRENNAN, T. J. & KEELEY, J. E. 2018a. Chaparral landscape conversion
- 571 in southern California. *Valuing Chaparral*. Springer.
- 572 SYPHARD, A. D., BRENNAN, T. J. & KEELEY, J. E. 2019a. Drivers of chaparral type
- 573 conversion to herbaceous vegetation in coastal Southern California. *Diversity and*574 *Distributions*, 25, 90-101.
- 575 SYPHARD, A. D., BRENNAN, T. J. & KEELEY, J. E. 2019b. Extent and drivers of vegetation
- 576 type conversion in Southern California chaparral. *Ecosphere*, 10, e02796.
- 577 SYPHARD, A. D., CLARKE, K. C. & FRANKLIN, J. 2007a. Simulating fire frequency and
- 578 urban growth in southern California coastal shrublands, USA. *Landscape Ecology*, 22,
 579 431-445.
- 580 SYPHARD, A. D., FRANKLIN, J. & KEELEY, J. E. 2006. Simulating the effects of frequent
- 581 fire on southern California coastal shrublands. *Ecological Applications*, 16, 1744-1756.
- 582 SYPHARD, A. D. & KEELEY, J. E. 2016. Historical reconstructions of California wildfires
 583 vary by data source. *International Journal of Wildland Fire*, 25, 1221-1227.
- 584 SYPHARD, A. D., RADELOFF, V. C., KEELEY, J. E., HAWBAKER, T. J., CLAYTON, M.
- 585 K., STEWART, S. I. & HAMMER, R. B. 2007b. Human influence on California fire 586 regimes. *Ecological Applications*, 17, 1388-1402.
- 587 SYPHARD, A. D., SHEEHAN, T., RUSTIGIAN-ROMSOS, H. & FERSCHWEILER, K.
- 588 2018b. Mapping future fire probability under climate change: Does vegetation matter?
 589 *PLoS One*, 13, e0201680.

- 590 THOMAS, C. & DAVIS, S. 1989. Recovery patterns of three chaparral shrub species after
 591 wildfire. *Oecologia*, 80, 309-320.
- 592 VAN DE WATER, K. M. & SAFFORD, H. D. 2011. A summary of fire frequency estimates for
- 593 California vegetation before Euro-American settlement. *Fire Ecology*, 7, 26-58.
- 594 VENTURAS, M. D., MACKINNON, E. D., DARIO, H. L., JACOBSEN, A. L., PRATT, R. B.
- 595 & DAVIS, S. D. 2016. Chaparral shrub hydraulic traits, size, and life history types relate
- to species mortality during California's historic drought of 2014. *PloS one*, 11, e0159145.
- 597 ZAMMIT, C. A. & ZEDLER, P. H. 1993. Size Structure and Seed Production in Even-Aged
- 598 Populations of Ceanothus Greggii in Mixed Chaparral. *Journal of Ecology*, 81, 499-511.
- 599 ZEDLER, P. H. 1995. Fire frequency in southern California shrublands: biological effects and
- 600 management options. Brushfires in California wildlands: ecology and resource
- 601 management. International Association of Wildland Fire, Fairfield, Washington, USA,
- 602 101-112.

603 ZEDLER, P. H., GAUTIER, C. R. & MCMASTER, G. S. 1983. Vegetation change in response

- 604 to extreme events: the effect of a short interval between fires in California chaparral and 605 coastal scrub. *Ecology*, 64, 809-818.
- 606
- 607
- 608
- 609
- 610
- 611
- 612

- **Table 1** Linear mixed-effects model results for all vegetation type covers at the plot level (N =
- 614 198). Each model had site as a random effect and was analyzed using the residuals of pre/post-

fire vegetation percent cover.

TT .	Model AIC					
Vegetation Class	Burn Interval	Aspect	Years After Fire	Distance to Coast	Five-year Average Rainfall	
chaparral	1519.907	1515.694*	1514.127*	1519.806	1519.095	
sage scrub	1525.744	1522.241*	1522.682*	1527.828	1527.542	
grass	1296.388	1296.534	1295.17*	1296.466	1296.417	
tree	1160.116*	1164.776	1164.47	1164.837	1164.873	

- 617 * indicates best model(s)

- -

631 Figure legends

632

- 633 Figure 1 Minimum fire intervals as reported by CalFire (frap.fire.ca.gov) for Ventura and Los
- Angeles counties from 1878 to 2009. Twelve sites where one polygon burned twice within five
- 635 years (twice-burned, black) and an adjacent polygon burned once within the same five-year
- 636 period (once-burned, white).

637

638 Figure 2 Linear regressions with confidence intervals of pre-fire and post-fire percent cover for

all four vegetation type covers. Each point represents a once-burned (gray) or twice-burned

640 (black) plot.

641

642 Figure 3 Linear regressions with confidence intervals of time since last fire with residuals (left

643 side) for chaparral ($P < 0.001^{***}$, $r^2 = 0.05$), sage shrub ($P = 0.005^{**}$, $r^2 = 0.03$), and grass (P =

644 0.16, $r^2 = 0.005$) cover. Distribution of the residuals (right side) of chaparral (ANOVA, P =

645 0.114), sage scrub (ANOVA, $P = 0.046^*$), and grass (ANOVA, P = 0.886) cover by aspect.

646

Figure 4 Distribution of residuals for tree cover (ANOVA, $P = 0.03^*$) by burn interval. When the outlier of tree percent lost in the twice-burned burn interval is removed, the relationship is still significant (ANOVA, P value = 0.048*).

650

Figure 5 Transitions for all pixels showing pre-fire (left axis) and pos-tfire (right axis) cover within (a) once-burned plots and (b) twice-burned plots. Gray bands depict the proportion of pixels that remained or transitioned to another cover type.



Figure 1 Minimum fire intervals as reported by CalFire (frap.fire.ca.gov) for Ventura and Los
Angeles counties from 1878 to 2009. Twelve sites where one polygon burned twice within five
years (twice-burned, black) and an adjacent polygon burned once within the same five-year
period (once-burned, white).



Figure 2 Linear regressions with confidence intervals of pre-fire and post-fire percent cover for
all four vegetation type covers. Each point represents a once-burned (gray) or twice-burned

- 669 (black) plot.
- 670

- 671
- 672





Figure 3 Linear regressions with confidence intervals of time since last fire with residuals (left side) for chaparral ($P < 0.001^{***}$, $r^2 = 0.05$), sage shrub ($P = 0.005^{**}$, $r^2 = 0.03$), and grass (P = 0.16, $r^2 = 0.005$) cover. Distribution of the residuals (right side) of chaparral (ANOVA, P = 0.114), sage scrub (ANOVA, $P = 0.046^{*}$), and grass (ANOVA, P = 0.886) cover by aspect.





679 **Figure 4** Distribution of residuals for tree cover (ANOVA, $P = 0.03^*$) by burn interval. When

680 the outlier of tree percent lost in the twice-burned burn interval is removed, the relationship is

681 still significant (ANOVA,
$$P$$
 value = 0.048*).



Figure 5 Transitions for all pixels showing pre-fire (left axis) and post-fire (right axis) cover

686 within (a) once-burned plots and (b) twice-burned plots. Gray bands depict the proportion of

687 pixels that remained or transitioned to another cover type.