

1 When Irrigation Cannot Keep Pace: Aridification, Crop Composition, and
2 the Spatial Concentration of Agricultural Water Use Efficiency Decline in
3 Central Chile

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6 **Abstract**

7 Water scarcity under sustained aridification is among the most consequential threats to agriculture in
8 Mediterranean-climate regions. Central Chile's ongoing megadrought (~2010-present), characterized by
9 persistent precipitation deficits and intensified atmospheric evaporative demand-provides a natural experiment
10 for examining how agricultural water use efficiency (WUE) responds to prolonged climatic stress. Using
11 remotely sensed estimates of net primary production (NPP) and evapotranspiration (ET) across the 76
12 agricultural watersheds (2001-2020), we characterize WUE dynamics along multiple dimensions of aridification
13 and assess the role of crop composition in modulating watershed-scale sensitivity.

14 Contrary to expectations that technological adaptation might buffer efficiency outcomes, we find a persistent,
15 spatially coherent decline in agricultural WUE across much of central Chile, with structural break points
16 clustering near 2010 in coincidence with megadrought onset. Hydroclimatic pressures-particularly reduced
17 water availability and elevated atmospheric demand-appear to have outweighed any efficiency gains from
18 improved irrigation practices, suggesting that under sustained drought, water and energy limitations constrain
19 agricultural productivity at regional scales beyond the reach of field-level adaptation. Precipitation deficit
20 was the dominant aridification signal explaining WUE variability, with atmospheric demand contributing
21 modest independent explanatory power. Crop composition explained more than half of the spatial variance
22 in drought sensitivity, with cereal-dominated systems showing lower WUE-SPEI sensitivity relative to forage
23 systems, while irrigated area share was positively associated with drought sensitivity.

24 These results challenge the assumption that irrigation efficiency improvements can adequately buffer
25 agriculture against intensifying hydroclimatic stress, and underscore the need for water governance frameworks

26 that operate at the landscape and regional scale.

27 *Keywords:* water use efficiency, megadrought, aridification, SPEI, Chile, NPP/ET, crop composition

28 1. Introduction

29 Global agriculture consumes approximately 70% of freshwater withdrawals (Hoekstra and Mekonnen, 2012),
30 yet water use efficiency (WUE) remains poorly quantified at regional scales under conditions of sustained
31 climatic stress. As aridification intensifies across dryland and Mediterranean-climate regions (Gebrechorkos
32 et al., 2025), understanding how agricultural WUE responds to different dimensions of water deficit becomes
33 critical for adaptive water governance (FAO y ONU Agua, 2025; Howden et al., 2007; Wallace, 2000).

34 Chile provides an unusually compelling natural laboratory for this question. Since approximately 2010,
35 central and southern Chile has experienced a persistent precipitation deficit that has been characterized as a
36 megadrought—a multi-year dry anomaly that exceeds the interannual variability expected from modes of
37 climate variability alone (Garreaud et al., 2017, 2020). The megadrought has reduced annual precipitation by
38 20-40% relative to the twentieth-century mean, depleted streamflows, and lowered groundwater levels across
39 the country’s most productive agricultural zones (Boisier et al., 2018). Parallel to precipitation reduction,
40 atmospheric evaporative demand has intensified as warming has elevated vapor pressure deficit (VPD),
41 compounding water stress beyond what precipitation-only indices capture (Li et al., 2023; Zambrano et al.,
42 2025).

43 Despite the severity of this hydroclimatic disruption, characterized by 20-40% precipitation reductions
44 relative to twentieth-century means, streamflow depletion, and intensified atmospheric evaporative demand
45 through elevated vapor pressure deficit (VPD), the agricultural response, specifically in terms of WUE, has
46 not been systematically characterized. WUE provides a systems-level signal of how efficiently agricultural
47 land converts water into biomass under changing conditions (Hoover et al., 2023; Ito and Inatomi, 2012; Jiang
48 et al., 2025). While Garreaud et al. (2017), Garreaud et al. (2020), and Boisier et al. (2018) document the
49 megadrought’s precipitation and hydrological impacts across Chile’s productive zones, no studies have been
50 conducted that provide a detailed analysis of cropland WUE dynamics under multidimensional aridification.
51 Global meta-analyses reveal drought typically increases crop WUE by ~1.5% through physiological adjustments

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52 (Yu et al., 2020), yet these effects vary unpredictably by crop type, drought timing, and duration without
53 regional mechanistic breakdowns (e.g., NPP vs. ET drivers). Changes in WUE during prolonged drought may
54 reflect multiple processes operating simultaneously: stomatal regulation under vapor pressure deficit stress,
55 shifts in phenology, compositional changes in cropping systems, and the partial buffering of irrigated systems
56 against precipitation deficits (Wang et al., 2024). Mediterranean irrigation studies estimate 35% potential
57 water savings via efficient systems but rarely quantify WUE responses to combined precipitation deficits
58 and VPD intensification in export-oriented perennial systems (Fader et al., 2016). Regional assessments in
59 Latin America report low agricultural water utilization (35-86%) but fail to disentangle drought-specific
60 drivers or crop-composition effects on sensitivity (Mekonnen and Hoekstra, 2011). This leaves critical gaps
61 in understanding how Chile’s export-oriented perennial horticulture responds to sustained megadrought
62 conditions.

63 Agricultural crop composition is likely to modulate the WUE response to aridification. Perennial fruit crops
64 (vineyards, orchards), which dominate the high-value export agriculture of central Chile, have different water
65 use strategies compared with annual grain and forage crops. Perennials maintain year-round root systems
66 and may sustain higher net primary production (NPP) under deficit conditions through deep root access,
67 whereas annuals experience sharper productivity collapses once critical thresholds are breached (Chakraborti
68 et al., 2023; Davis et al., 2017). The degree to which crop composition moderates WUE sensitivity to
69 aridification signals is investigated here using agricultural census data linked to watershed-level WUE and
70 drought records.

71 A key conceptual contribution of this paper is to treat aridification as a multi-process phenomenon rather
72 than a single drought signal. Previous analyses have often used a single index, typically a precipitation-based
73 measure, to characterize drought impacts on vegetation. However, SPEI (Standardized Precipitation-
74 Evapotranspiration Index, Vicente-Serrano et al. (2010)) captures the combined effect of precipitation deficits
75 and elevated atmospheric demand, SPI (Standardized Precipitation Index, Mckee et al. (1993)) isolates
76 precipitation deficits, and EDDI (Evaporative Demand Drought Index, Hobbins et al. (2016), McEvoy et al.
77 (2016)) captures anomalies in atmospheric evaporative demand independently of precipitation. Whether WUE
78 responds differently to these distinct aridification dimensions has not been established for Chilean agriculture.
79 A methodologically distinctive feature of this study is the use of a paired-census quasi-experimental design
80 to disentangle structural crop-composition effects from drought-driven land-use adaptation. By comparing
81 crop-composition-sensitivity relationships estimated from the 2007 (pre-drought) and 2021 (post-drought)

82 agricultural censuses, and by fitting spatial error models that account explicitly for the geographic clustering of
83 crop systems alongside aridification exposure, the study can assess whether elevated WUE-drought sensitivity
84 in particular watershed types reflects an enduring structural property of those crop systems or a post-drought
85 artefact of compositional change. This integration of paired-census design with spatial statistical inference,
86 rather than treating OLS crop-composition coefficients as causally independent, constitutes the primary
87 methodological contribution of the study alongside its multi-dimensional aridification framework.

88 This paper addresses the following primary question: How does agricultural WUE respond to different
89 dimensions of aridification during the Central Chile Megadrought? Secondary questions ask: (i) Do zNPP
90 (NPP z-score) and SETI (Standardized Evapotranspiration Index) trends reveal whether observed WUE
91 change is driven primarily by declining vegetation productivity or by changes in actual evapotranspiration
92 (ET)? (ii) Did WUE undergo a structural break coincident with megadrought onset around 2010? (iii) Which
93 long-term aridification signal, SPEI, SPI, or EDDI, explains the greatest share of WUE variability? (iv) Does
94 crop composition modulate WUE sensitivity to aridification? (v) Are WUE responses spatially clustered
95 across Chilean watersheds?

96 **2. Data and Methods**

97 *2.1. Study area*

98 Continental Chile extends from approximately 17°S to 56°S, encompassing some of the world’s most extreme
99 latitudinal climate gradients within a single country (Aceituno et al., 2021). The northern Atacama region is
100 hyperarid; the semi-arid Norte Chico (approximately 26°S-32°S) transitions into the Mediterranean-climate
101 zone of central Chile (32°S-38°S), which concentrates the majority of the country’s irrigated agriculture,
102 viticulture, and fruit export production. South of the Biobío River (~38°S), a temperate humid zone supports
103 rainfed cereal and forage production transitioning to temperate rainforests further south (Figure 1).

104 The megadrought has most severely affected the 30°S-42°S band, where precipitation deficits of 25-45%
105 relative to the twentieth-century mean have persisted since 2010 (Boisier et al., 2018; Garreaud et al., 2020).
106 The study uses watershed-level spatial units based on the Chilean National Water Authority (DGA, General
107 Directorate of Water) hydrological boundaries, which provide physically meaningful aggregation units and
108 align with the country’s water allocation governance framework (Rivera et al., 2016).

109 *2.2. WUE data*

110 Annual cropland WUE expressed in $g \cdot C \cdot kg^{-1} H_2O \cdot^{-1}$, is defined as:

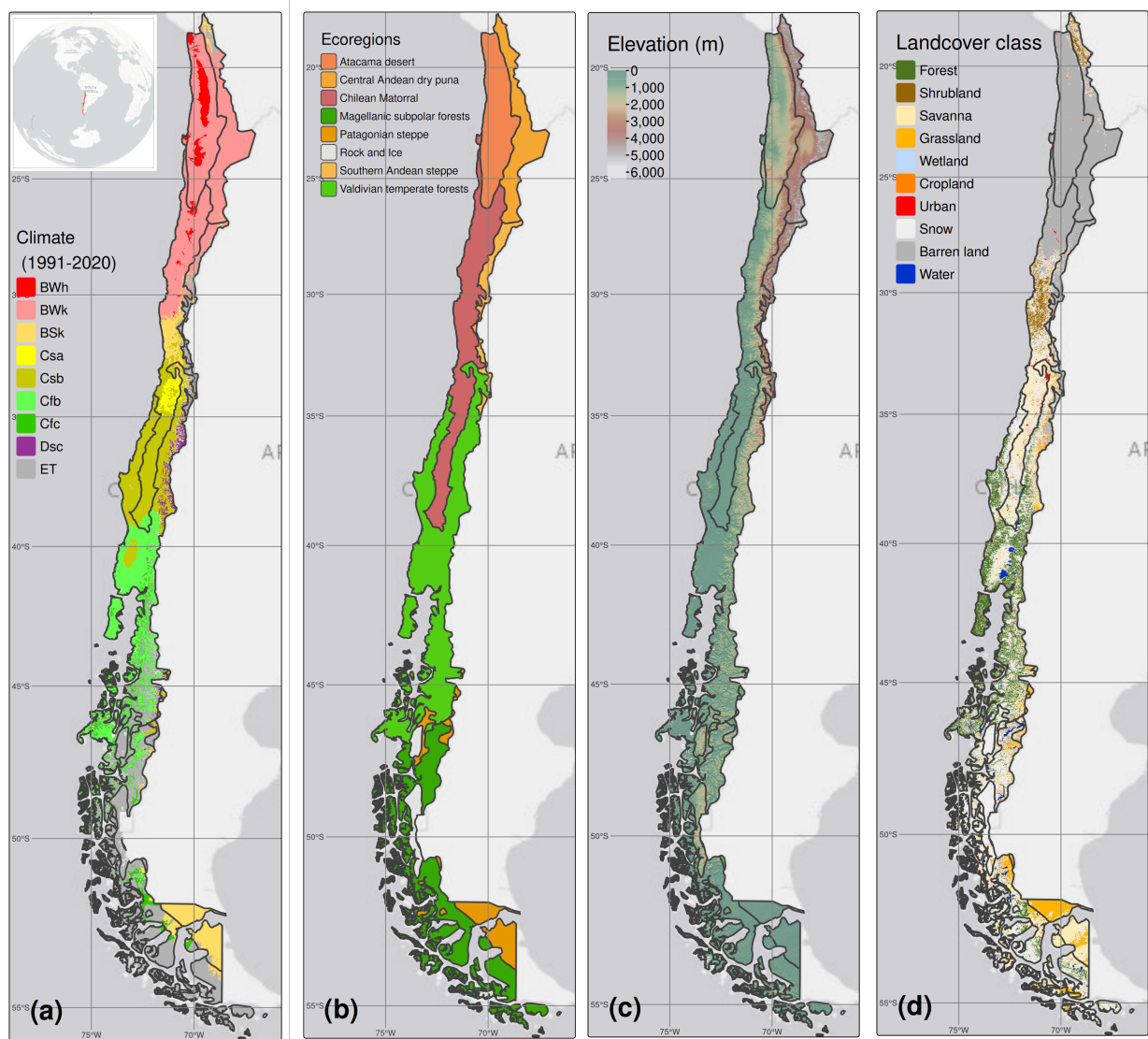


Figure 1: Climate, topography, and land cover classes across continental Chile. Koppen-Geiger climate classes (a), ecoregions (b), topography (c), and persistent land cover classes (> 80%) for 2001-2023 (d) across continental Chile. Reproduced from Zambrano et al. (2025) under CC BY 4.0 license.

$$WUE = \frac{NPP}{ET}$$

111 It was obtained from the global cropland WUE dataset of [Jiang et al. \(2025\)](#), which provides 1-km grid
 112 annual rasters for 2001-2020 derived specifically for agricultural areas. NPP was estimated using an improved
 113 Evaporative Fraction Light-Use Efficiency (EF-LUE) model driven by ERA5-Land reanalysis meteorology
 114 (temperature, solar radiation, vapour pressure deficit; bilinearly resampled from the native ~9 km resolution
 115 to 1 km as described in [Jiang et al. \(2025\)](#)), GLASS fAPAR and LAI (500 m, 8-day; interpolated to daily),
 116 and cropland distribution from ESA CCI-LC land cover (300 m, aggregated to 1 km). The EF-LUE model
 117 parameterises stomatal water stress via an evaporative fraction term calibrated against FLUXNET2015 flux
 118 tower observations across multiple Köppen-Geiger climate zones. Annual NPP was computed as:

$$NPP = 0.8 \times \sum_d GPP_d - \sum_d MR_d$$

119 where MR_d is temperature-dependent maintenance respiration of leaves and fine roots. ET was sourced
 120 from the ETMonitor product (1 km, daily, [Hu and Jia \(2015\)](#)), which integrates energy balance, water balance,
 121 and plant physiological processes and was validated against flux towers with correlation > 0.75 . Rasters
 122 were cropped to Chile's extent (76°W-66.24°W, 56.54°S-17.49°S), then monthly, and annual mean WUE was
 123 extracted per watershed by spatial averaging. The 20-year time series (2001-2020) spans a pre-megadrought
 124 reference period (2001-2009) and the megadrought period (2010-2020).

125 WUE values were aggregated as arithmetic means across cropland pixels within each watershed boundary
 126 ([Hoover et al., 2023](#); [Jiang et al., 2025](#)).

127 *2.3. Drought drivers and WUE mechanistic components*

128 The variables used in this study serve two analytically distinct roles. Drought drivers (SPEI, SPI, EDDI)
 129 characterize the external aridification forcing imposed on agricultural systems. WUE mechanistic components
 130 (zNPP, SETI) track the internal responses within the WUE identity, representing the numerator (NPP) and
 131 denominator (ET) separately, and are therefore not treated as additional drought indices.

132 *2.3.1. Aridification drivers*

133 Three complementary drought indices were used to characterize distinct dimensions of aridification, following
134 the framework established in [AghaKouchak et al. \(2015\)](#), [Gebrechorkos et al. \(2023\)](#), and [Zambrano et al.
135 \(2025\)](#):

136 **SPEI (Standardized Precipitation-Evapotranspiration Index, 36-month scale)** represents the
137 primary aridification driver. SPEI captures the hydroclimatic balance by incorporating both precipitation
138 inputs and atmospheric evaporative demand into a standardized anomaly, making it sensitive to the combined
139 effect of precipitation deficit and VPD intensification that characterizes the megadrought ([Li et al., 2023](#);
140 [Zambrano et al., 2025](#)). SPEI was calculated from ERA5-Land monthly precipitation, and ET was calculated
141 based on FAO-56 Penman-Monteith reference evapotranspiration following [Vicente-Serrano et al. \(2010\)](#).

142 **SPI (Standardized Precipitation Index, 36-month scale)** isolates precipitation deficits, providing a
143 comparison signal that reflects the purely pluviometric dimension of drought without evaporative demand
144 effects ([Mckee et al., 1993](#)).

145 **EDDI (Evaporative Demand Drought Index, 36-month scale)** characterizes anomalies in atmospheric
146 evaporative demand independently of precipitation, thereby capturing the thermally driven component of
147 aridification ([Hobbins et al., 2016](#); [McEvoy et al., 2016](#)). The EDDI implementation follows a sign convention
148 aligned with SPEI and SPI: positive values indicate below-normal evaporative demand (wet-like conditions)
149 and negative values indicate above-normal demand (drought-like conditions). This convention inverts the
150 original [Hobbins et al. \(2016\)](#) polarity and allows direct cross-index comparison with a shared interpretation
151 (negative = drought).

152 All three indices were computed at the 36-month accumulation scale to capture long-term signals of
153 aridification ([Zambrano et al., 2025](#)), regularly associated with hydrological impacts. Monthly values were
154 extracted per watershed polygon and aggregated to annual means before analysis.

155 *2.3.2. WUE mechanistic components*

156 Two variables were incorporated to decompose WUE variability in terms of the identity $WUE = NPP/ET$,
157 providing insight into whether observed WUE changes are driven by the productivity (NPP) or water use
158 (ET) side of the ratio.

159 **zNPP (z-score NPP)** tracks the NPP numerator of WUE directly. Annual NPP estimates from

160 MOD17A3HGF (MODIS Terra, 500 m, Collection 6.1) were z-score standardised within the 2001-2020
161 reference period, yielding an anomaly series that reflects interannual variability in vegetation productivity
162 relative to the historical mean.

163 **SETI-36 (Standardized Evapotranspiration Index, 36-month scale)** captures multi-year anomalies
164 in actual evapotranspiration relative to the historical baseline, representing the ET denominator of WUE.
165 Drought-induced reductions in actual ET reflect the combined effect of stomatal closure, reduced soil water
166 availability, and phenological shifts. The imbalance between EDDI (atmospheric evaporative demand) and
167 SETI (actual evapotranspiration) identifies watersheds where potential demand increasingly exceeds actual
168 water use, a signal of progressive vegetation-atmosphere decoupling and physiological water stress. SETI
169 was derived from MOD16A2GF (MODIS Terra, 500m 8-day, Collection 6.1) actual evapotranspiration data
170 (Running et al., 2021). Because zNPP (MOD17A3HGF) and SETI (MOD16A2GF) are derived from products
171 independent of the ETMonitor/EF-LUE pipeline used by Jiang et al. (2025) to construct the WUE rasters,
172 this mechanistic decomposition functions as a cross-product consistency check rather than an exact algebraic
173 decomposition of the WUE numerator and denominator.

174 *2.4. Agricultural census data*

175 Crop composition data were obtained from two Chilean Agricultural and Forestry Censuses: the VII Census
176 (2007, pre-drought baseline) (INE, 2007) and the VIII Census (2020-2021, endpoint) (INE, 2021). Both
177 censuses report cultivated area (ha) by crop category and management type (irrigated/rainfed) at commune
178 level. Crop categories included: cereals, legumes and tubers, industrial crops, vegetables and aromatics,
179 fruit trees, vineyards and pisco grapes, cut flowers, seed crops, nurseries, and forage crops. Categories were
180 harmonised across census editions to ensure comparability.

181 Raw census data were processed to assign commune identifiers by normalizing names and cross-referencing
182 with spatial boundary databases, including manual correction of combined entries for communes grouped for
183 statistical confidentiality. Crop areas were preserved separately by irrigation condition (irrigated, rainfed) as
184 reported directly in the census. Where the irrigation split was not reported directly, it was imputed from
185 subsidiary files using residual ratios and flagged accordingly. Of the 3,807 crop-category records in the 2007
186 census long-format dataset, 2,535 (66.6%) used imputed irrigation splits; the 2021 census had no imputed
187 records (all irrigation splits directly reported). Imputed values preserve national agricultural totals but
188 may misallocate irrigation at the individual commune level, potentially introducing attenuation bias in the
189 percentage irrigated area regression coefficient for the 2007 analysis. Given that the 2007 baseline is used

190 only for directional consistency checks rather than primary inference, this limitation does not affect the main
191 crop composition results. For the 2021 analysis, which underpins the primary SEM results, the irrigated area
192 share variable is derived entirely from directly reported data. Commune-level data were spatially aggregated
193 to watershed level using area-weighted means via a spatial join between commune boundaries and watershed
194 polygons. Each watershed therefore receives a crop composition vector reflecting the agricultural structure of
195 the communes it contains.

196 From the per-watershed crop areas, the following proportional indicators were derived:

- 197 • *pct_fruits*: proportion of area under fruit trees
- 198 • *pct_vineyards*: proportion under vineyards and pisco grapes
- 199 • *pct_cereals*: proportion under cereals
- 200 • *pct_vegetables*: proportion under vegetables and aromatics
- 201 • *pct_forages*: proportion under forage crops
- 202 • *pct_irrigated*: irrigated share of total agricultural area (irrigated / (irrigated + rainfed)), reflecting the
203 buffering capacity of irrigation infrastructure against precipitation deficits

204 Three functional aggregates were computed: perennial horticultural crops ($pct_perennials = pct_fruits$
205 $+ pct_vineyards$), annual cropland ($pct_annuals = pct_cereals + pct_vegetables$), and forage/pasture
206 ($pct_forage = pct_forages$). Dominant crop type was assigned to each watershed using a 40% area threshold:
207 if any functional group exceeded 40% of total agricultural area, the watershed was classified accordingly;
208 otherwise it was classified as mixed.

209 2.5. Statistical analyses

210 2.5.1. Trend analysis

211 Mann-Kendall trend tests and Sen's slope estimates were computed per watershed using the annual WUE
212 time series (2001-2020), which accounts for autocorrelation through prewhitening. Equivalent analyses
213 were performed for zNPP trends, SETI trends, and atmospheric evaporative demand (EDDI) to provide
214 comparative multi-variable context.

215 2.5.2. Structural break detection

216 The Pettitt test (Pettitt, 1979) was applied to the annual WUE series of each watershed to detect the
217 most probable year of a shift in the mean level. The Pettitt test is a non-parametric rank-based method
218 that does not require distributional assumptions and is widely used for hydrological change-point detection.

219 Benjamini-Hochberg (BH) false-discovery-rate (FDR) correction was applied to Pettitt structural break
220 p-values (76 simultaneous watershed-level tests). Watersheds were classified by: (i) significance of the detected
221 break ($p < 0.05$), and (ii) proximity to megadrought onset, defined as break years within ± 2 years of 2010
222 (i.e., 2008-2012).

223 2.5.3. Before/after comparison

224 Mean WUE was computed separately for the pre-megadrought (2001-2009) and megadrought (2010-2020)
225 periods per watershed. The difference $\Delta WUE = WUE_{post} - WUE_{pre}$ was mapped spatially. Wilcoxon
226 signed-rank tests (paired by spatial unit) tested the significance of the mean shift at each spatial aggregation
227 level. Distributions were visualized as violin-boxplot combinations faceted by ecoregion, using the WWF
228 Terrestrial Ecoregions of the World classification (Dinerstein et al., 2017) to stratify results by ecological
229 context.

230 2.5.4. WUE-aridification relationships

231 Spearman rank correlations (ρ) were computed between annual WUE and each of the three indices (SPEI,
232 SPI, EDDI), applied per watershed across the 20-year time series. Per-watershed Spearman correlation and
233 OLS regression results are reported without FDR correction, as each watershed p-value describes a distinct
234 independent time series rather than constituting a family of tests on the same null hypothesis. Contemporaneous
235 linear regressions ($WUE \sim \text{Index}$) were fitted per watershed to estimate the slope coefficient, which
236 we interpret as the local sensitivity of WUE to each aridification signal ($Sensitivity = \Delta WUE / \Delta Index$).
237 Spearman ρ values were mapped spatially and the mean ρ across watersheds was plotted as a function of
238 each index. Sensitivity maps were produced for SPEI as the primary index.

239 A two-way fixed-effects panel regression model was estimated:

$$WUE_{it} = \beta \cdot Index_{it} + \mu_i + \tau_t + \varepsilon_{it} \quad (1)$$

240 where μ_i represents watershed fixed effects and τ_t represents year fixed effects, thereby controlling for
241 unobserved time-invariant differences among watersheds and common annual shocks. An interaction model was
242 additionally estimated to test whether within-watershed WUE–drought sensitivity changed after megadrought
243 onset:

$$WUE_{it} = \beta_1 \cdot Index_{it} + \beta_2 \cdot Index_{it} \times \mathbb{1}(year \geq 2010) + \mu_i + \tau_t + \varepsilon_{it} \quad (2)$$

244 *2.5.5. Spatial dependence analysis*

245 Moran’s I was computed for WUE trend slope, Pettitt break year, and R^2 of the WUE-SPEI relationship,
 246 using queen contiguity spatial weights. Getis-Ord Gi^* local statistics were computed for WUE trend slope to
 247 identify spatial hotspots ($Gi^* > 1.96$) and coldspots ($Gi^* < -1.96$) at the 95% significance level. Queen
 248 contiguity was used as the primary weighting scheme because it preserves physical adjacency between
 249 watershed polygons.

250 *2.5.6. Crop composition and WUE sensitivity*

251 The per-watershed OLS slope of WUE regressed on SPEI ($\Delta WUE/\Delta SPEI$) was joined with census-derived
 252 crop composition data, and a multiple OLS regression was fitted:

$$Sensitivity_i = \beta_0 + \beta_1 \cdot pct_fruits_i + \beta_2 \cdot pct_vineyards_i + \beta_3 \cdot pct_cereals_i \quad (3)$$

$$+ \beta_4 \cdot pct_vegetables_i + \beta_5 \cdot pct_irrigated_i + \varepsilon_i$$

253 Forage and pasture proportions (*pct_forages*) served as the reference category and were omitted from
 254 the model to avoid the dummy variable trap; regression coefficients therefore express the change in WUE
 255 sensitivity relative to a watershed whose agricultural area is entirely under forage or pasture. Variance
 256 inflation factors (VIF) were computed to verify the absence of harmful multicollinearity (threshold: $VIF > 5$).
 257 The variable *pct_irrigated* was included as a covariate to control for the buffering effect of irrigation access
 258 on the WUE-drought relationship. Because WUE trends exhibit strong spatial autocorrelation (Moran’s I =
 259 0.627), Moran’s I was also computed on the OLS residuals using queen-contiguity spatial weights; where
 260 residuals were spatially autocorrelated ($p < 0.05$), a spatial error model was estimated and reported alongside
 261 OLS results.

262 A sensitivity regression using pre-drought (2007) census proportions as predictors was also estimated to
 263 address temporal confounding concerns: if results are directionally consistent with the 2021 regression, the
 264 crop composition-sensitivity relationship can be interpreted as a structural property rather than an artefact
 265 of drought-driven adaptation (see Section 4.3 and Supplementary Figure S-1).

266 2.6. Software

267 All statistical analyses and spatial processing were conducted in R (v4.5.2; R Core Team (2025)). Raster
268 operations and spatial aggregation used `terra` (v1.8.54; Hijmans (2025)). Vector geometries and spatial
269 joins used `sf` (v1.0.23; Pebesma (2018), Pebesma and Bivand (2023)). Spatiotemporal array operations used
270 `stars` (v0.6.8; Pebesma and Bivand (2023)). Mann-Kendall trend tests and Sen’s slope estimation with
271 prewhitening used `modifiedmk` (v1.6; Patakamuri and O’Brien (2021)) and `trend` (v1.1.6; Pohlert (2023)).
272 Spatial autocorrelation (Moran’s I, Getis-Ord G_i^*) and spatial weight matrices used `spdep` (v1.4.1; Bivand
273 and Wong (2018)). Spatial Error Model was made using `{spatialreg}` (Bivand et al., 2021). Panel regression
274 with fixed effects used `plm` (Croissant and Millo, 2008). Variance inflation factors used `car` (v3.1.3; Fox and
275 Weisberg (2019)). General-purpose data manipulation and reshaping used `tidyverse` (v2.0.0; Wickham
276 et al. (2019)), including `dplyr`, `tidyr`, `readr`, `stringr`, `purrr`, and `lubridate`. Thematic maps used `tmap`
277 (v4.2; Tennekes (2018)). Statistical graphics used `ggplot2` (v3.5.2; Wickham (2016)). Multi-panel figure
278 layout used `patchwork` (v1.3.1; Pedersen (2025)).

279 3. Results

280 3.1. Long-term WUE trends

281 Trend analysis across $n = 76$ Chilean agricultural watersheds with complete WUE data revealed spatially
282 heterogeneous WUE dynamics over the 2001-2020 period (Figure 2). WUE trends followed a predominantly
283 negative pattern across the Mediterranean-climate zone of central Chile (approximately 30°S-35°S), consistent
284 with the onset and intensification of the megadrought.

285 Decomposing WUE trends into their mechanistic components, $zNPP$ representing the NPP numerator
286 and $SETI$ representing the ET denominator, reveals the distinct contributions of productivity decline and
287 evapotranspiration change to the observed WUE dynamics. $zNPP$ trends exhibited strong spatial covariance
288 with WUE, with negative anomalies concentrated in the same central Mediterranean watersheds, indicating
289 that NPP suppression is the primary driver of WUE deterioration in those zones (Garreaud et al., 2017;
290 Zambrano et al., 2025). $SETI$ trends showed more spatially complex patterns: watersheds where actual ET
291 declined less steeply than NPP correspond to WUE decline, while watersheds where ET reduction exceeded
292 NPP reduction show stable or improving WUE.

293 Declining WUE in the Mediterranean zone is thus driven primarily by NPP suppression, while ET responses
294 are spatially heterogeneous and modulated by irrigation access and crop type.

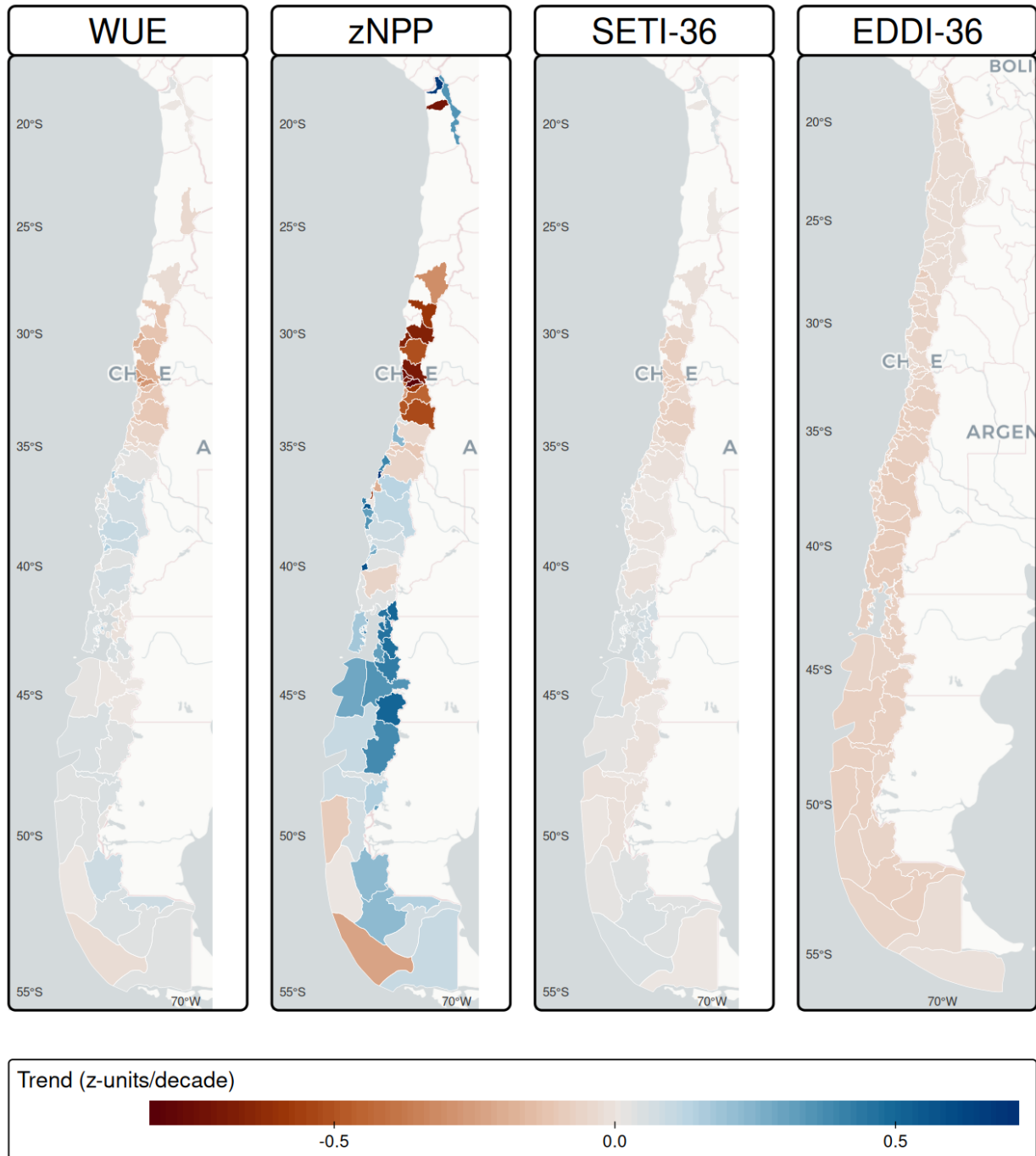


Figure 2: Spatial distribution of long-term trends in WUE, zNPP (NPP anomaly), SETI-36 (long-term ET anomaly), and EDDI-36 (long-term atmospheric evaporative demand) across Chilean watersheds (2001-2020). Trend magnitudes are expressed as z-score standardized Sen's slope estimates.

295 *3.2. Structural breaks in WUE*

296 The Pettitt test detected statistically significant structural breaks ($p < 0.05$) in 17 of 76 Chilean watersheds
297 (22.4%). Of these 17 raw-significant breaks, 12 (71%) occurred within ± 2 years of 2010 (the recognized onset
298 of the megadrought) suggesting that the megadrought is temporally associated with structural shifts in WUE
299 dynamics in a subset of watersheds (Figure 3 a). After applying Benjamini-Hochberg false-discovery-rate
300 (FDR) correction across all 76 per-watershed tests, none of the breaks survived at the 0.05 threshold (all
301 adjusted $p > 0.05$), reflecting the limited power of individual non-parametric tests applied across spatially
302 correlated watersheds with only 20 annual observations each. The geographic concentration of near-2010
303 breaks in the central Mediterranean zone — precisely where the megadrought’s hydroclimatic footprint is
304 strongest — provides qualitative corroboration that these detections are not randomly distributed in space,
305 even if they cannot be treated as statistically independent given the documented spatial autocorrelation in
306 WUE dynamics (Moran’s I for break year = 0.164, $p = 0.011$).

307 Spatially, megadrought-proximate break years were concentrated in the central and north-central Mediter-
308 ranean zone, supporting the interpretation that the megadrought represents a structural discontinuity in
309 agricultural water use efficiency for those regions. Watersheds in the extreme north (hyperarid Atacama)
310 and the wet south showed either no significant breaks or break years unrelated to the 2010 megadrought
311 onset, consistent with the spatial footprint of the megadrought as documented by Garreaud et al. (2020) and
312 Boisier et al. (2018).

313 *3.3. WUE before and after the megadrought*

314 Comparison of mean WUE between the pre-megadrought (2001-2009) and megadrought (2010-2020) periods
315 revealed a predominantly negative ΔWUE across central Chilean watersheds, indicating that the megadrought
316 period was associated with reduced agricultural water use efficiency relative to the reference period (Figure 3
317 b). The spatial pattern of ΔWUE closely mirrors the megadrought’s hydroclimatic footprint, with the largest
318 negative shifts concentrated in Mediterranean-climate watersheds where the precipitation deficit has been
319 most severe.

320 The aggregate Wilcoxon signed-rank test comparing pre-megadrought (2001-2009) and megadrought (2010-
321 2020) mean WUE across all 76 watersheds did not reach statistical significance ($p = 0.330$). This aggregate
322 non-significance reflects the spatial heterogeneity of WUE responses across ecologically diverse watersheds:
323 pooling Mediterranean-climate zones (where declines are concentrated) with southern humid zones (where
324 WUE responses are weaker or mixed) attenuates the overall effect. Ecoregion-stratified Wilcoxon tests

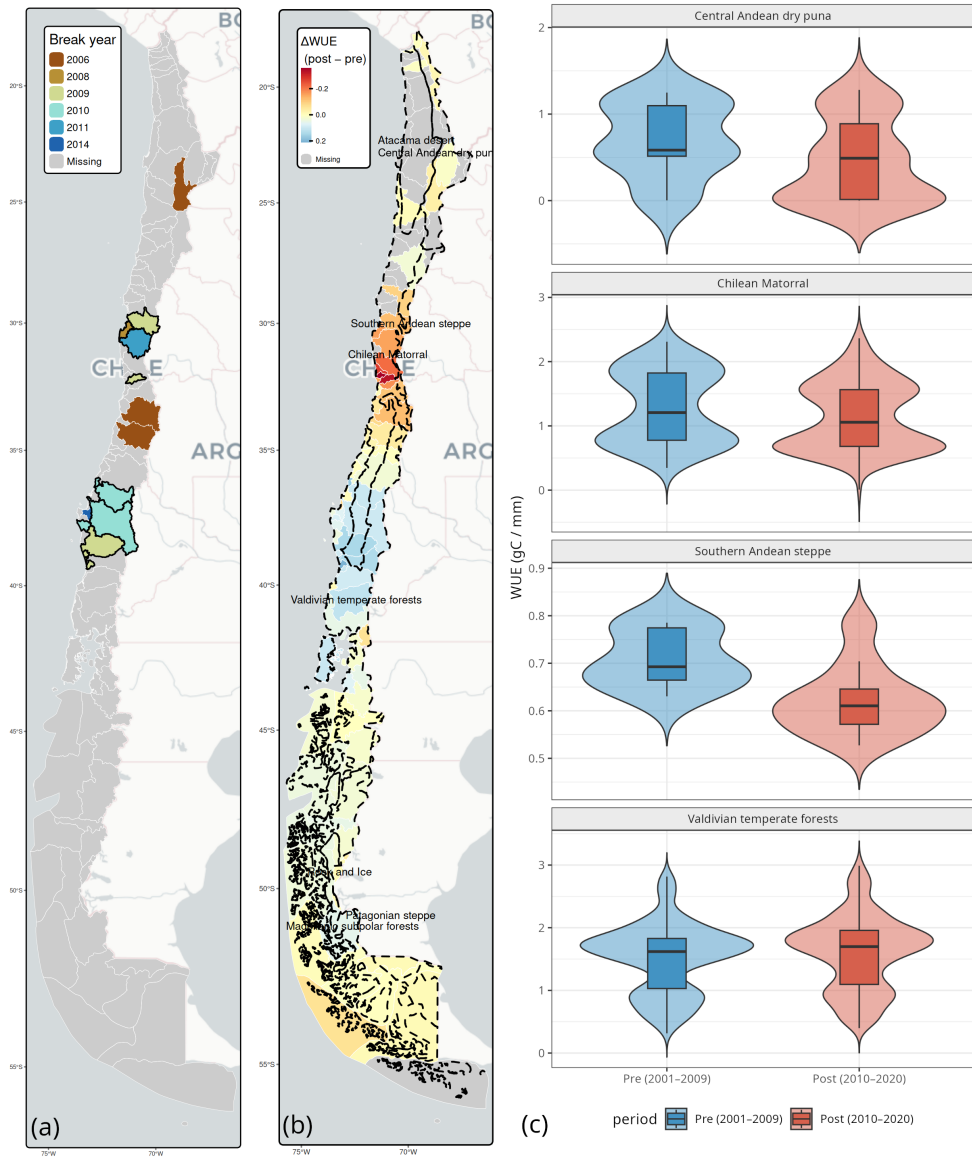


Figure 3: (a) Spatial distribution of structural break years in annual WUE per watershed, detected by the Pettitt test (significant breaks at $p < 0.05$). Watersheds with break years within ± 2 years of the megadrought onset (2010) are highlighted with bold borders, indicating megadrought-proximate structural change. Non-significant breaks are shown in grey. (b) Spatial map of WUE change between the megadrought period (2010-2020) and the pre-megadrought period (2001-2009) ($\Delta WUE = WUE_{post} - WUE_{pre}$) per watershed. Positive values (blue) indicate watersheds where mean WUE increased; negative values (red) indicate WUE decline. Ecoregion boundaries (dashed lines) are overlaid for ecological context. (c) Violin-boxplot distributions of annual WUE for the pre-megadrought (blue, 2001-2009) and megadrought (red, 2010-2020) periods, stratified by ecoregion. Each panel corresponds to one ecoregion; y-axes are free-scaled.

325 (Supplementary Table S-2) confirmed that the WUE shift was strongest in the Chilean Matorral ecoregion,
326 the primary Mediterranean-climate agricultural zone, with a Hodges-Lehmann median difference of -0.132
327 $g \cdot C \cdot kg^{-1} H_2O \cdot^{-1}$ (95% CI: -0.203 to -0.073 ; $p < 0.001$), consistent with the documented hydroclimatic
328 footprint. Violin-boxplot distributions stratified by ecoregion (Figure 3 c) confirm that the strongest negative
329 WUE shifts are concentrated in central Chile’s Mediterranean-climate watersheds, while southern transitional
330 zones show more variable or positive responses.

331 3.4. WUE responses to aridification signals

332 Spearman correlation analysis revealed positive associations between annual WUE and SPEI across most
333 Chilean watersheds (Figure 4), indicating that wetter years (higher SPEI) are generally associated with
334 higher WUE, while drought years (lower SPEI) correspond to lower WUE. The strength of this relationship
335 varied spatially and by drought index.

336 SPEI and SPI showed broadly similar spatial correlation patterns with WUE across Chilean watersheds,
337 with mean temporal R^2 of 0.181 (SPEI) and 0.172 (SPI), a difference of only 0.009 that was not statistically
338 significant in paired comparisons across per-watershed R^2 values (Wilcoxon signed-rank test: $p = 0.504$,
339 median difference = 0.002). Both SPEI and SPI significantly outperformed EDDI in temporal explanatory
340 power (mean $R^2 = 0.113$; SPEI vs. EDDI: $p < 0.001$, median difference = 0.064). This result does
341 not support a claim of SPEI superiority over SPI for this dataset; rather, it suggests that incorporating
342 atmospheric evaporative demand (through SPEI’s PET term) provides negligible additional predictive power
343 over precipitation-alone (SPI) for annual WUE variability at the 36-month accumulation scale. EDDI showed
344 correspondence to the correlations with SPI/SPEI, although there were some differences observed in the
345 central and southern regions of the country, which aligns with the more complex role of atmospheric demand
346 anomalies across various ecological contexts.

347 3.5. Panel regression: spatial versus temporal WUE-drought relationships

348 A two-way fixed-effects panel regression model (Equation 1), controlling for watershed-level time-invariant
349 differences (μ_i) and common annual shocks (τ_t), yielded non-significant coefficients for all three drought
350 indices when estimated across all 76 watersheds: SPEI ($\beta = -0.009$, SE = 0.017, $\rho = 0.589$), SPI ($\beta =$
351 -0.015 , SE = 0.016, $\rho = 0.363$), and EDDI ($\beta = -0.002$, SE = 0.014, $\rho = 0.860$). In contrast, zNPP was
352 strongly significant ($\beta = 0.124$, SE = 0.016, $\rho < 0.001$).

353 This contrast indicates that the WUE-drought index relationship is primarily driven by *between-watershed*

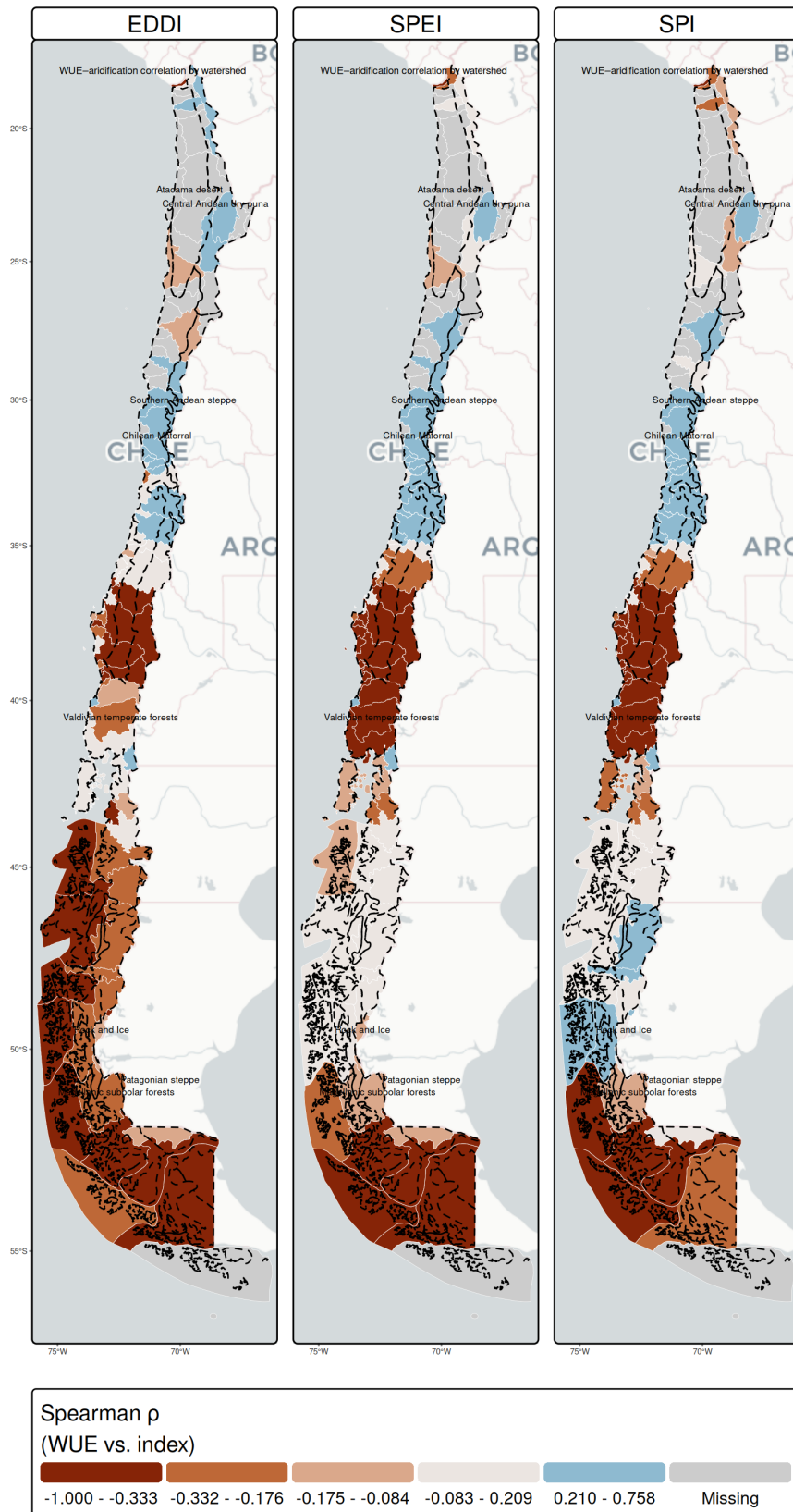


Figure 4: Spatial maps of Spearman ρ (WUE vs. drought index) per watershed, faceted by drought index (SPEI, SPI, EDDI). Diverging colour scale (blue = positive correlation, red = negative correlation).

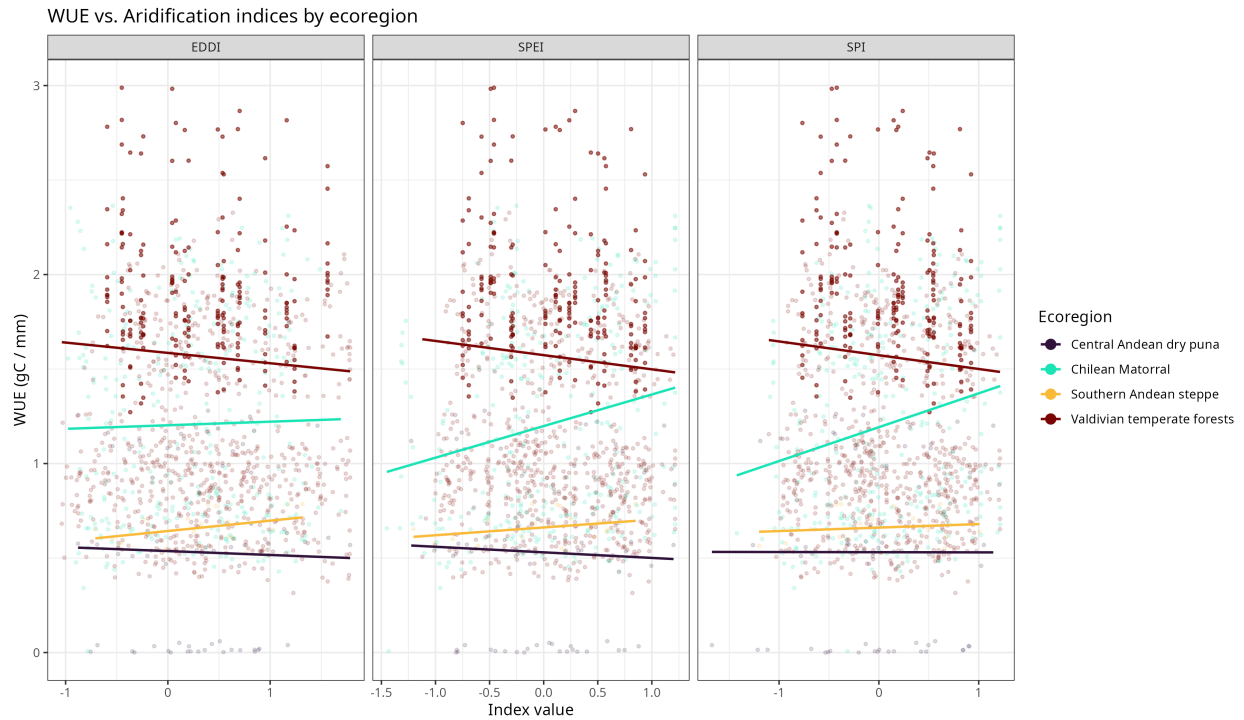


Figure 5: Scatter plots of annual WUE versus index values for SPEI, SPI, EDDI, and zNPP, stratified by ecoregion (colour). Grey line shows the overall linear fit with 95% confidence interval.

354 spatial differences, watersheds in drier areas structurally have both lower WUE and more negative drought
 355 index values, rather than by *within-watershed* year-to-year covariation of WUE and drought stress. The
 356 panel fixed effects absorb all time-invariant watershed characteristics (aridity zone, crop system, soil type,
 357 irrigation access), leaving only the within-watershed temporal variation to be explained. That zNPP retains
 358 significance in this design is consistent with the $WUE = NPP/ET$ identity: as a direct measure of annual
 359 vegetation productivity, zNPP tracks the NPP component of WUE at the within-watershed annual scale.
 360 SPEI, SPI, and EDDI, by contrast, are distal aridification drivers whose association with WUE operates
 361 primarily through the chronic spatial aridity gradient — which is absorbed by watershed fixed effects.

362 Restricting the panel model to the Mediterranean-zone watersheds (Chilean Matorral ecoregion, $n = 16$),
 363 where the megadrought’s hydroclimatic signal is strongest, yielded significant within-watershed WUE-drought
 364 associations for SPEI ($\beta = 0.124$, $\rho = 0.006$) and SPI ($\beta = 0.085$, $\rho = 0.012$), while EDDI remained
 365 non-significant ($\beta = 0.001$, $\rho = 0.979$). This finding confirms that the all-watershed non-significance reflects
 366 geographic dilution of the drought signal by drier northern and wetter southern watersheds rather than a
 367 genuine absence of within-watershed temporal coupling: once the analysis is restricted to the core megadrought
 368 zone, both precipitation-based indices show significant positive within-watershed WUE-drought associations.

369 The interaction model (Equation 2) tested whether within-watershed WUE–drought sensitivity changed
370 structurally after megadrought onset. The SPEI interaction coefficient was marginal ($\beta = 0.079$, $\rho = 0.056$),
371 while SPI ($\beta = 0.043$, $\rho = 0.269$) and EDDI ($\beta = -0.010$, $\rho = 0.742$) were non-significant. The marginal
372 SPEI result suggests a modest intensification of within-watershed sensitivity to hydroclimatic balance during
373 the megadrought period, but does not constitute strong evidence of a structural shift: the drought-period
374 WUE decline operates predominantly through the spatial aridity gradient rather than through an intensified
375 within-watershed temporal mechanism.

376 *3.6. Spatial distribution of WUE sensitivity*

377 The WUE-SPEI sensitivity map (Figure 6 a) shows the OLS regression slope ($\Delta WUE/\Delta SPEI$) per
378 watershed. Sensitivity broadly follows the latitudinal aridity gradient. Central Mediterranean watersheds
379 exhibited higher values than humid southern watersheds, with locally high sensitivity in perennial-dominated
380 irrigated zones (Figure 6 c and d).

381 *3.7. Spatial clustering of WUE trends*

382 Moran’s I statistics confirmed significant positive spatial autocorrelation in WUE trends, Pettitt break
383 years, and WUE-SPEI R^2 values across Chilean watersheds. Moran’s I for WUE trend slope was $I = 0.627$
384 ($p < 0.001$), for Pettitt break year $I = 0.164$ ($p = 0.011$), and for WUE-SPEI R^2 $I = 0.296$ ($p < 0.001$), all
385 indicating that WUE response to the megadrought is not spatially random but exhibits strong geographic
386 clustering. Adjacent watersheds tend to show similar trend directions, break timing, and drought sensitivity
387 (Figure 6 b).

388 Getis-Ord G_i^* hotspot analysis of WUE trend slopes identified coherent clusters of declining WUE (coldspots)
389 in the central Mediterranean zone and localized clusters of increasing or stable WUE (hotspots) in transitional
390 zones north and south of the primary drought impact corridor (Figure 6 b). The spatial clustering of WUE
391 responses points to large-scale hydroclimatic forcing—specifically the megadrought—as the dominant driver
392 of WUE dynamics, rather than localized management or land-use effects.

393 *3.8. Agricultural crop composition and WUE sensitivity*

394 The maps of dominant crop type (Figure 6 c) and perennial crop share (Figure 6 d), provide spatial
395 context for interpreting the WUE sensitivity patterns. Integration of census-derived crop composition with
396 WUE-SPEI sensitivity estimates ($n = 75$ watersheds with complete WUE, irrigation, and census data)
397 revealed a significant positive association between perennial crop share and drought sensitivity (Figure 7),

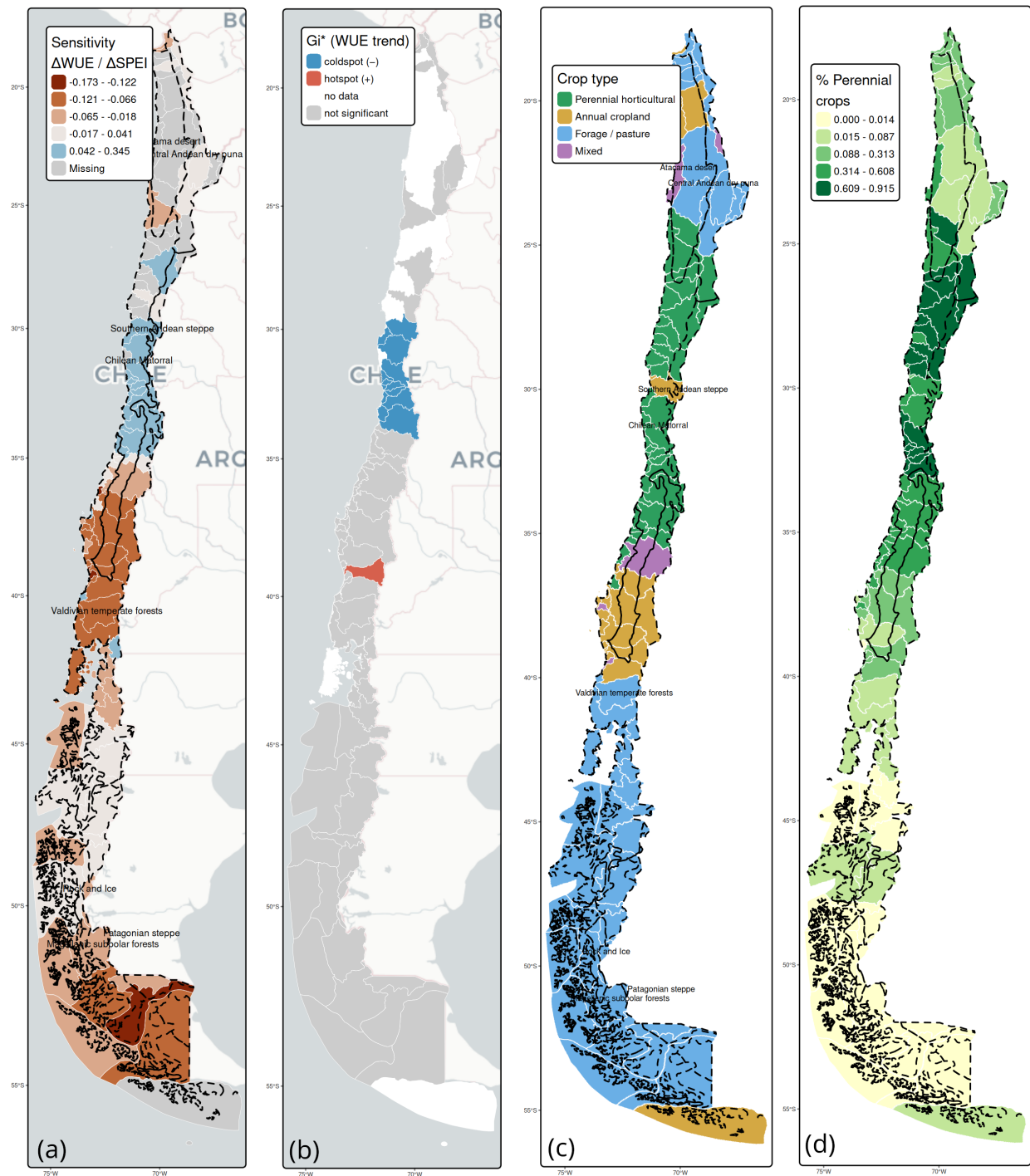


Figure 6: (a) Spatial distribution of WUE-SPEI sensitivity ($\Delta WUE / \Delta SPEI$, expressed as the OLS slope of the annual WUE ~ SPEI regression per watershed, 2001-2020). High positive values indicate strong positive WUE response to SPEI; near-zero or negative values indicate weak or inverted coupling. (b) Getis-Ord G_i^* hotspot analysis of WUE trend slope across Chilean watersheds. Red watersheds are statistically significant hotspots (positive WUE trend clusters; $G_i^* > 1.96$); blue are coldspots (negative WUE trend clusters; $G_i^* < -1.96$); grey are not significant at the 95% level. (c) Choropleth map of dominant crop type per watershed, classified by the agricultural functional group exceeding 40% of total agricultural area (Perennial horticultural, Annual cropland, Forage/pasture, Mixed). (d) Choropleth map of perennial crop share (% of total agricultural area under fruit trees and vineyards) per watershed.

398 suggesting that this is a crop-system property rather than an artifact of ecoregion geography. This implies
 399 that perennial agricultural systems in Chile are more sensitive to aridification fluctuations, both benefiting
 400 more in wet years and suffering more in dry years-compared with annual crop systems. Watersheds classified
 401 as “perennial horticultural” (40% of agricultural area under fruit trees and vineyards) exhibited the highest
 402 median WUE-SPEI sensitivity, whereas “forage/pasture” watersheds showed the lowest. “Annual cropland”
 403 watersheds showed intermediate but spatially variable sensitivity, reflecting heterogeneous responses among
 404 cereal and vegetable crops.

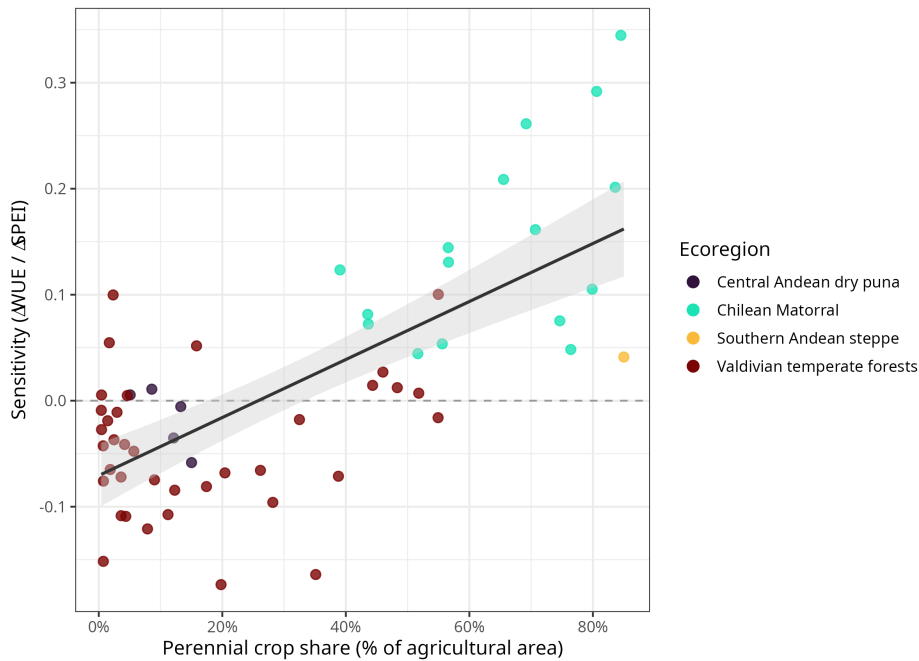


Figure 7: Scatter plot of WUE-SPEI sensitivity versus the perennial crop share (% of total agricultural area) per watershed, coloured by ecoregion, with an overall linear fit.

405 The multivariate OLS regression of WUE-SPEI sensitivity on 2021 crop proportions, controlling for irrigated
 406 area share, was fitted to the 75 watersheds with agricultural land, complete WUE data, and non-missing
 407 irrigation data ($R^2 = 0.576$, adj. $R^2 = 0.545$, $F(5,69) = 18.74$, $p < 0.001$, $n = 75$; Figure 8). Fruit tree
 408 proportion was the strongest positive predictor ($\beta = 0.213$, 95% CI [0.117, 0.310], $\rho < 0.001$), confirming
 409 that perennial horticultural systems are the most drought-sensitive land use type. Cereal proportion was
 410 a significant negative predictor ($\beta = -0.207$, 95% CI [-0.322, -0.093], $\rho < 0.001$), indicating that annual
 411 grain-crop watersheds exhibit substantially lower WUE-SPEI sensitivity than the forage/pasture reference
 412 category. Vineyard proportion was positive but not statistically significant ($\beta = 0.092$, 95% CI [-0.105, 0.289],
 413 $p = 0.354$). Vegetables were also non-significant ($\beta = -0.056$, 95% CI [-0.205, 0.093], $\rho = 0.456$). Irrigated

414 area share ($pct_irrigated$) did not reach statistical significance after controlling for crop composition ($\beta =$
 415 0.046 , 95% CI $[-0.032, 0.123]$, $p = 0.241$), suggesting that the irrigation-buffering effect is largely captured
 416 by the crop type variables. Forage/pasture served as the reference category (omitted). All variance inflation
 417 factors were below 5, confirming the absence of harmful multicollinearity.

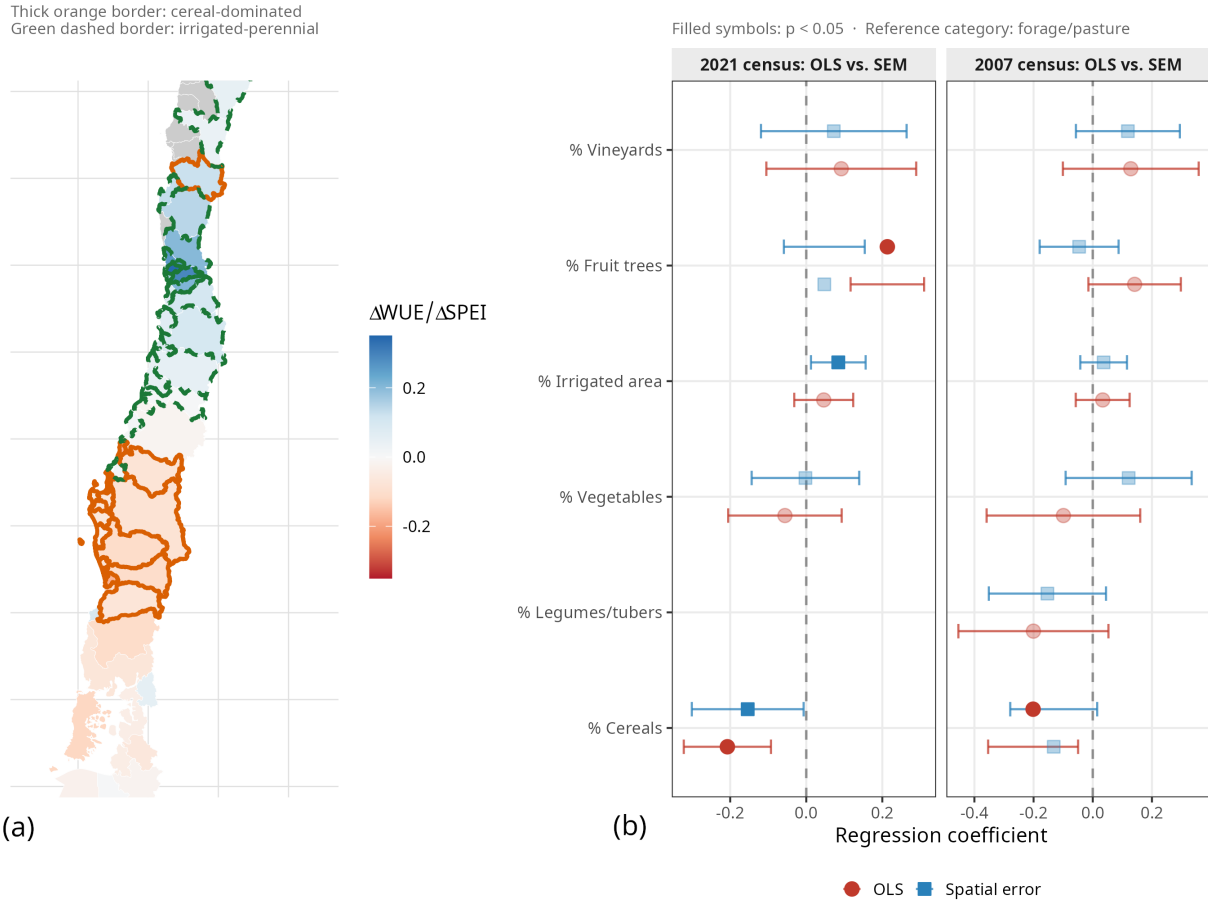


Figure 8: (a) Choropleth map of WUE-SPEI sensitivity ($\Delta WUE/\Delta SPEI$, OLS slope per watershed, 2001–2020). Thick orange borders denote cereal-dominated watersheds (Annual cropland dominant); green dashed borders denote irrigated-perennial watersheds (Perennial horticultural dominant). (b) Regression coefficients of crop-composition variables predicting WUE-SPEI sensitivity, faceted by census year. Left facet: 2021 census OLS (red circles) vs. spatial error model (blue squares); right facet: 2007 pre-drought census OLS vs. SEM. Filled symbols indicate $p < 0.05$. Reference category: forage/pasture.

418 Moran's I computed on OLS residuals was $I = 0.309$ ($p < 0.001$), indicating significant positive spatial
 419 autocorrelation in the residuals. A spatial error model (queen-contiguity weights) was therefore fitted as the
 420 primary inferential model (spatial autoregressive error parameter $\lambda = 0.615$, $\rho < 0.001$). The spatial error
 421 model partially modified the OLS conclusions: fruit tree proportion was no longer statistically significant
 422 ($\beta = 0.047$, $\rho = 0.382$), suggesting that its OLS significance was partly driven by the geographic clustering
 423 of perennial orchards in the Mediterranean zone rather than an independent crop-composition effect on

424 sensitivity. Cereal proportion remained a significant negative predictor ($\beta = -0.154$, $\rho = 0.040$), and irrigated
425 area share became a significant positive predictor ($\beta = 0.084$, $\rho = 0.022$), indicating that after accounting for
426 spatial clustering, irrigation access is associated with higher WUE-SPEI sensitivity, possibly because irrigated
427 watersheds sustain production and thus show stronger drought coupling through the NPP component of
428 WUE.

429 A pre-drought (2007) baseline regression (watershed-level, $n = 75$; $R^2 = 0.306$, adj. $R^2 = 0.234$, $F(7,67) =$
430 4.23 , $p < 0.001$; Supplementary Figure S-1) was directionally consistent: cereal proportion was a significant
431 negative predictor ($\beta = -0.202$, 95% CI $[-0.354, -0.050]$, $p = 0.010$), while fruit tree proportion was positive
432 but did not reach statistical significance at $p < 0.05$ ($\beta = 0.142$, 95% CI $[-0.015, 0.298]$, $p = 0.075$). The
433 2007 spatial error model ($\rho = 0.731$, $p < 0.001$) also showed a non-significant fruit tree coefficient ($\beta = -0.046$,
434 $p = 0.498$) and a marginally non-significant cereal coefficient ($\beta = -0.132$, $p = 0.078$). The directional
435 consistency of the cereal effect across census years supports a structural rather than adaptive interpretation
436 of the crop composition–sensitivity relationship (see Section 4.3).

437 To assess whether the WUE-SPEI sensitivity associations are confounded by natural vegetation (matorral)
438 contamination within the agricultural mask, natural vegetation fraction per watershed was computed from
439 ESA CCI-LC (2016; classes 40, 120–122, 130) and added as a covariate to the spatial error model. Median
440 natural vegetation fraction within agricultural pixels was 0.267 (range 0.00–0.60), reflecting the substantial
441 fragmentation of the agricultural landscape in central Chile. When included in the spatial error model, the
442 natural vegetation fraction was non-significant ($\beta = -0.017$, $p = 0.810$), and the primary crop composition
443 coefficients were directionally consistent with the base model, indicating that matorral contamination does
444 not substantially confound the WUE sensitivity associations.

445 To examine whether the positive WUE-SPEI sensitivity of perennial-dominated watersheds reflects the NPP
446 numerator, the ET denominator, or both components of WUE, per-watershed OLS sensitivity slopes were
447 computed separately for zNPP~SPEI and for SETI~SPEI (the MODIS evapotranspiration anomaly index)
448 and then regressed on the same crop composition variables. For the NPP component (zNPP-SPEI; OLS
449 $R^2 = 0.519$, $n = 99$), vineyard proportion was the strongest positive predictor (OLS: $\beta = 1.841$, $p < 0.001$;
450 SEM: $\beta = 0.769$, $p = 0.036$), and cereal proportion was a significant negative predictor in OLS ($\beta = -0.614$,
451 $p = 0.016$) but not in the spatial error model ($\beta = 0.017$, $p = 0.953$). For the ET component (SETI-SPEI;
452 OLS $R^2 = 0.348$, $n = 100$), vineyard proportion was also positive (OLS: $\beta = 1.061$, $p = 0.005$) and cereal
453 proportion was negative (OLS: $\beta = -0.837$, $p = 0.001$), but after spatial error correction, all coefficients

454 were non-significant ($SEM = 0.668$). These results indicate that the positive WUE-SPEI sensitivity of
 455 vineyard-dominated watersheds is primarily driven by greater NPP suppression under drought rather than
 456 ET maintenance, while the crop composition associations for the ET component do not survive spatial
 457 autocorrelation correction.

458 3.9. Cross-sectional analysis and model comparison

459 Cross-sectional regressions using the census year 2020 (where WUE, drought index, and crop composition
 460 are contemporaneous) confirmed the temporal patterns. Adding crop composition proportions to the cross-
 461 sectional models increased R^2 substantially for all indices: from 0.0004 to 0.432 for SPEI, from 0.078 to 0.459
 462 for SPI, and from 0.234 to 0.431 for EDDI (Figure 9). Mean temporal R^2 (averaged across the full 2001-2020
 463 per-watershed regressions) was 0.181 for SPEI, 0.172 for SPI, and 0.113 for EDDI. The substantially larger R^2
 464 gains from adding crop composition in the 2020 cross-section compared with temporal models suggest that
 465 crop structure explains a greater share of the spatial distribution of drought sensitivity than of its interannual
 466 variability within each watershed. Sample sizes for each analytical step are documented in Supplementary
 467 Table S-1 ($n = 75$ watersheds with complete WUE, irrigation, and census data used in crop composition
 468 analyses).

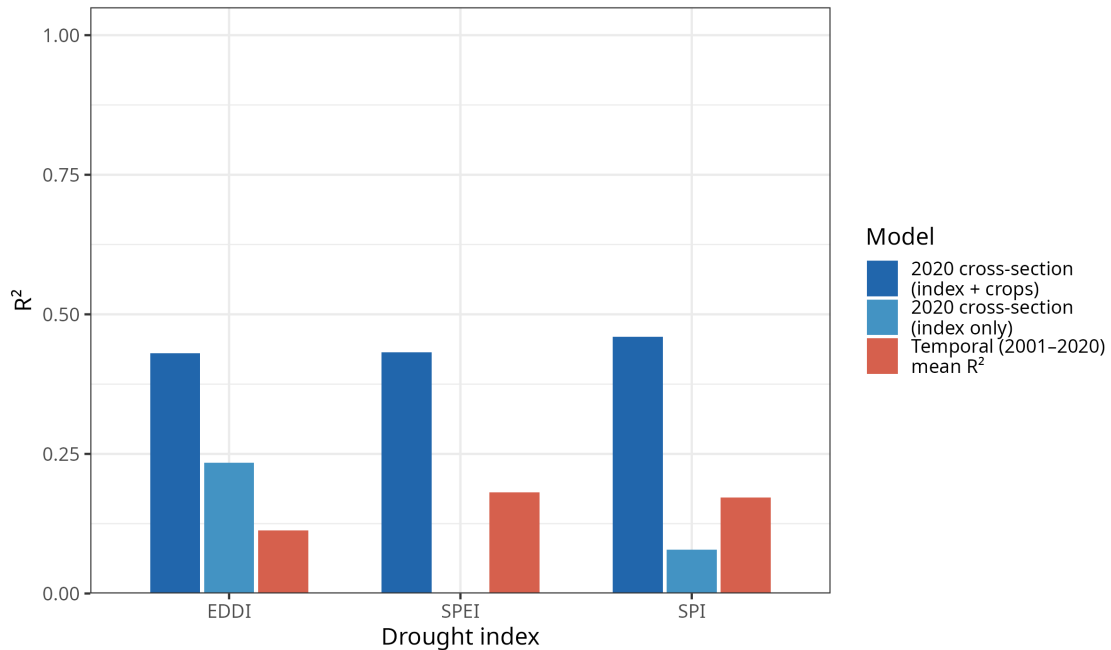


Figure 9: Bar chart comparing R^2 values across three model types for each drought index: (i) 2020 cross-sectional regression (index only: SPEI = 0.0004, SPI = 0.078, EDDI = 0.234), (ii) 2020 cross-sectional regression (index + crop proportions: SPEI = 0.432, SPI = 0.459, EDDI = 0.431), and (iii) temporal regression (mean R^2 : SPEI = 0.181, SPI = 0.172, EDDI = 0.113).

469 4. Discussion

470 4.1. The megadrought as a structural driver of WUE change

471 The Pettitt test results indicate that 17 of 76 watersheds (22.4%) showed raw-significant structural breaks
472 ($p < 0.05$) in WUE dynamics, with 71% of those breaks concentrated around 2010. While none survived
473 Benjamini-Hochberg FDR correction, reflecting the limited statistical power of individual tests applied
474 to spatially correlated watersheds with only 20 annual observations, the near-2010 concentration of raw-
475 significant breaks is geographically coherent and consistent with the onset of the megadrought. This finding
476 is best interpreted as evidence that WUE dynamics are *temporally associated with* the megadrought onset
477 rather than as definitive statistical proof of a structural break: the spatial concentration of change points
478 in the central Mediterranean zone, precisely where the megadrought’s hydroclimatic footprint is strongest
479 (Boisier et al., 2018; Garreaud et al., 2020), provides geographic corroboration. The geographic coherence
480 of near-2010 breaks in this zone supports a megadrought-associated interpretation, though the spatial
481 autocorrelation among watersheds (Moran’s I for break year = 0.164, $p = 0.011$) means the 17 Pettitt results
482 cannot be treated as independent observations, which precludes formal inference about temporal clustering
483 from independence-based tests.

484 This finding is consistent with the broader literature on regime shifts in ecological and agricultural systems
485 under sustained stress (Folke et al., 2004), and reinforces the interpretation of the megadrought as a
486 transformative stress event rather than an incremental trend (Kates et al., 2012). The implications for water
487 management are significant: if WUE has undergone a step-change downward in central Chile, agricultural
488 water requirements per unit of production have increased, placing additional pressure on already-stressed
489 water allocation systems (Budds, 2020; Rivera et al., 2016).

490 4.2. Multidimensional aridification: why index choice matters

491 SPEI and SPI showed statistically indistinguishable temporal explanatory power ($p = 0.504$), indicating
492 that incorporating atmospheric evaporative demand (SPEI’s PET term) provides negligible predictive gain
493 over precipitation-alone (SPI) at the 36-month accumulation scale, likely because precipitation deficits and
494 evaporative demand anomalies are strongly correlated over multi-year megadrought periods (Li et al., 2023;
495 Zambrano et al., 2025). EDDI was significantly weaker than both, suggesting that atmospheric demand
496 anomalies in isolation are less directly coupled to WUE at the watershed scale.

497 EDDI’s distinct spatial correlation pattern indicates partially independent effects of atmospheric demand on

498 WUE. Semi-partial Spearman correlations (controlling for SPEI) confirmed a modest independent EDDI signal
499 (mean partial $\rho = 0.041$ vs. unconditional $\rho = 0.069$), consistent with strong thermodynamic coupling between
500 precipitation deficit and VPD (Li et al., 2023). The EDDI effect is therefore a statistical association rather
501 than a causally isolated demand pathway; in irrigation-buffered watersheds, near-zero EDDI correlations
502 reflect sustained production when water supply is maintained.

503 This result has practical implications for drought monitoring and agricultural water management: because
504 SPEI and SPI show statistically equivalent explanatory power at the 36-month scale, precipitation-only
505 monitoring (SPI) may be sufficient for characterizing annual WUE variability under conditions similar to
506 the megadrought. However, in shorter-accumulation or sub-seasonal contexts where evaporative demand
507 anomalies decouple from precipitation deficits, relying solely on SPI could underestimate WUE impacts in
508 regions experiencing intensified atmospheric demand under anthropogenic warming (Boisier et al., 2018).

509 4.3. Spatial versus temporal variation in WUE-drought relationships

510 The two-way fixed-effects panel regression revealed an important analytical distinction: the cross-sectional
511 associations between drought indices and WUE (mean temporal $R^2 = 0.172$ - 0.181 for SPI and SPEI) reflect
512 the chronic spatial gradient in aridity across Chile more than they reflect year-to-year within-watershed
513 dynamics. Once watershed fixed effects absorb time-invariant characteristics, aridity zone, crop system, soil
514 type, irrigation access; the within-watershed temporal association between drought indices and WUE is
515 non-significant for SPEI, SPI, and EDDI across all 76 watersheds. Only zNPP, which directly tracks the
516 NPP numerator of WUE, retains significance in the within-watershed temporal dimension.

517 This spatial-versus-temporal distinction has direct implications for how the results should be interpreted.
518 The primary contribution of this paper is in characterizing the *spatial* structure of WUE drought sensitivity:
519 where the megadrought has concentrated its impacts (central Mediterranean zone), which crop systems
520 amplify that impact (perennial horticultural), and what the spatial extent and clustering of vulnerability
521 looks like (Moran's $I = 0.627$). The panel regression does not invalidate these spatial findings; it clarifies that
522 the aridification signal is spatially organized rather than temporally reactive at the annual within-watershed
523 scale.

524 Restricting the panel to Chilean Matorral watersheds ($n = 16$) yielded significant SPEI ($\beta = 0.124$, $\rho =$
525 0.006) and SPI ($\beta = 0.085$, $\rho = 0.012$) associations while EDDI remained non-significant ($\beta = 0.001$, $\rho =$
526 0.979), confirming geographic dilution as the source of all-watershed non-significance. A post-2010 interaction

527 term showed only a marginal SPEI effect ($\beta = 0.079$, $\rho = 0.056$), indicating that the megadrought WUE
528 decline operates predominantly through the chronic spatial aridity gradient rather than an abrupt shift in
529 within-watershed temporal sensitivity.

530 The sign reversal between the all-watershed SPEI coefficient ($\beta = -0.009$, NS) and the Mediterranean-
531 subset coefficient ($\beta = +0.124$, $\rho = 0.006$) reflects confounding by southern watersheds, where cloudier
532 wet years suppress NPP relative to ET, the opposite of the moisture-limited Mediterranean response. The
533 Mediterranean-subset result is the ecologically meaningful within-watershed WUE-SPEI sensitivity; the
534 all-watershed null reflects geographic heterogeneity rather than absent drought-WUE coupling.

535 4.4. Crop composition as a moderator of drought sensitivity

536 The finding that perennial horticultural watersheds exhibit higher WUE-SPEI sensitivity than annual
537 or forage systems is interpreted as a spatial association between crop composition and drought sensitivity,
538 not as evidence of direct causation. Higher sensitivity means that WUE fluctuates more strongly with the
539 hydroclimatic balance signal, it increases more in wet years and decreases more in dry years. Under prolonged
540 drought such as the megadrought, this implies that perennial-dominated zones face disproportionately large
541 WUE penalties. Given that Chile's export-oriented fruit and wine sectors are concentrated in precisely
542 these Mediterranean-climate watersheds, the economic exposure to aridification is amplified by the crop
543 composition of these zones (Fernández et al., 2023).

544 This spatial association is unlikely to be solely an artefact of drought-driven crop composition change.
545 A potential concern is that the 2020-2021 census composition may partly reflect adaptive responses to
546 the preceding megadrought, for example, abandonment of water-intensive annual crops and retention of
547 higher-value perennials. However, the pre-drought (2007) watershed-level baseline regression ($n = 75$; R^2
548 $= 0.306$; Supplementary Figure S-1) was directionally consistent with the 2021 results. Cereal proportion
549 was a significant negative predictor in both the 2007 OLS ($\beta = -0.202$, $\rho = 0.010$) and the 2021 SEM (β
550 $= -0.154$, $\rho = 0.040$). Fruit tree proportion was positive in both years (2021 OLS: $\beta = 0.213$, $\rho < 0.001$;
551 2007 OLS: $\beta = 0.142$, $\rho = 0.075$) but did not reach significance in the spatial error models for either year.
552 The directional consistency of the cereal effect, the most robust predictor across all model specifications,
553 supports the interpretation that the crop composition-sensitivity relationship reflects a structural property of
554 agricultural land use rather than a purely adaptive reorganization in response to drought. The evidence for
555 the perennial (fruit tree) effect is weaker under spatial error correction and should be interpreted cautiously.

556 This association is consistent with physiology: perennials sustain large permanent canopy structures
557 that amplify WUE fluctuations, while annuals buffer GPP loss through shorter growing cycles and earlier
558 senescence; forage systems show lower base WUE sensitivity (Chakraborti et al., 2023; Davis et al., 2017).
559 The NPP/ET decomposition analysis confirms that this perennial sensitivity operates primarily through the
560 NPP numerator: vineyard proportion was a significant predictor of zNPP-SPEI sensitivity in the spatial
561 error model ($\beta = 0.769$, $p = 0.036$) but the ET denominator response (SETI-SPEI SEM: $\beta = 0.582$, $p =$
562 0.137) did not survive spatial correction, suggesting that droughts suppress vine productivity more strongly
563 than they reduce vineyard evapotranspiration, consistent with deep-rooted perennials maintaining partial
564 water uptake under moisture stress.

565 These results complement the global literature on crop-water relationships (Boser et al., 2024; Wang et al.,
566 2024) by demonstrating that the composition of agricultural land use fundamentally shapes the WUE
567 response to aridification at the regional scale. Policies aimed at improving agricultural water use efficiency in
568 Chile during prolonged drought should account for this crop-composition modulation effect, recognizing that
569 perennial-dominated watersheds require more adaptive water governance than annual crop systems (FAO y
570 ONU Agua, 2025; Hellegers and Van Halsema, 2021).

571 4.5. Spatial structure and governance implications

572 The strong spatial autocorrelation in WUE trends—confirmed by Moran’s I and the G_i^* hotspot analysis—
573 indicates that the megadrought’s impact on agricultural WUE is not atomistic at the watershed level but
574 organized into large coherent spatial patterns. This spatial structure has direct implications for water
575 governance: the most drought-affected zones form geographic clusters in which water stress is likely to
576 compound across watersheds simultaneously, reducing the potential for spatial redistribution of water
577 resources to buffer impacts (Budds, 2020; Rivera et al., 2016).

578 The spatial clustering of WUE coldspots in the central Mediterranean zone, where Chile’s most eco-
579 nomically important irrigated agriculture is concentrated, represents a critical vulnerability nexus. Under
580 continued aridification, the convergence of high crop sensitivity (perennial systems), strong hydroclimatic
581 stress (megadrought), and spatial clustering of WUE decline could accelerate the economic and ecological
582 consequences of insufficient adaptive response (Fernández et al., 2023; MMA, MINAGRI, 2013).

583 *4.6. Methodological limitations and future research*

584 Several limitations should be acknowledged. First, WUE estimated from satellite-derived *NPP/ET*
585 integrates all vegetation cover within agricultural areas and may include natural vegetation patches within
586 watershed boundaries, introducing noise. The agricultural mask used to define WUE pixels was held static
587 over the 2001-2020 study period. In central Chile, agricultural lands are fragmented and interspersed with
588 native Mediterranean shrubland (matorral), which may exhibit different NPP and ET responses to drought.
589 Because matorral distribution varies systematically along the latitudinal aridity gradient, the proportion of
590 natural vegetation within the agricultural mask could potentially confound the observed spatial differences
591 in WUE sensitivity. We addressed this directly by computing the natural vegetation fraction per watershed
592 from ESA CCI-LC (2016) and including it as a control covariate in the spatial error model. Median natural
593 vegetation fraction within agricultural pixels was 0.267 (range 0.00–0.60). The natural vegetation covariate
594 was non-significant ($\beta = -0.017$, $p = 0.810$) and did not alter the primary crop composition coefficients,
595 indicating that matorral contamination is not a material confounder of the WUE sensitivity patterns reported
596 here.

597 Second, the static agricultural mask cannot distinguish active cultivation from drought-driven land aban-
598 donment. If irrigated parcels transitioned to bare soil or sparse cover during the megadrought, the resulting
599 decline in both NPP and ET would produce WUE signals indistinguishable from active-area efficiency loss.
600 This confound is most likely to affect rainfed cereal watersheds, where water stress can cause crop failure
601 and partial abandonment. Future studies should cross-reference the WUE time series with annual dynamic
602 crop extent products (e.g., GLAD Global Cropland Extent) to identify pixels that transitioned out of active
603 agriculture during the megadrought period.

604 Third, the use of two cross-sectional agricultural censuses (2007 and 2020-2021) captures pre- and post-
605 megadrought composition but cannot resolve gradual compositional change within the study period; longitu-
606 dinal census data would allow analysis of how cropping system adaptation responds incrementally to drought
607 stress. The directional consistency between 2007 and 2021 regression results reduces but does not eliminate
608 this concern.

609 Fourth, central Chile’s agricultural valleys depend heavily on Andean snowmelt-fed rivers for irrigation water.
610 SPEI and SPI are calculated from local precipitation and reference evapotranspiration, and do not capture
611 the contribution of upstream snowpack to actual irrigation water availability. In years with strong snowmelt
612 input, irrigated crops may be buffered from meteorological drought signals that SPEI would classify as severe.

613 This decoupling between local meteorological drought and actual irrigation water availability is particularly
614 relevant for perennial horticultural crops, which are predominantly irrigated by snowmelt-fed river systems,
615 and may cause the WUE-SPEI sensitivity of these systems to be underestimated or mischaracterized. Future
616 research should explicitly integrate hydrological drought indices such as Snow Water Equivalent (SWE)
617 anomalies or streamflow-based indices such as the Streamflow Drought Index (Lema et al., 2025) to capture
618 the buffering effect of upstream snowpack. This would allow separation of meteorological from hydrological
619 drought effects on WUE in Andean-fed agricultural systems and provide a more logically coherent framework
620 for evaluating crop-specific vulnerability in these irrigation-dependent regions.

621 Future research should examine WUE dynamics at sub-seasonal scales to capture phenological shifts,
622 integrate irrigation infrastructure data (excluded from this paper per scope definition) to more precisely
623 distinguish irrigated from rainfed WUE responses, and extend the analysis into the 2021-2025 period to
624 assess whether the megadrought’s hydroclimatic signal has intensified or abated. The census-based crop
625 composition analysis presented here provides a methodological template that can be applied to evaluate
626 whether agricultural adaptation—through crop switching or intensification—is modifying WUE trajectories
627 over time (Chakraborti et al., 2023; Xie et al., 2023).

628 5. Conclusions

629 This study provides a multi-method characterization of agricultural WUE responses to the Central Chile
630 Megadrought (2010-2020) across the country’s hydrological watershed network. The principal conclusions
631 are:

- 632 1. **WUE trends are spatially heterogeneous and predominantly negative in central Chile.**
633 Mann-Kendall trend analysis confirmed declining WUE across the Mediterranean-climate zone ($n = 76$
634 watersheds), consistent with the megadrought’s documented hydroclimatic footprint.
- 635 2. **WUE change points are temporally associated with megadrought onset.** Pettitt tests
636 detected raw-significant change points in 17 of 76 watersheds, with 71% of those breaks occurring
637 within ± 2 years of 2010. None survived FDR correction, reflecting the limited power of individual
638 non-parametric tests on spatially correlated 20-year series. The spatial concentration of near-2010
639 breaks in the central Mediterranean zone provides geographic corroboration of megadrought-associated
640 timing, though spatial autocorrelation in break years (Moran’s $I = 0.164$, $p = 0.011$) prevents formal
641 inference about temporal clustering from independence-based tests.

- 642 **3. SPEI and SPI show comparable temporal explanatory power for WUE variability; both**
643 **outperform EDDI.** Mean temporal R^2 was 0.181 (SPEI) and 0.172 (SPI), a difference not statistically
644 significant across watersheds. EDDI was significantly weaker (mean $R^2 = 0.113$, $\rho < 0.001$ vs. both).
645 This indicates that at the 36-month accumulation scale, the precipitation signal captures most of the
646 WUE variance, with the atmospheric demand component adding small but detectable independent
647 explanatory power (mean partial ρ with SPEI controlled = 0.041 vs. unconditional $\rho = 0.069$).
- 648 **4. The WUE-drought relationship is primarily spatial, not temporal, but the core**
649 **megadrought zone exhibits within-watershed coupling.** Two-way fixed-effects panel re-
650 gressions show non-significant within-watershed associations across all 76 watersheds, indicating
651 that cross-sectional correlations reflect the chronic aridity gradient rather than year-to-year within-
652 watershed dynamics. Only zNPP is significant in the temporal panel dimension. When restricted
653 to Mediterranean-zone watersheds (Chilean Matorral, $n = 16$), SPEI and SPI become significant,
654 confirming geographic dilution as the source of all-watershed non-significance. The primary analytical
655 contribution is characterization of the *spatial* structure of WUE sensitivity, concentrated in the central
656 Mediterranean zone where within-watershed temporal coupling is also detectable.
- 657 **5. Crop composition is spatially associated with WUE sensitivity, with cereal proportion**
658 **as the most robust predictor.** Watersheds dominated by perennial horticultural crops exhibited
659 higher WUE-SPEI sensitivity than annual or forage systems in the OLS model. A spatial error model
660 attenuated the fruit-tree coefficient to non-significance, while cereal proportion remained a significant
661 negative predictor ($\beta = -0.154$, $p = 0.040$) and irrigated area became a significant positive predictor (β
662 $= +0.084$, $p = 0.022$). The cereal effect — lower WUE-SPEI sensitivity relative to forage/pasture — is
663 the most robust finding under the spatial error specification. Cereal proportion was also a significant
664 negative predictor in the pre-drought 2007 watershed-level regression ($\beta = -0.202$, $p = 0.010$), supporting
665 a structural rather than purely adaptive interpretation of this effect. NPP/ET component decomposition
666 shows that vineyard sensitivity operates primarily through the NPP numerator (zNPP-SPEI SEM: $\beta =$
667 0.769 , $p = 0.036$), while the ET denominator response does not survive spatial correction, indicating
668 that droughts suppress vine productivity more than vineyard evapotranspiration. Natural vegetation
669 contamination within the agricultural mask (median fraction = 0.267) was tested as a control covariate
670 and was non-significant ($\beta = -0.017$, $p = 0.810$), confirming that matorral fragmentation does not drive
671 the crop composition–sensitivity associations.
- 672 **6. WUE responses are spatially clustered.** Moran’s I and G_i^* statistics confirm that WUE trends

673 and drought sensitivity form coherent geographic clusters, with the highest drought impact concentrated
674 in the central Mediterranean zone.

675 These findings collectively demonstrate that the Central Chile Megadrought has not only reduced mean
676 WUE but has reorganized the statistical structure of agricultural water use efficiency in ways that reflect
677 persistent aridification rather than episodic drought. Managing agricultural water use under continued
678 aridification will require governance frameworks that account for the multidimensional nature of drought,
679 the spatial association between crop system composition and drought sensitivity, and the spatially clustered
680 geography of vulnerability.

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688 **7. Code and Data Availability**

689 All analysis code is provided as Supplementary Material for reviewer access (R scripts for all preprocessing,
690 statistical analysis, and figure generation steps, with package versions managed via `renv` for reproducibility).
691 A permanent citable archive with a DOI will be deposited at Zenodo upon acceptance, and the full repository
692 will be made publicly available at that time. https://github.com/frzambra/WUE_article

693 Annual cropland WUE rasters (NPP/ET, 1 km, 2001-2020) were obtained from Jiang et al. (2025), available
694 at the CASEarth Data Platform (<https://data.casearth.cn/dataset/640f0132819aec3f2b52a4bb>). NPP data
695 (zNPP) were obtained from MOD17A3HGF (MODIS Terra, 500 m, Collection 6.1) and ET data (SETI)
696 from MOD16A2GF (MODIS Terra, 500 m 8-day, Collection 6.1), both available from NASA Earthdata
697 (<https://earthdata.nasa.gov>). Drought index raster time series (SPEI, SPI, EDDI) were computed from ERA5-
698 Land monthly meteorological reanalyses. Agricultural census microdata for the 2007 and 2020-2021 censuses
699 are publicly available from the Chilean Instituto Nacional de Estadísticas (INE; <https://www.ine.gob.cl>).
700 Full software version details are listed in §2.6.

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857 **Supplementary Material**

858 *Supplementary Table S-1 — Sample sizes by analytical step*

Analysis	n	Spatial unit	Notes
WUE trend (Mann-Kendall)	76	Watershed	Agricultural watersheds with complete 2001-2020 WUE data

Analysis	n	Spatial unit	Notes
WUE-SPEI sensitivity (OLS slope)	76	Watershed	Same set; one per-watershed regression per index
Panel regression (two-way FE)	76	Watershed	Balanced panel; 76×20 = 1,520 watershed-year obs. minus missing
Panel regression — Mediterranean subset	16	Watershed	Chilean Matorral ecoregion watersheds; panel $N = 16 \times 20 =$ 320 watershed-year obs.
Crop composition OLS (2021 census)	75	Watershed	76 minus 1 watershed with missing irrigation data
Crop composition spatial error model (2021)	75	Watershed	Same set; <code>spatialreg::errorsarlm,</code> queen contiguity
Crop composition OLS (2007 baseline)	75	Watershed	2007 VII Census proportions aggregated to watershed level via <code>comcuen.rds</code> spatial join; $R^2 = 0.306$
Crop composition SEM (2007 baseline)	75	Watershed	<code>spatialreg::errorsarlm,</code> queen contiguity; = 0.731, $p < 0.001$

⁸⁵⁹ *Supplementary Table S-2 — Ecoregion-stratified Wilcoxon signed-rank tests*

⁸⁶⁰ Wilcoxon signed-rank tests comparing mean WUE in the pre-megadrought (2001-2009) versus
⁸⁶¹ megadrought (2010-2020) periods, stratified by ecoregion. Hodges-Lehmann median difference and
⁸⁶² 95% confidence interval are reported alongside p-values. Full numerical results are available in

863 outputs/tables/wilcox_by_ecoregion.csv.

864 *Supplementary Figure S-1 — Pre-drought (2007) crop composition and WUE sensitivity*

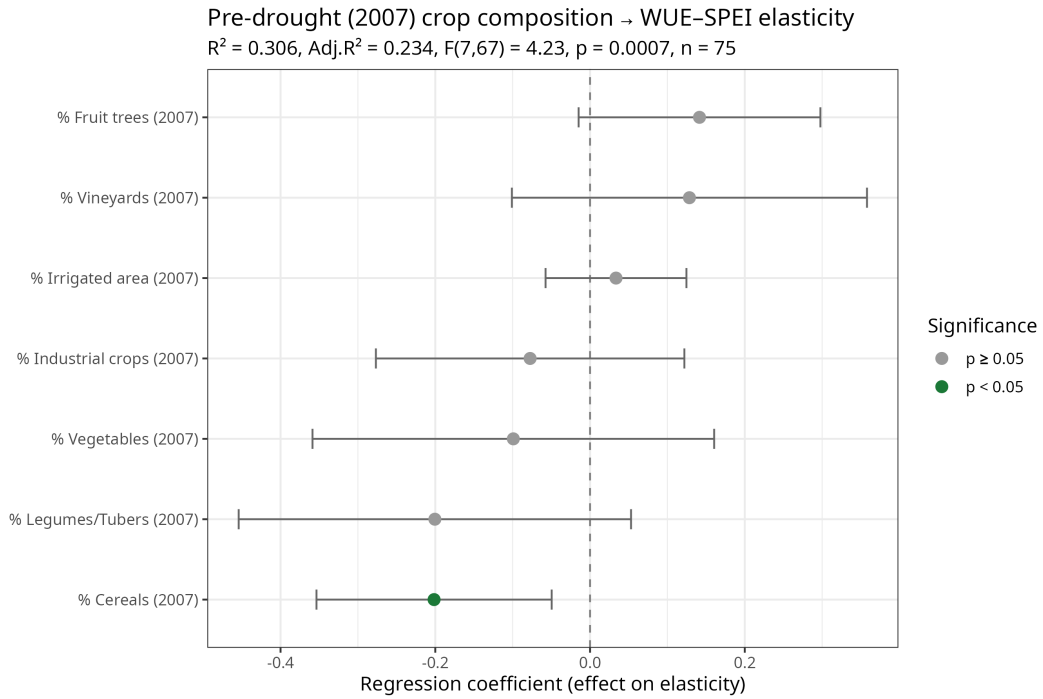


Figure 10: Pre-drought (2007) crop composition as a predictor of WUE-SPEI sensitivity. OLS and spatial error model (SEM) regression coefficient plots using VII Agricultural and Forestry Census (2007) proportions aggregated to watershed level as predictors (reference category: forage/pasture; $n = 75$ watersheds). The cereal coefficient is directionally consistent with the 2021 SEM result, supporting a structural rather than purely adaptive interpretation. OLS: $R^2 = 0.306$, $F(7,67) = 4.23$, $p < 0.001$. SEM: $\beta = 0.731$, $p < 0.001$.