

INFESTATION MAPS AND SPATIAL STABILITY OF MAIN WEED SPECIES IN MAIZE CULTURE¹

Mapas de Infestação e de Estabilidade Espacial de Infestantes da Cultura do Milho

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ABSTRACT - A study on the spatial distribution of the major weeds in maize was carried out in 2007 and 2008 in a field located in Golegã (Ribatejo region, Portugal). The geo-referenced sampling focused on 150 points of a 10 x 10 m mesh covering an area of 1.5 ha, before herbicide application and before harvest. In the first year, 40 species (21 botanical families) were identified at seedling stage and only 22 during the last observation. The difference in species richness can be attributed to maize monoculture favouring reduction in species number. Three of the most representative species were selected for the spatial distribution analysis: *Solanum nigrum*, *Chenopodium album* and *Echinochloa crus-galli*. The three species showed an aggregated spatial pattern and spatial stability over both years, although the herbicide effect is evident in the distribution of some of them in the space. These results could be taken into account when planning site-specific treatments in maize.

Keywords: geo-statistics, kriging, precision agriculture, *Chenopodium album*, *Echinochloa crus-galli*, *Solanum nigrum*

RESUMO - Analisou-se a distribuição e estabilidade espacial de *Chenopodium album*, *Echinochloa crus-galli* e *Solanum nigrum* em um campo de milho do Ribatejo (Portugal) durante os anos de 2007 e 2008. A amostragem georreferenciada incidiu sobre 150 pontos em uma malha de 10 x 10 m, cobrindo uma área de 1,5 ha, antes da aplicação de herbicidas e antes da colheita. No primeiro ano foram identificadas 40 espécies (21 famílias botânicas), em fase de plântula; no final do segundo ano, apenas 22. A diferença no número de espécies pode ser atribuída à monocultura de milho, que contribui para a redução da riqueza específica. As três espécies mais representativas foram analisadas mediante um estudo geoestatístico, o que permitiu determinar o grau de variabilidade espacial e obter mapas de infestação por kriging baseados nos modelos que melhor se ajustavam a cada caso. Os resultados mostraram estabilidade espacial em todos os casos, exceto para *C. album* e *S. nigrum* em um dos períodos de tempo. Esses mapas, em conjunto com a estabilidade espacial geral registrada para as três espécies, são úteis para desenvolver programas de aplicação localizada de herbicidas, visando reduzir o impacto ambiental do recurso a esses produtos fitofarmacêuticos.

Palavras-chave: geoestatística, kriging, agricultura de precisão, *Chenopodium album*, *Echinochloa crus-galli*, *Solanum nigrum*.

INTRODUCTION

Weeds are usually aggregated in patches within the crop (Wiles et al., 1992; Gonzalez-Andujar and Saavedra, 2003). This is important from the point of view of management because herbicide use could be reduced by spraying only patches with weed infestation or by adjusting

herbicide rate (Barroso et al., 2004). In order to implement localized applications, it is necessary to know the spatial distribution map of the weed species, either identifying emerged weed flora or weed seed banks (Monquero et al., 2008; Izquierdo et al., 2009). Due to the high cost associated with the generation of these maps is important to

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minimize the frequency of the mapping, therefore, the spatial and temporal stability of the weed patches is critical. Species with a stable spatial distribution will not require the frequent generation of new maps (Barroso *et al.*, 2006; Longchamps *et al.*, 2012).

The aim of this study was to analyze the spatial distribution and spatio-temporal stability of the main weed species present in a maize field, using geostatistical techniques.

MATERIAL AND METHODS

The sample plot was situated in Golegã (Ribatejo) (4380373N, 467433E) in Portugal, with an area of 2.5 ha of which 1.5 ha was sampled with a total sample taking of 150 points (each point with an area of 10 m x 10 m) (Figure 1). The plot was under a beet-maize rotation, with maize planted in 2007 and 2008, when the sampling was made. Samples (seedlings plants m⁻²) were taken twice per year, one at the beginning of the cropping season (before herbicide application) and at its end (one week before harvest). Sampling points were georeferenced. Weed control in 2007 was conducted in postemergence (before completion of sampling) with atrazine, sulfonylureas, and bromoxynil. In the subsequent campaign, the herbicide treatment for weed control was sprayed again. Atrazine-resistant populations of *Chenopodium album* were present in the field.

A relative importance index (RI) was calculated for the major species for each sampling time, as $RI = (rD + rF) / 2$, where rD is the relative density and rF is the relative frequency (absolute frequency of each weed / total absolute frequency of all weeds) (Derksen *et al.*, 1993)

Geostatistical techniques (Gonzalez-Andújar *et al.*, 2001; Jurado-Exposito *et al.*, 2003; López-Granados, 2011) were used to describe and to map the spatial distribution of *Echinochloa crus-galli*, *Chenopodium album* and *Solanum nigrum*. A cross-validation procedure was performed to check model validity.

In order to determine the spatiotemporal stability, the statistical test proposed by Syrjala (1996) was used to detect whether the spatial distribution of the population has changed

over time. The null hypothesis was that there was no difference in the spatial distributions of two populations against the alternative hypothesis that some unspecified difference existed between the two distributions. This test is specifically designed to be insensitive to differences in the total abundances in the study area, but sensitive to differences in the distributions given the relative sizes of the two populations. The statistic to test the null hypothesis is the square of the difference between the two cumulative distribution functions, summed over all sampling locations, that is:

$$\Psi = \sum_{k=1}^k [\Gamma_1(x_k, y_k) - \Gamma_2(x_k, y_k)]^2 \quad (\text{eq. 1})$$

where $\Gamma_i(x_k, y_k)$ is the cumulative distribution function for the i population at the k sampling location and is defined as:

$$\Gamma_1(x_k, y_k) = \sum_{\forall x \leq x_k, \forall y \leq y_k} y_i(x, y) \quad (\text{eq. 2})$$

where $\Sigma \Gamma_i(x, y)$ is the sum of all normalized density observations, whose location (x, y) is such that $x \leq x_k$ and $y \leq y_k$, see Eqn 3.

$$y_i(x_k, y_k) = \frac{d_i(x_k, y_k)}{D_i} \quad (\text{eq. 3})$$

$$D_i = \sum_{k=1}^k d_i(x_k, y_k) \quad (\text{eq. 4})$$

where being $d_i(x_k, y_k)$ the value of the weed species density in a given sampling point (x_k, y_k) , for the i population and D_i the sum of all density observations for that species, which defines the normalized density observation.

RESULTS AND DISCUSSION

Fourty *taxa* were recorded and distributed over 21 families with predominance of *Poaceae*, *Asteraceae* and *Caryophyllaceae*. Eighty six percent of all species were annuals. Eighteen species were present on both years. Table 1 shows the relative importance index of the species evaluated in post-crop emergence,

Table 1 - Prevailing life cycle and relative importance index (rI) of weed species during two growing seasons (2007 and 2008)

Species	BAYER code	Botanical family	Life cycle	May 2007	Sep. 2007	May 2008	Sep. 2008
<i>Abutilon theophrasti</i> Medicus	ABUTH	Malvaceae	annual	5.8	1.6	4.7	1.1
<i>Amaranthus blitoides</i> S. Watson	AMABL	Amarantaceae	annual	1.8	0.8	0.0	0.0
<i>Amaranthus deflexus</i> L.	AMADE	Amarantaceae	annual	5.5	6.1	20.5	27.4
<i>Ammi majus</i> L.	AMIMA	Apiaceae	annual	1.0	1.0	0.9	0.0
<i>Anagallis arvensis</i> L.	ANGAR	Primulaceae	annual	5.1	1.3	0.0	0.0
<i>Atriplex patula</i> L.	ATPPA	Chenopodiaceae	annual	0.8	0.0	0.0	0.0
<i>Beta vulgaris</i> L.	BEAVX	Chenopodiaceae	annual	2.6	1.0	1.0	0.0
<i>Cerastium glomeratum</i> Thuill.	CERGL	Caryophyllaceae	annual	0.0	0.0	1.3	0.0
<i>Chenopodium album</i> L.	CHEAL	Chenopodiaceae	annual	6.6	3.1	5.1	3.1
<i>Convolvulus arvensis</i> L.	CONAR	Convolvulaceae	perennial	1.7	0.9	0.7	1.5
<i>Conyza bonariensis</i> (L.) Cronq.	ERIBO	Asteraceae	annual	0.0	1.4	0.0	0.0
<i>Cynodon dactylon</i> (L.) Pers.	CYNDA	Poaceae	perennial	0.8	0.0	0.7	0.8
<i>Cyperus esculentus</i> L.	CYPLO	Cyperaceae	perennial	1.0	0.0	0.0	0.0
<i>Cyperus longus</i> L.	CYPLO	Cyperaceae	perennial	0.0	0.0	0.0	0.0
<i>Cyperus rotundus</i> L.	CYPLO	Cyperaceae	perennial	1.4	0.0	0.3	0.0
<i>Datura stramonium</i> L.	DATST	Solanaceae	annual	5.9	2.2	3.8	3.8
<i>Digitaria sanguinalis</i> (L.) Scop.	DIGSA	Poaceae	annual	3.4	4.0	4.2	3.9
<i>Diplotaxis catholica</i> (L.) DC.	DIPCA	Brassicaceae	annual	1.8	0.6	3.3	0.0
<i>Echinochloa crus-galli</i> (L.) P. Beauv.	ECHCG	Poaceae	annual	7.0	6.7	14.7	23.8
<i>Kickxia spuria</i> (L.) Dumort.	KICSI	Scrophulariaceae	annual	3.1	1.5	0.5	0.0
<i>Lamium purpureum</i> L.	LAMPU	Lamiaceae	annual	0.5	0.0	0.0	0.8
<i>Oxalis acetosella</i> L.	OXACO	Oxalidaceae	annual	0.0	0.0	0.9	1.7
<i>Oxalis corniculata</i> L.	OXAPC	Oxalidaceae	annual	1.4	9.6	0.8	0.0
<i>Picris echioides</i> L.	PICEC	Asteraceae	biennial	0.6	1.0	0.4	0.8
<i>Piptatum miliaceum</i> (L.)	ORZMI	Poaceae	perennial	0.5	0.5	0.0	0.0
<i>Plantago major</i> L.	PLAMA	Plantaginaceae	perennial	0.5	0.0	0.0	0.0
<i>Poa annua</i> L.	POAAN	Poaceae	annual	0.9	20.2	4.8	11.9
<i>Polycarpon tetraphyllum</i> (L.) L.	POYTE	Caryophyllaceae	annual	3.3	3.3	0.5	0.0
<i>Polygonum persicaria</i> L.	POLPE	Polygonaceae	annual	0.0	0.0	0.2	0.0
<i>Portulaca oleracea</i> L.	POROL	Portulacaceae	annual	0.5	0.6	0.7	0.0
<i>Pseudognaphalium luteo-album</i> (L.) Hilliard		Asteraceae	annual	0.0	0.0	0.0	0.4
<i>Ranunculus muricatus</i> L.	RANMU	Ranunculaceae	annual	1.9	0.6	0.8	0.0
<i>Ranunculus trilobus</i> Desf.	RANTB	Ranunculaceae	annual	1.0	1.0	3.5	0.0
<i>Rumex crispus</i> L.	RUMPC	Polygonaceae	perennial	0.9	2.2	0.7	0.0
<i>Rumex pulcher</i> L.	RUMCR	Polygonaceae	perennial	1.4	0.8	0.5	0.0
<i>Setaria verticillata</i> (L.) P. Beauv.	SETVE	Poaceae	annual	0.0	0.0	7.5	1.9
<i>Solanum nigrum</i> L.	SOLNI	Solanaceae	annual	13.9	8.5	0.6	5.1
<i>Sonchus asper</i> (L.) Hill	SONAG	Asteraceae	biennial	2.3	2.5	9.8	0.0
<i>Sonchus oleraceus</i> L.	SONOL	Asteraceae	biennial	6.2	9.8	0.4	7.3
<i>Sorghum halepense</i> (L.) Pers.	SORHA	Poaceae	perennial	0.0	0.0	0.0	0.8
<i>Spergula arvensis</i> L.	SPRAR	Caryophyllaceae	annual	0.5	0.5	1.1	0.0
<i>Stellaria media</i> (L.) Vill.	STEME	Caryophyllaceae	annual	0.8	3.1	0.0	1.8
<i>Trifolium campestre</i> Schreber	TRFCA	Fabaceae	annual	0.5	0.5	0.6	0.0
<i>Verbena officinalis</i> L.	VEBOF	Verbenaceae	perennial	0.8	0.8	3.8	0.0
<i>Veronica persica</i> Poiret	VERPE	Scrophulariaceae	annual	4.2	3.9	0.3	2.6
<i>Xanthium strumarium</i> L.	XANST	Asteraceae	annual	2.1	0.0	0.0	0.0
Total density (seedlings m ⁻²)				113	64	236	71



the first of May (before application of herbicide) and the second of September (one week before harvest) for 2007 and 2008. Considering only the most important species ($rI > 1$) in 2007, the number of species was higher, 25-22 from May to September and the total density in the first observation was lower (113 seedlings m^{-2}) than in 2008. In the latter, the number of species from May to September and the total density were 15-14 species and 236 seedlings m^{-2} , respectively (Table 1). A modification of the cropping system from crop rotation (beet-maize) to maize monocropping could explain the decline in diversity from 2007 to 2008.

The three most abundant species (over 100 seedlings m^{-2}) during the first cropping season were *E. crus-galli*, *C. album* and *S. nigrum*. These results are in accordance with studies in maize crops in different areas from Spain and Portugal (Dorado et al., 2009).

The spatial distribution of *E. crus-galli*, *C. album* and *S. nigrum* was analysed and different models were fitted to each weed species (Table 2). There was no evidence of directional effect (anisotropy) in accordance with Halstead et al. (1990) and Cardina et al. (1995).

The results indicate that the three species showed an aggregated spatial distribution with varying degrees of spatial dependence over time. The nugget, the sill and the range varied among species and within species between

different sampling times (Table 2) possibly due to the effect of herbicide action.

The three weed species declined significantly during the last season, particularly *C. album* and *S. nigrum*. For the last sampling, made at the end of the cropping season of 2008, populations of *C. album* and *S. nigrum* virtually disappeared because of herbicide action.

A decrease in weed density of 70%, between the first and second observation, was registered in 2008 and could be attributed to herbicide efficacy. Weed density within the year decreased from 236 seedlings m^{-2} , before herbicide application, to 71 seedlings m^{-2} at harvest. The *C. album* population was resistant to atrazine and two herbicides were applied in post-emergence to increase control.

C. album patches were located mainly next to field margins and remained considerably stable across the period of two years of study (Figure 1). The concentration of patches near field edges is common in maize fields (Colbach et al., 2000) and could be attributed to seed emigration, lower levels of crop establishment and poor control in these areas.

S. nigrum showed a large decline in population at the end of the second cropping season following the herbicide application, which could significantly affect their spatial distribution (Figure 3).

Table 2 - Semivariograms adjusted to data for major maize weed species at different sampling dates

Species	Date	Fitted Model	Nugget	Sill	Range (m)	RSS
<i>Echinochloa crus-galli</i>	May 07	Linear Model	0.571	0.751	2262	0.0084
	Sep. 07	Gaussian Model	0.312	0.810	2193	0.0049
	May 08	Exponential Model	0.746	1.493	1651	0.0237
	Sep. 08	Exponential Model	0.669	1.339	612	0.0409
<i>Chenopodium album</i>	May 07	Spherical Model	0.408	1.069	866	0.0397
	Sep. 07	Spherical Model	0.163	0.327	827	0.0109
	May 08	Exponential Model	0.087	1.020	116	0.0661
	Sep. 08	Exponential Model	0.008	0.059	71	0.0002
<i>Solanum nigrum</i>	May 07	Gaussian Model	0.089	0.509	135	0.0141
	Sep. 07	Exponential Model	0.095	0.689	88	0.0200
	May 08	Spherical Model	0.412	0.825	520	0.0248
	Sep. 08	Spherical Model	0.013	0.142	132	0.0007

Nugget - the y-intercept of the model; *Sill* - the model asymptote; *Range* - the separation distance over which spatial dependence is apparent; *RSS* - Residual Sums of Squares.

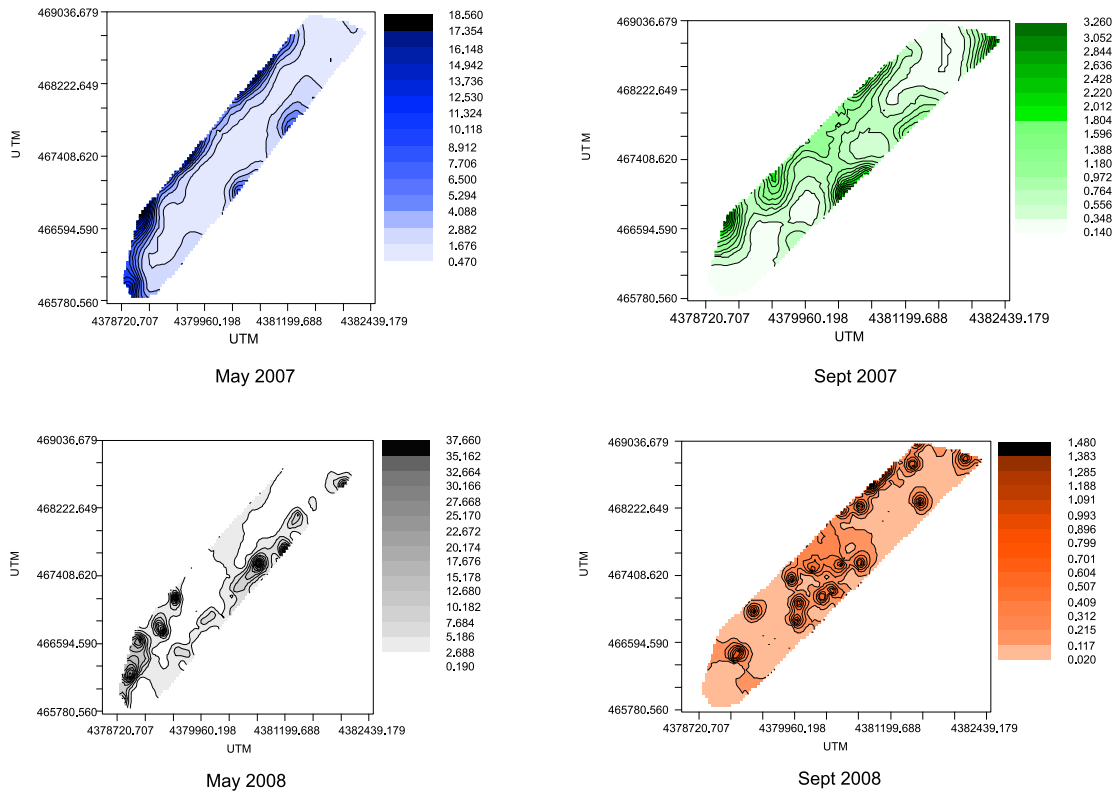


Figure 1 - Isoline kriged maps of *Chenopodium album* density (seedlings m⁻²) in 2007 and 2008.

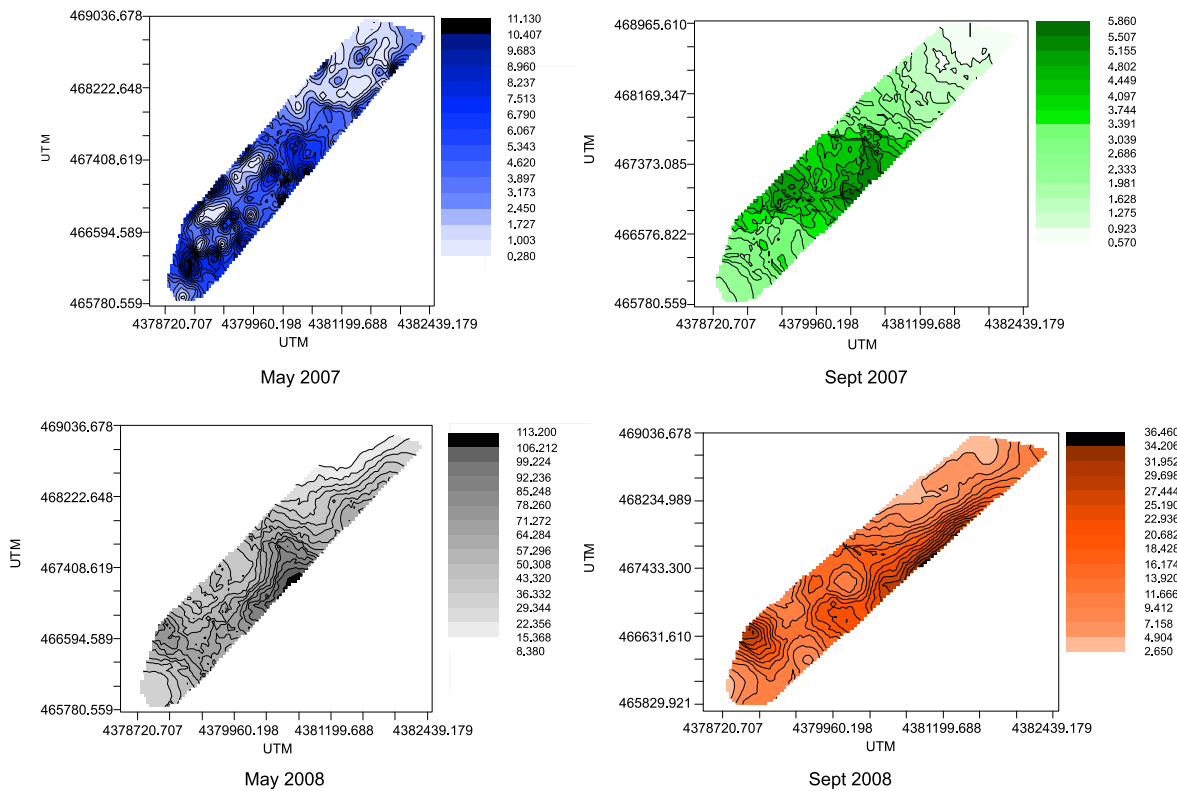


Figure 2 - Isoline kriged maps of *Echinochloa crus-galli* density (seedlings m⁻²) in 2007 and 2008.



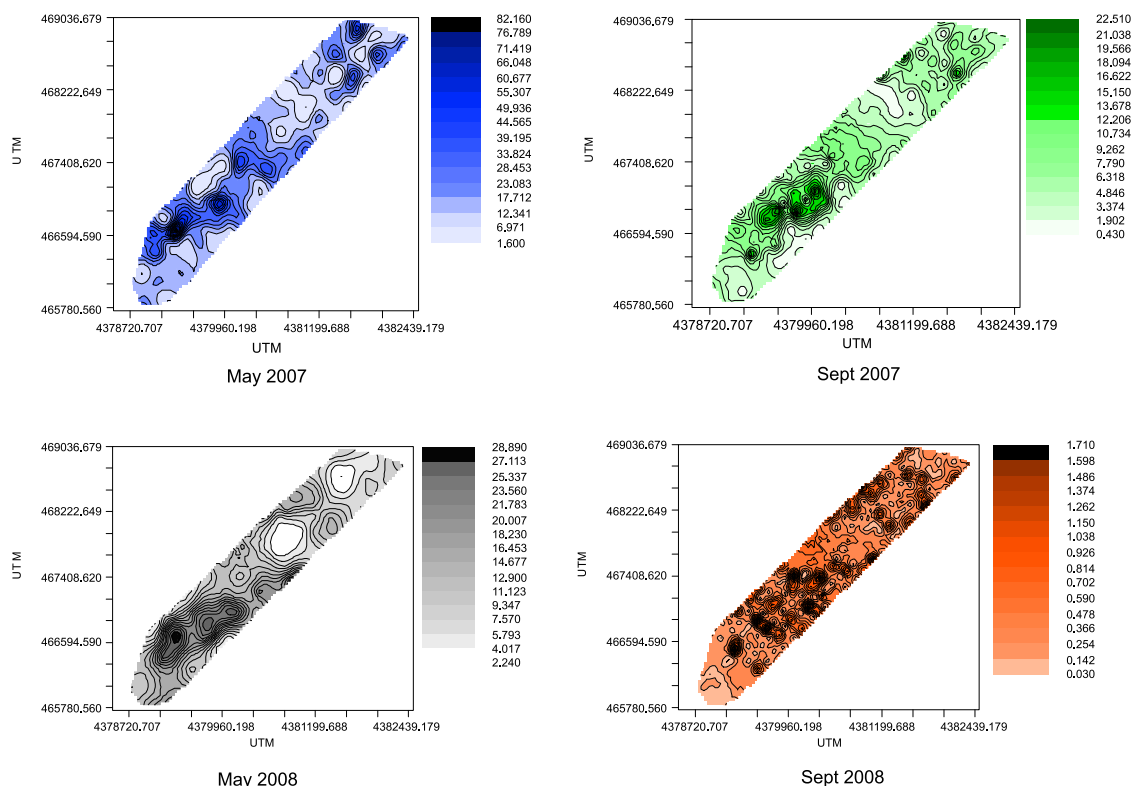


Figure 3 - Isoline kriged maps of *Solanum nigrum* density (seedlings m^{-2}) in 2007 and 2008.

The success of *E. crus-galli* (Figure 2) may be attributed to the production of large numbers of small, easily dispersed seeds per plant, possession of seed dormancy, rapid ability to flower under a wide range of photoperiods and relative tolerance of mature seeds to herbicide application (Maun & Barret, 1986).

By using the Syrjala's test to evaluate spatial and temporal persistence (Table 3), it can be concluded that *E. crus-galli* showed within and between years spatial stability. *C. album* showed spatial stability in the two last time periods and *S. nigrum* in the two first time periods (Table 3). It should be noted that the three species showed spatial stability between years (September 07/May 2008). The results of patch persistence over time vary between species depending on their bioecological characteristics, but need to be combined with weed management practices in the field, such as herbicide application and direction of machine work at cultivation and harvesting or soil fertility (Shiratsuchi et al., 2005).

In general, patch persistence is greater for perennial (*S. nigrum*) compared to annual weed species (*C. album* and *E. crus-galli*). However, some annuals with high seed production such as *Alopecurus myosuroides* is also spatially stable. This pattern could be due to the use of effective herbicides that do not allow weeds to produce seeds and contribute to restrain the dispersion of weed seed across the field.

In general, the three species showed spatiotemporal stability, suggesting that weed seedling distribution mapped in the first year are good predictors of future seedling distributions. This can open the door to use site-specific weed management for these species in maize in Portugal. Further studies are needed in Mediterranean conditions to allow for site-specific application of selective herbicides in the field. Recent technology using gadgets such as small unmanned aircraft systems (UAS) could contribute to the cost-effectiveness of sampling methods, one of the main constrains to practical application of mapping to weed management (Zhang & Kovacs, 2012; Ramussen et al., 2013).

Table 3 - Comparison of sample maps for the three main weed species according to statistics ϕ (Syrjala test)

Species	May-Sep. 07	Sep. 07-May 08	May-Sep. 08
<i>Echinochloa crus-galli</i>	0.185 ns	0.158 ns	0.693 ns
<i>Chenopodium album</i>	1.505*	0.972 ns	2.063 ns
<i>Solanum nigrum</i>	0.069 ns	0.239 ns	2.124**

ns: No significant; *: significant (P>0.05); **: significant (P>0.001).

Echinochloa crus-galli, *Chenopodium album* and *Solanum nigrum* were the main weeds found in the first year of study in the maize field, although the last two suffered a drastic decline during time and almost have disappeared after the second year. The three species showed an aggregated spatial pattern and showed spatial stability over both years, although the herbicide effect is evident in the distribution of some of them in space. These results could be taken into account when planning site-specific treatments in maize.

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It is with regret that we report that Edite Sousa (1953 - 2010), friend, colleague and co-author died in 2010.

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