

Abstract

Survey200: CAP LTER's approach to extensive field monitoring

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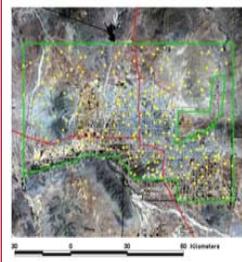
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“To what extent do human variables contribute to explaining spatial variation in the basic ecological properties of an urban ecosystem, and how do these relationships change through time?”

Study Design

Field survey

Field sampling scheme: We used a dual-density, randomized, tessellation-stratified design (see point distribution at right) to obtain a spatially dispersed, representative and unbiased sample of the study area. Sampling density inside the developed urban core was 3x that outside. Our application of probability-based sampling at this scale across an urban area is unique.

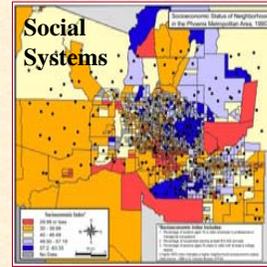


Typical survey sites

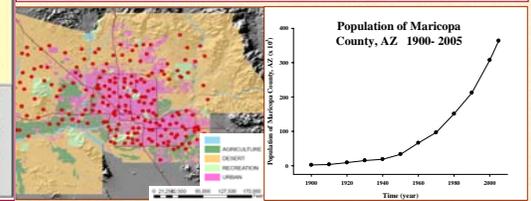
A synoptic integrated field inventory of key biotic, abiotic, and some human variables is carried out in 30x30m field plots (n≈200) over a 3-month period. Beginning in 2000, surveys are conducted every 5 years.



Additional data



Geographic and socioeconomic variables are obtained from existing CAP LTER databases and from other sources, such as the US Census block group immediately surrounding each of the survey sites.



Soil chemistry: Heavy metals

Spatial Analysis of Pb concentration (ppm) by Kriging



Spatial Analysis of V concentration (ppm) by Kriging



Soil samples (top 10 cm) were analyzed by inductively coupled plasma mass spectrometry for concentrations of trace elements, and distributions were plotted with GIS. Elements such as Pb, Cd, Cu, Ag show strong urban influence while V, As, Zn, Mn might have complex sources from geology and human activity.

Spatial Analysis of As concentration (ppm) by Kriging



Soil chemistry: nutrients

• **Surface soil (0 – 10 cm depth) was sampled to determine whether a legacy of past agricultural use is detectable in the soil chemistry of contemporary residential lawns**

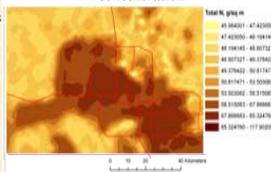
► Organic matter, carbon, nitrogen, and soluble ion concentrations were twice as great in yards that were previously farmed as in yards that were developed from desert, and the pools remained elevated 40 y after urbanization.

► Nitrogen accumulation rates (1.5 g m⁻² y⁻¹) in residential soils were not affected by prior land use, suggesting that home owners do not adjust rates of residential fertilizer application to account for the high fertility of previously farmed soils.

► Bioavailable, inorganic phosphorus (P) was elevated in soil with a recent agrarian past, but this signal disappeared after 10-30 y of residential use because P accumulated in yards developed on native desert soils.

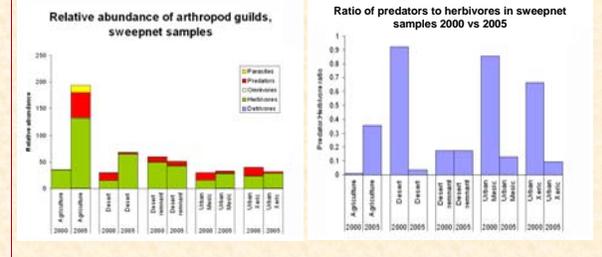
► Our results suggest a “direct agrarian legacy,” wherein agricultural impacts on soil chemistry endure urbanization, more so than an “indirect legacy,” wherein contemporary land management is shaped by prior land use.

Hierarchical spatial modeling was employed to map surface-soil nutrient concentration



Arthropods: 2000 to 2005

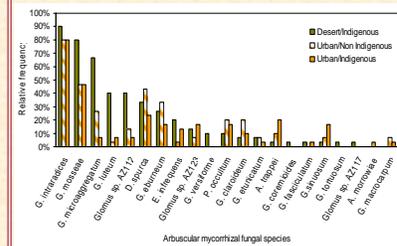
Although this is a long-term study and we can only compare two sampled years so far, we find three interesting results. First, the urban sites, particularly the desert remnants, appear much more stable from 2000 to 2005 compared to the desert sites, which fluctuated widely. Differences in productivity and rainfall provide a reasonable explanation for the fluctuations: 2000 was a drought year whereas 2005 which received much rainfall during the winter. The result lend further support to the hypothesis that urban sites are buffered from seasonal or yearly fluctuations in productivity owing to human management.



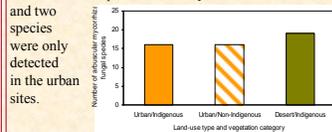
Arbuscular mycorrhizal fungi

A selected subset of Survey 200 sites were sampled for AM fungi. Soil samples were taken at from the roots systems of plants at 10 sites of each of the following types: urban, non-indigenous plants; urban, indigenous plants; and desert, indigenous plants. Trap cultures initiated from these soil samples were used to obtain AM fungal spores for identification.

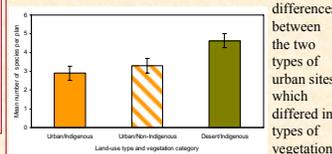
Relative Frequency: There were differences between desert and urban sites in the relative frequency of some AM fungal species. Desert sites had more species with a relative frequency > 50%. Three relatively common species at desert sites (*Glomus microaggregatum*, *G. luteum* and *Glomus* sp. AZ112) were detected less frequently at urban sites.



Species Composition: 21 AM fungal species were detected (urban=17, desert=19). Roughly 70% of the species were detected at both urban and desert locations. Four species were only detected in the desert and two species were only detected in the urban sites.



species/sample: A greater number of AM fungal species per plant detected at desert vs. urban sites (F=5.874, df=2, P=0.004). There were no significant differences between the two types of urban sites which differed in types of vegetation.



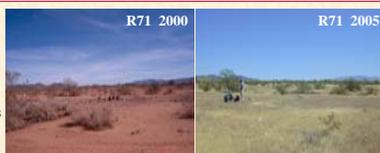
Vegetation: 2000 to 2005

Annuals
Annuals were not quantified in 2000 but visual inspection suggests considerable differences between assemblages of annuals at survey sites in 2000 and 2005. These differences may reflect above-average precipitation in the winter of 2004/2005.

Perennials
• Differences among compositions of perennial plants in 2000 and 2005 were assessed by calculating pair-wise, ecological dissimilarities (Sorenson distance metric) between sites.

► Assemblage structure of perennial plants was highly variable across both urban and desert sites but pair-wise ecological dissimilarities were significantly greater ($p < 0.001$) for urban sites, suggesting there was a greater change in perennial plant assemblages at urban sites compared to desert sites.

► Changes in community composition at desert sites were generally correlated to soil characteristics whereas changes in community composition at urban sites were more closely associated with anthropogenic characteristics.



Significant correlates to changes in composition of perennial plants between 2000 and 2005 at desert and urban sites

Desert sites	p	r
Extractable phos.	0.019	0.285
Organic carbon	≤ 0.037	≥ 0.253
Slope	0.004	0.347

Urban sites	p	r
Dist. to urban center	0.042	0.236
Population density	0.030	0.251