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### ► To cite this version:

Abir Oudi, Mohamed Ali Chokri, Abdessalem Hammouda, Rim Chaabane, Riadh Badraoui, et al.. Physiological impacts of pollution exposure in seabird's progeny nesting in a Mediterranean contaminated area. *Marine Pollution Bulletin*, 2019, 142, pp.196-205. 10.1016/j.marpolbul.2019.02.056 . hal-02619725

**HAL Id: hal-02619725**

**<https://hal.inrae.fr/hal-02619725v1>**

Submitted on 26 Oct 2021

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1     **Physiological impacts of pollution exposure in seabird's progeny**  
2             **nesting in a Mediterranean contaminated area**

3

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14

15 **Abstract**

16

17 Aquatic wildlife is exposed through trophic transfer of hazardous substances to several threats  
18 inducing physiological impairments. We aimed at assessing the impact of contamination in one  
19 of the hot spots of pollution along Mediterranean coasts, the gulf of Gabes in Tunisia, on  
20 Common tern *Sterna hirundo*, a piscivorous top predator bird. Firstly, we compared the  
21 reproductive effort of breeding adults through clutch size distribution in three sites with  
22 different levels of pollution. Then, a battery of genotoxicity and oxidative stress biomarkers  
23 was carried out to assess physiological impairments in chicks. Whilst defense mechanisms  
24 showed a depletion, lipid peroxidation and genotoxicity increased significantly according to  
25 pollution level. The multi-biomarker approach used here, discriminated chicks according to  
26 contamination degree of their nesting sites. Increase in genotoxicity and oxidative stress were  
27 correlated to a decrease in chick body mass known to lead to long-term impacts on juvenile  
28 survival and recruitment in birds.

29

30 **Keywords:** biomarker, antioxidant, marine pollution, oxidative stress, *Sterna hirundo*,  
31 genotoxicity, Gulf of Gabes, reproductive impairment, chick body mass.

32

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## 36 1. Introduction

37 Pollution of the marine environment is a global issue owing to the discharge of urban, industrial  
38 and agricultural effluents into rivers, estuaries and coastal areas (Cravo *et al.*, 2012). Due to  
39 their ubiquity and highly persistent nature, a wide array of pollutants can accumulate in tissues  
40 of numerous marine organisms and are transferred across the food chain (Wu *et al.*, 2009). This  
41 often leads to rising concentrations (biomagnification) in top predators of marine trophic webs  
42 which may cause a prominent threat while their concentration in the environment remains  
43 below policy thresholds (Wu *et al.*, 2009; Nendza *et al.*, 1997). Indeed, dietary exposure  
44 represents the main source of contamination of predating marine birds and mammals, when  
45 direct uptakes of environmental pollutants (water, sediment, air) are of minor contribution  
46 (Nendza *et al.*, 1997). Combined with high trophic status and long-life span, life history traits  
47 of upper predators may exacerbate their vulnerability to chronic contamination and their  
48 biological costs (Rowe, 2008). On the one hand, long-lived species are prone to accumulate  
49 high loads of persistent chemicals during their life cycle including early developmental stages  
50 and on the other hand, they have a low capacity for genetic adaptation due to slow renewal of  
51 generations (Rowe, 2008).

52 Several biological defects have been associated with high body burdens of chemicals in aquatic  
53 wildlife and fish species ultimately affecting their fitness. Reduced eggshell thickness  
54 (Cortinovis *et al.*, 2008), offspring developmental defects (e.g., abnormalities, impaired growth,  
55 developmental instability) (Hoffman *et al.*, 1986; Jenssen *et al.*, 2010; Letcher *et al.*, 2010) and  
56 physiological disturbances of the exposed organisms such as endocrine disruption and  
57 altered vitamin homeostasis have been demonstrated (Letcher *et al.*, 2010; Rolland, 2000).  
58 There is increasing awareness of the ubiquitous role of oxidative stress inducing physiological  
59 disturbances due to to chemicals exposure (Valavanidis *et al.*, 2006). When reactive oxygen  
60 species (ROS) production overwhelms antioxidants, a state of physiological imbalance known  
61 as “oxidative stress” occurs (Valavanidis *et al.*, 2006). This altered redox homeostasis may  
62 significantly alter biomolecules (DNA damage, lipid peroxidation and protein oxidation), affect  
63 individual growth, survival and reproduction of the exposed organisms which in turn might lead  
64 to population declines (Koivula and Eeva, 2010; Viarengo *et al.*, 2007; Goutte *et al.*, 2014).  
65 Nevertheless, studies addressing the relationship between contaminant-induced stress and  
66 individual fitness alterations in wild populations remain scarce (Provencher *et al.*, 2016). While  
67 oxidative stress related to pollutants (especially metals) has been demonstrated in experimental  
68 studies in birds, it has rarely been studied in wild free-ranging bird species (Koivula and Eeva,

69 2010; Espín *et al.*, 2014 but see Costantini *et al.*, 2014). Therefore, there is an increasing interest  
70 in assessing the fate of chemicals, individual impacts and ecological risks of coastal pollution  
71 on bird populations. Expanding our comprehension about tolerance and detoxification ability  
72 of birds is particularly important as it may help in predicting how other species respond to  
73 chemical stressors in their environment. It is of main interest to identify most sensitive species  
74 to chemical pollution and to understand mechanisms that may be responsible of populations  
75 decline in polluted areas (Provencher *et al.*, 2016; Koivula and Eeva, 2010).

76 Biomarkers are “biochemical, cellular, physiological or behavioral variation that can be  
77 measured in tissue or body-fluid samples or at the level of whole organisms providing evidence  
78 of exposure to, and/or effects of, one or more chemical pollutants (and/or radiation)” (Depledge  
79 and Fossi, 1994). They are considered as early warning tools, which used together in  
80 combination, investigating several relevant physiological functions, provide an integrated  
81 measure of exposure to, and/or effects of, contaminants in the aquatic environment (Cajaraville  
82 *et al.*, 2000; Van derOost *et al.*, 2003). They could be predictive of population dynamic  
83 disturbances, and thus allow anticipating detrimental changes at higher levels of biological  
84 organization (population, community or ecosystem) (Cajaraville *et al.*, 2000; Van derOost *et al.*  
85 *et al.*, 2003; Vasseur and Cossu-Leguille, 2003; Amiard-Triquet *et al.*, 2013).

86 Yet, since various antioxidants can be employed differently depending on species, in defense  
87 against oxidative damage, one biomarker of oxidative stress is not reliable enough to reflect  
88 changes in the complex homeostasis (Koivula and Eeva, 2010; Espín *et al.*, 2014). Thus, it is  
89 recommended to investigate simultaneously a set of biomarkers of exposure and effect,  
90 especially when individuals are exposed to complex mixtures of environmental stressors  
91 (Cajaraville *et al.*, 2000; Cravo *et al.*, 2009). Often used in multi-criteria approaches  
92 investigating several relevant biological endpoints, biomarkers are efficient tools to assess the  
93 health status and ecological risks of organisms exposed to environmental pollutant mixtures.  
94 Such multi-biomarker approaches have been successful in covering various relevant infra-  
95 individual responses, including oxidative stress with invertebrates and fish species (Chèvre *et al.*  
96 *et al.*, 2003; Viarengo *et al.*, 2007; Damasio *et al.*, 2011; Benali *et al.*, 2015; Santos *et al.*, 2016).

97 The gulf of Gabes (southeastern coasts of Tunisia) is one of the main hotspot of pollution along  
98 Mediterranean coasts where high levels of polycyclic aromatic hydrocarbons (PAH) and heavy  
99 metals have been detected in water and sediments (Zaghden *et al.*, 2014; El Zrelli *et al.*, 2015;  
100 Rabaoui *et al.*, 2015; Fourati *et al.*, 2017). In this area, several common tern colonies (*Sterna*  
101 *hirundo*) are established during the breeding season. This seabird used in this work as a model  
102 species is common over a wide geographical range with low annual reproductive output. As a

103 long-lived species and a top-level predator of aquatic ecosystems consuming almost exclusively  
104 fish, this bird is prone to bioaccumulate xenobiotics inducing adverse impacts including  
105 oxidative stress and potentially changes in reproductive activity (Van derOost *et al.*, 2003;  
106 Rowe, 2008). In order to achieve a comprehensive evaluation of adverse outcomes of pollution  
107 on birds we conducted a comparative study on tern colonies breeding in three sites subjected to  
108 different level of pressures. As income breeders, like common terns, depend on locally acquired  
109 resources to form eggs rather than on reserves, contamination may be a major constraint for  
110 reproduction (Ezard *et al.*, 2007). So, our first objective was to detect the impact of pollution  
111 on the reproductive potential of tern females. We hypothesized that chemical pressures may  
112 prevent females from increasing their reproductive effort through laying reduced clutches  
113 compared to their conspecifics breeding in a relatively clean area. Our second objective was to  
114 assess the alteration of physiological status and genotoxic damage of chicks by carrying out a  
115 multibiomarker approach using a combination of oxidative stress and genotoxicity biomarkers.  
116 The use of chicks presents two advantages. First, the effect of age-related bioaccumulation is  
117 avoided, since the exposure time is known and similar for all individuals (Quirós *et al.*, 2008).  
118 Second, because common terns are known to have a limited foraging range around their colony  
119 site (maximum foraging range of 9 km; Rock *et al.*, 2007) and all the chicks are provisioned  
120 with prey caught by parents in the surroundings of the colony, the results should reflect the  
121 impact of local pollution (Quirós *et al.*, 2008). Hence, cellular antioxidant defenses, namely  
122 activities of two antioxidant enzymes (superoxide dismutase (SOD) and catalase (CAT)), and  
123 two exogenous antioxidants (vitamin A and vitamin E) besides a biomarker of impact (TBARS)  
124 were selected as oxidative stress biomarkers. These biomarkers have been widely used in many  
125 field studies in order to assess the extent of pollution in different environments (An *et al.*, 2012;  
126 Cravo *et al.*, 2009, 2012). The micronucleus assay (MN) reflecting chromosomal damages was  
127 used as a biomarker of genotoxicity. Although investigating genotoxicity is valuable in wildlife  
128 risk assessment, a few studies have been conducted on wild birds especially for biomonitoring  
129 purposes (e.g., Quirós *et al.*, 2008; Skarphedinsdottira *et al.*, 2010). In overall, the present work  
130 aims at investigating whether contamination affects reproduction of common tern on a pollution  
131 gradient impacting parental nesting capacities. We also investigated physiological disturbances  
132 and body mass of offspring reared and fed in nesting sites under contrasted chemical pressures.

133

## 134 **2. Materials and methods**

### 135 **2.1. Study sites**

136 Field work was carried out in three sites on the Eastern coasts of Tunisia which reflect different  
137 levels of pollution (Figure 1). The first nesting site (33°54'N 10°06'E) is located in the Gulf of  
138 Gabes close to an industrial complex in the region of Chatt-Essalem known to be one of the  
139 most contaminated area in Gabes (Zaghden *et al.*, 2014; El Zrelli *et al.* 2015; Rabaoui *et al.*,  
140 2015). Local industrial activities in the area led to the contamination of the aquatic ecosystems  
141 by pollutant mixtures from multiple effluents rejected by the phosphate treatment complex and  
142 the Fluor Chemical Industries (ICF) (Rabaoui *et al.*, 2015) as well as those from the wastewater  
143 treatment plant of Chatt-Essalem and urban activities. Chatt-Essalem is located between the  
144 fishing harbor and the commercial harbor, making the water renewal low due to very weak  
145 currents and hydrodynamics (Rabaoui *et al.*, 2015) which minimize chemicals dilution and  
146 promote local accumulation. The second site, in northern Gabes gulf (34°39'N 10°42 E), is  
147 located about 12 km south Sfax city. The southern coastal area of Sfax is subjected to industrial  
148 pollution due to the discharge of chemical effluents from the phosphoric acid plant (SIAPE)  
149 besides a municipal wastewater treatment plant. The approximate quantities of phosphogypsum  
150 dumped from the phosphate treatment plants are 135 million tons in Chatt-Essalem and 30  
151 million tons in Sfax (Ben Amor and Jomaa, 2012).

152 The Monastir salina (35°45'N 10°42'E) is in the gulf of Hammamet. This area was considered  
153 as a reference site because industrial or urban effluents are much lower than the other sites.

154 To describe the level of anthropogenic pressures on these sites, we investigated the landscape  
155 use in the area. A radius of 10 km around each nesting area was chosen to measure the surface  
156 of agricultural and natural habitats, of urban areas and the surface of industrial activities. Maps  
157 were originally produced from a Landsat TM image with 30 m resolution taken in 2013. Image  
158 processing included atmospheric and geometric corrections was performed using ENVI Flash.  
159 Identification and digitization of geographic features were performed through ArcCatalog and  
160 ArcMap modules incorporated in ARCGIS 9.0. The surface of each area was determined using  
161 the calculate geometry tool of ARCGIS.

162

## 163 **2.2.Data collection and blood sampling**

164 Colonies of common tern were visited in the three sites throughout the nesting period from end  
165 of April to early July 2015. However, only synchronous nests and chicks from these sites were  
166 considered in our study and blood sampled. Each nest was marked with a small numbered  
167 wooden stick 40 to 50 cm apart from the nest. All nests were visited every 2-3 days in Chatt-  
168 Essalem, every 2-7 days in Sfax salina and every 2-12 days in Monastir Salina to check nest

169 content. Visits were more frequent during laying eggs to determine laying date and clutch size  
170 of nests. Clutch size was considered as completed when the number of eggs did not change  
171 between at least two visits. Since some chicks could have been predated between our field  
172 sessions, we could not rigorously assess hatching success for several clutches. In both Monastir  
173 and Sfax salina, chicks were easily detected because they hide between rocks of a narrow dike  
174 formed by soil and small rocks and surrounded from both sides by water. However, to prevent  
175 wandering of chicks or their escape after hatching in Chatt-Essalem site, nests were fenced  
176 individually with fishing nets (about 0.30×0.75m). The enclosures included natural or added  
177 cover so that chicks can hide and find shade and were open at the top to allow free access by  
178 adult birds which continued normal care of chicks even in presence of fences.  
179 Three weeks after hatching, chicks were weighed by a fixed spring scale (Model Kern HDB  
180 5k5) or a Pesola and blood samples (volume equivalent to 1% of body weight) were taken from  
181 the brachial vein. Samples were then transported to the laboratory. A droplet of whole blood  
182 was taken to prepare smears (2 smears per chick) for the micronucleus assay while the rest of  
183 blood was centrifuged at 3500 rpm for 10 min. Both plasma and cell pellets were stored at -  
184 20°C until analyzed. Samples were taken from 25, 21 and 19 chicks respectively in Chatt-  
185 Essalem, Sfax salina and Monastir salina nesting sites.

186

## 187 **2.3. Biomarker analyses**

188

### 189 **2.3.1. Thiobarbituric acid reactive substances (TBARS)**

190 The TBARS assay is commonly used to quantify oxidative stress by measuring peroxidative  
191 damages to lipids that result from free radical generation (Valavanidis *et al.*, 2006). The  
192 intensity of lipid peroxidation can be approximated as the level of endproducts such as  
193 malondialdehyde (MDA) and other aldehydes which are appraised with thiobarbituric acid  
194 hence the name of the assay, thiobarbituric acid-reactive substances (TBARS) (Valavanidis *et*  
195 *al.*, 2006). Biological samples were combined with 1.5 ml TBA 0.8%, 1.5 ml trichloroacetic  
196 acid (TCA) 20% and 800 µl distilled water in the presence of 200 µl of sodium dodecyl sulfate  
197 (SDS) 8.1%. The incubation mixture was heated for 30 min at 90°C. After cooling, it was  
198 centrifuged (4000 rpm, 15 min) and the TBARS concentration was determined based on the  
199 absorbance at 532 nm. Results were expressed as nmol/ml.

200

### 201 **2.3.2. Antioxidant enzymes activities**



202 The Superoxide dismutase (SOD) activity was assessed, spectrophotometrically at 505 nm  
203 according to [McCord and Fridovich \(1969\)](#), by determining its ability to inhibit the  
204 photoreduction of cytochrome C by the superoxide anion. One unit of SOD represents the  
205 amount that inhibits the photoreduction of cytochrome C by 50%. SOD activity was expressed  
206 as U/ml.

207 Catalase (CAT) activity was measured spectrophotometrically in ultraviolet accordingly to the  
208 protocol from [Aebi \(1984\)](#) by following the consumption of H<sub>2</sub>O<sub>2</sub> at 240 nm. Decays in  
209 absorbance were recorded in the reaction mixture consisted of 500 µl of H<sub>2</sub>O<sub>2</sub> (0.03M), 950 µl  
210 of potassium phosphate buffer (0.05M; pH = 7.0) and 50 µl of sample.  
211 One unit of CAT was defined as the amount of enzyme that decomposes 1 µmol of H<sub>2</sub>O<sub>2</sub> per  
212 minute.

### 213 **2.3.3. Plasmatic antioxidant vitamins**

214 Plasmatic vitamins A (Vit A) and E (Vit E) were estimated by High Performance Liquid  
215 Chromotography (HPLC) in isocratic mode using the methodology reported by [Ferns et al](#)  
216 [\(2000\)](#). Plasma was firstly deproteinized with ethanol containing internal standards (retinyl  
217 acetate and  $\alpha$ -tocopheryl acetate) and then it underwent lipid extraction with hexane. After  
218 evaporation, the dry residue was dissolved and diluted with methanol. 50 µl of sample was  
219 injected onto a C18 (15 cm; 4.6 mm) column using a mixture of methanol *n*-butanol and water  
220 as mobile phase (v/v/v; 98, 5:5:5) at a flow rate of 1.7 ml/min. Detection was carried out at 330  
221 nm for the retinol acetate and at 292 nm for the  $\alpha$ -tocopherol. Results were expressed in mg/l.

### 222 **2.3.4. Micronucleus assay**

223  
224 Micronucleus assays were performed as described by [Skarphedinsdottir et al \(2010\)](#). Briefly,  
225 the blood smears were stained with 4 % Giemsa (> 10 min). The slides were rinsed in distilled  
226 water, air-dried, and coded for blind scoring under optical microscope at 100× magnification.  
227 Specimens of 25, 20 and 19 chicks respectively from Chatt-Essalem, Sfax and Monastir were  
228 randomly analyzed by a single observer. The frequency of micronucleated erythrocytes as well  
229 as several erythrocytes nuclear abnormalities described in Figure 2 were scored for 5,000  
230 erythrocytes in each investigated specimen following the zig-zag model (to avoid crossing the  
231 same field more than once). The criteria for identifying micronucleus were: (1) same color and  
232 intensity as the cell nucleus (2) size smaller than or equal to 1/5 of the main nucleus (3) rounded  
233 shape with a nuclear membrane (4) clearly detached from the cell nucleus with intra-  
234 cytoplasmatic location.

## 2.4. Statistical analysis

235  
236 The first objective was to compare clutch size distribution between sites. A chi-square test was  
237 conducted comparing proportion of nests with 3 eggs, 2 eggs and 1 egg among sampled sites.  
238 For biomarkers data, inter-site differences of biomarkers responses and chick mass were tested  
239 using non-parametric Kruskal-Wallis and Wilcoxon pairwise comparison tests. We investigated  
240 the micronucleus probability of detection between sites using Generalized Linear Mixed  
241 Models (GLMM) with a binomial error. All erythrocytes without micronucleus were coded as  
242 “0” while erythrocytes with micronucleus were coded “1” so that we modeled the probability  
243 of exhibiting a micronucleated erythrocyte. An ‘individual’ random effect was added to the  
244 model linking the 5,000 erythrocytes analyzed to the same individual. The fitted model was  
245 compared to the null model based on an Analysis of Deviance (ANODEV). To summarize  
246 biomarker responses, we implemented a principal component analysis (PCA) to analyze  
247 correlations between biomarker responses of the six biomarkers measured in chicks. The site  
248 location was added as a supplementary information which has no influence on the principal  
249 components but add supplementary information for a better interpretation of the inertia. Finally,  
250 the first axis of the PCA discriminating chicks according to their biomarker responses was used  
251 to assess the relationship between chick biomarker responses hereafter considered as “chick  
252 physiological status” and their masses three-weeks post-hatching using a linear regression.  
253 All data are represented as the mean  $\pm$  standard error (SE) and statistical significance was  
254 defined at  $p \leq 0.05$ . The entire statistical analysis was carried out using R.3.2.2 software (R  
255 Development Core Team, 2011) with the packages “lme4” for GLMM, “PMCMRplus” for  
256 pairwise multiple comparisons (Pohlert *et al.*, 2018) and “FactoMineR” for multivariate  
257 analysis (Husson *et al.*, 2012).

## 3. Results

### 3.1. Evaluation of landscape use around the monitored sites

260  
261 Data showing land use and the ratio of different types of pressures present 10 km around each  
262 site are presented in Table 1 and Supplementary materials 1. Land cover results showed that  
263 Chatt-Essalem nesting site has less natural and agricultural zone (77.17%) than Sfax (84.34%)  
264 and Monastir salina (88.35%). Additionally, there is a slight gradual increase in urban area from  
265 Monastir (11.39%) to Sfax (13.16%) and Chatt-Essalem (18.59%). The site of Chatt-Essalem  
266 (4.24% of industrial area) is conspicuously closer (distance = 0 km) and 1.70 times more

267 industrialized than Sfax (distance = 1.8 km; 2.5% of industrial area) and 16.31 times more  
268 industrialized than Monastir salina (distance = 4.7 km; 0.26% of industrial area).

269

### 270 **3.2.Clutch size distribution**

271 Clutch size ranged from 1 to 3 eggs but there was a predominance of nests with 3 eggs in all  
272 areas (Table 2). Frequencies of clutches of 1, 2 and 3 eggs were 4.9%, 9.7% and 85.4% in  
273 Monastir, 6.3%, 16.3% and 77.4% in Sfax and 5.8%, 10.6% and 83.6% in Chatt-Essalem  
274 respectively. No significant difference was detected on clutch size distribution among the 3  
275 sites ( $\chi^2 = 9.1$ ;  $df = 4$ ;  $p = 0.06$ ).

276

### 277 **3.3.Biomarker analyses**

#### 278 **3.3.1. Biomarkers of impacts: Lipid peroxidation and DNA damage**

279 Peroxidative damage measured as TBARS in chicks showed significant differences between  
280 the three sites (KW- $\chi^2 = 32.65$ ;  $df = 2$ ;  $p < 0.001$ ). The level of lipid peroxidation was  
281 significantly higher in chicks from Chatt-Essalem and Sfax ( $103.99 \pm 4.52$  and  $74.89 \pm 2.55$   
282  $\text{nmol.ml}^{-1}$  respectively) compared to chicks from the reference site ( $64.31 \pm 2.64 \text{ nmol.ml}^{-1}$ ;  $W$   
283  $= 331$ ,  $p < 0.001$  and  $W = 316.5$ ,  $p < 0.01$  respectively; Figure 3). Concerning micronucleus,  
284 the probability to detect micronucleus between sites was significantly recorded ( $p < 0.01$ ). The  
285 between sites fitted model was significantly different from the null model (AIC = 373.12 and  
286 383.51), confirmed by ANODEV comparison of the fitted model to the null model ( $\chi^2 = 14.38$ ;  
287  $df = 2$ ;  $p < 0.01$ ). Coefficients of the fitted model highlighted significant lower probability of  
288 observing a micronucleated erythrocyte in the reference site ( $0.13 \pm 0.03 \%$ ; Figure 3) than in  
289 Sfax salina ( $0.50 \pm 0.15 \%$ ;  $p < 0.01$ ) and Chatt-Essalem ( $0.36 \pm 0.11 \%$ ;  $p < 0.01$ ).

290

#### 291 **3.3.2. Biomarkers of defense: antioxidant enzymes (SOD and** 292 **CAT)**

293 Highly significant differences between sites were recorded for both SOD (KW- $\chi^2 = 11.39$ ;  $df =$   
294  $2$ ;  $p < 0.01$ ) and CAT activities (KW- $\chi^2 = 11.26$ ;  $df = 2$ ;  $p < 0.01$ ). These two enzymes exhibited  
295 lower activities at Chatt-Essalem (SOD =  $7.62 \pm 0.52 \text{ U.ml}^{-1}$ ; CAT =  $68.98 \pm 3.82 \text{ U.ml}^{-1}$ ;  
296 Figure 4) and Sfax (SOD =  $8.71 \pm 0.28 \text{ U.ml}^{-1}$ ; CAT =  $76.95 \pm 3.55 \text{ U.ml}^{-1}$ ) in comparison with  
297 the Monastir reference site (SOD =  $9.74 \pm 0.37 \text{ U.ml}^{-1}$ ; CAT =  $88.45 \pm 4.04 \text{ U.ml}^{-1}$ ). However,  
298 this decrease was significant only between Chatt-Essalem and Monastir ( $W = 274$ ,  $p < 0.01$  for  
299 SOD and  $W = 275$ ,  $p < 0.01$  for CAT; Figure 4).

300

301 **3.3.3. Biomarkers of defense: plasmatic antioxidant vitamins**  
302 **(Vit.A and Vit.E)**

303 Significant differences between sites were observed in Vitamin A ( $KW-\chi^2 = 9.03$ ;  $df = 2$ ;  $p =$   
304  $0.01$ ) and Vitamin E ( $KW-\chi^2 = 9.68$ ;  $df = 2$ ;  $p < 0.001$ ). The level of Vit.A demonstrated that  
305 the content in chicks from Chatt-Essalem ( $0.154 \pm 0.010 \text{ mg.L}^{-1}$ ) was significantly depleted  
306 compared to Monastir ( $0.188 \pm 0.009 \text{ mg.L}^{-1}$ ;  $W = 287.5$ ,  $p < 0.01$ ). In Sfax salina, although a  
307 lower value was reported ( $0.179 \pm 0.007 \text{ mg.L}^{-1}$ ), no significant difference was observed neither  
308 with the Monastir reference site nor with Chatt-Essalem (Figure 5). The level of Vit. E  
309 demonstrated that the content in chicks from Sfax salina ( $1.84 \pm 0.06 \text{ mg.L}^{-1}$ ) was significantly  
310 depleted compared with Monastir ( $2.19 \pm 0.07 \text{ mg.L}^{-1}$ ;  $W = 311$ ,  $p < 0.01$ ). In Chatt-Essalem  
311 although a lower value was reported ( $1.93 \pm 0.08 \text{ mg.L}^{-1}$ ), no significant difference was  
312 observed neither with the Monastir reference site nor with the Sfax salina (Figure 5).

313  
314 **3.3.4. Three-week-old chick mass**

315 Significant differences between sites were observed in chick masses ( $KW-\chi^2 = 6.87$ ;  $df = 2$ ;  $p$   
316  $= 0.03$ ). The pairwise comparison demonstrated that three-week-old chick masses from Chatt-  
317 Essalem ( $117.54 \pm 13.53 \text{ g}$ ) were significantly lower compared to Monastir ( $126.55 \pm 9.20 \text{ g}$ ;  
318  $W = 101.5$ ,  $p = 0.05$ ). Masses of three-week-old chicks from Sfax ( $117.19 \pm 10.47 \text{ g}$ ) were  
319 significantly lower compared to Monastir ( $W = 98.5$ ,  $p = 0.01$ ) but no significant differences in  
320 chick mass were observed between Chatt-Essalem and Sfax sites.

321  
322 **3.4. Correlations between biomarker responses in chicks and their body mass**

323 The first two components of the PCA accounted for 55.13% of the total dataset (Figure 6). The  
324 first component explained 34.64% of the variance and pointed out that SOD and CAT were  
325 positively correlated and negatively correlated with TBARS. The second component explained  
326 20.48% of the overall data variance, mainly explained by the level of Vit. A and Vit. E which  
327 were positively correlated and inversely correlated to the level of MN. Based on the biomarker  
328 responses, PCA allowed a clear discrimination of chicks from Monastir to Chatt-Essalem  
329 according to their physiological status in a gradual increase of anthropogenic pressure (Figure  
330 6). Finally, an increase of the PCA-Axis 1 values representing a decrease in physiological stress  
331 detected in chicks was significantly correlated to an increase in three week-old chick mass ( $R^2$   
332  $= 0.20$ ;  $F\text{-value} = 13.85$ ;  $df = 55$ ;  $p < 0.01$ ; Figure 7).

333  
334 **4. Discussion**

335 To our knowledge, this study is the first to provide evidences of physiological and genotoxic  
336 disturbances in wild common tern (*Sterna hirundo*) under natural conditions using a multi-  
337 biomarker approach. No significant difference was found in the reproductive investment of  
338 breeding adults, reported as the number of eggs laid per nest and their distribution among sites  
339 with contrasted level of contamination. We carried out a battery of relevant biomarkers  
340 investigating antioxidant enzymes, dietary exogenous vitamins (A and E), a biomarker of lipid  
341 peroxidation and a biomarker of genotoxicity. This multi-biomarker approach outlined  
342 physiological redox homeostasis disturbances and genotoxicity in *Sterna hirundo* chicks reared  
343 and fed in polluted areas which could imperil in the long-term their development. We  
344 demonstrated that chicks with disturbed physiological status also presented a decrease in body  
345 mass compared to reference conditions. As chick body mass is a useful fitness-related trait  
346 known as the predictor most commonly associated with post-fledging survival besides hatching  
347 date in birds, this work outline that the measured biomarker responses could affect their fitness  
348 over the course of their life span and lead to long-term changes in demographic traits and  
349 thereby in population growth and dynamics.

#### 350 **4.1.Clutch size**

351 As a significant part of the energy of birds under chemical pressure would be allocated in  
352 xenobiotic metabolization and costly defenses, and since reproduction is physiologically an  
353 energy demanding process, one may expect that increased energy expenditure and chemically-  
354 induced stress might divert resources away from reproductive investment in tern populations  
355 which in turn reduce chances for fueling laying efforts. In our study, the reproductive output of  
356 common tern, investigated in term of number of eggs laid per nest was similar with  
357 predominance of 3-egg clutches in all monitored areas. This may suggest that pollution does  
358 not represent a constraint for reproduction considering the number of eggs in polluted areas and  
359 that females seem to be resistant or balancing their energy requirements between these critical  
360 biological traits. [Mateo et al \(2004\)](#) highlighted that one of two common tern colonies they  
361 studied in a contaminated area from the Ebro delta (Spain) tend to have smaller clutch size,  
362 lower hatching success and higher yolk organochlorine compounds. These authors suspected  
363 that this is due to differences in fish consumed as terns from Banyà's colony feed on demersal  
364 and benthic fish species that are more exposed to contaminants from sediments whereas terns  
365 from Fangar's colony would feed mostly on pelagic species of small clupeiformes that are less  
366 exposed to contaminants. [Wiersma et al \(2004\)](#) have demonstrated the negative compromise  
367 between reproduction and oxidative protection in zebra finches. They showed that birds trading-

368 off their oxidative protection against the reproductive output, suffer decreased antioxidant  
369 enzymes activities when having experimental increased clutch size. However, [Markó et al](#)  
370 [\(2011\)](#) did not find a significant correlation between clutch size (number of eggs) and the  
371 oxidative status of collared flycatcher (*Ficedula albicollis*). In our work, we did not assess the  
372 oxidative status of females but it would be of great interest to further analyze correlations  
373 between clutch sizes and parental physiological status. An alternative explanation that may  
374 explain no differences in clutch sizes could be that even if the clutch size was not affected in  
375 polluted areas, a trade off may be mirrored in other functions ([Markó et al., 2011](#)) such as egg  
376 volume or the quantity of maternal antioxidants and xenobiotic defenses invested into eggs.  
377 Thus, further work will be needed to investigate these points. Furthermore, one limitation to  
378 our work is that it was restricted to the progeny during early life stages, so that it might be  
379 biased toward good quality individuals that succeed in hatching and chick rearing. Yet this  
380 potential bias is conservative, it should reduce the differences between our study sites if only  
381 individuals with lower level of contamination succeed in breeding. Future works, studying the  
382 entire population, notably adults including non-breeding ones or those which fail breeding early  
383 in the season, and potentially over several years for instance using Capture-Recapture approach  
384 are definitely needed to further elucidate this point.

385  
386

#### 4.2. Biomarker analyses

387 A significant rise in the level of TBARS was measured according to the degree of pressures  
388 surrounding nesting sites. The highest value was observed in chicks from Chatt-Essalem and  
389 Sfax in a lesser extent while those from Monastir showed the lowest value. An increase in  
390 TBARS levels mean an excessive production of free radicals damaging polyunsaturated lipids.  
391 After hatching, common tern parents feed their progeny with fish caught locally in the feeding  
392 habitats ([Burger and Gochfeld, 1993](#)). Thus, overgeneration of free radicals might be promoted  
393 both directly or indirectly by xenobiotic metabolization entering the organisms mostly due to  
394 food intakes in aquatic bird top predators ([Limón-Pacheco and Gonsebatt, 2009](#); [Valavanidis et](#)  
395 [al., 2006](#); [Quirós et al., 2008](#)). In order to make accurate inferences about the level of oxidative  
396 stress experienced by chicks, we measured the TBARS as a biomarker of impact which can  
397 reflect the level of oxidative stress and biomarkers of defenses that allow organisms to cope  
398 with the presence of pollutants in their environment ([Valavanidis et al., 2006](#); [Amiard-Triquet,](#)  
399 [2013](#)). Thus, we took in account the cooperative tasks of the antioxidant system, choosing  
400 enzymatic biomarkers and non-enzymatic ROS scavenger antioxidants ([Espín et al., 2014](#)).  
401 Antioxidant molecules play a key role in defense mechanisms by counteracting free radicals

402 formation and their harmful effects. In this study, both enzymatic and non-enzymatic  
403 antioxidants exhibited a significant decline at contaminated sites in comparison with the  
404 reference site. They revealed a balanced and coordinated work between each other in their  
405 biological role of detoxifying. For instance, SOD and CAT were positively correlated between  
406 each other. Indeed, SOD-CAT is the first line of defense against free radicals proliferation that  
407 causes damages to lipid membrane (Van der Oost *et al.*, 2003). They act synergistically, firstly  
408 SOD catalyzes superoxide ( $O_2^{\bullet-}$ ) to hydrogen peroxide ( $H_2O_2$ ), which is then transformed by  
409 CAT to  $H_2O$  and  $O_2$  (Vijayavel *et al.*, 2004). An excess of  $H_2O_2$  related to accumulation of  
410 reactive oxygen species (ROS), can inhibit SOD activity (Jodynis-Liebert *et al.*, 2005) and CAT  
411 activity can be likewise inhibited by an excess of superoxide radicals (Jodynis-Liebert *et al.*,  
412 2005). This was conspicuous in our data where negative correlation was found between the  
413 level of reactive species presented here as TBARS and the inhibition of SOD-CAT in  
414 contaminated sites. This inhibition of enzymes can be explained by the high level of  
415 contamination previously highlighted in the gulf of Gabes by toxic metals and polycyclic  
416 aromatic hydrocarbons inhibiting SOD and/or CAT activities (Atli and Canli, 2010; Romeo *et*  
417 *al.*, 2000; Vijayavel *et al.*, 2004). More interestingly, in some cases, contaminants can even  
418 inhibit the gene expression of antioxidants in chicks which can therefore promote reactive  
419 oxygen species propagation (Limón-Pacheco and Gonsebatt, 2009).

420 Similar to antioxidant enzymes pattern, vitamins A and E were positively correlated  
421 between each other. This was not surprising since both vitamins are lipophilic chain breaking  
422 antioxidants, concentrated in cellular membranes and acting in tandem to inhibit  
423 lipoperoxidation by scavenging ROS (Koivula and Eeva, 2010). Their decreased level suggests  
424 their intense mobilization as an antioxidantizing agent consequent to an abnormal production of  
425 free radicals as it is displayed by the level of lipid peroxidation. As a consequence of this  
426 depleted stock, chicks from Gabes gulf (Chatt-Essalem and Sfax sites) will have exacerbated  
427 lipid radicals spread, inducing further disturbances in coherence with other reports indicating  
428 repercussions of vitamins A and E deficiency (neurological abnormalities, abnormal  
429 erythrocyte membrane morphology, oxidative stress, altered immune system, malformations;  
430 Rolland, 2000; Clarke *et al.*, 2008). For instance, in the work of Fernie *et al* (2005) a lack of  
431 vitamin A in American kestrel was associated with oxidative stress and immunopathological  
432 effects.

433 Unlike all antioxidants measured in the present work, concentration of vitamin E was  
434 found higher in the heavily contaminated site of Chatt-Essalem than in Sfax and was not  
435 significantly different from the reference site. We suspect that this discrepancy may be due to



436 disparity in the exogenous intake of vitamin E. In fact, according to prey items found in nests,  
437 parents in Sfax salina and Monastir salina feed their chicks with several fish preys such as  
438 annular seabream, sardines and sand steenbras but in Chatt-Essalem, parents bring to their  
439 chicks mostly sardines (unpublished results) which are well known to be highly rich in  
440 polyunsaturated lipids and omega 3 fatty acids (Bandarra *et al.*, 1997). Consequently, we  
441 hypothesize that this richness involves an increase in serum content of vitamin E to prevent  
442 their oxidation. A similar pattern of increased vitamin E was observed in eggs of four  
443 piscivorous bird species consuming fish rich in polyunsaturated fatty acids (Surai *et al.*, 2001).

444 As a final point, biomarker responses of *Sterna hirundo* chicks integrated in multivariate  
445 analysis differed considerably between Gabes gulf sites and the reference site as we detected  
446 inter-site differences. Chicks were discriminated according to their physiological status in a  
447 way reflecting chemical pollution gradient and landscape use in each nesting area even if we  
448 cannot exclude that chicks from the three studied sites could have been exposed to other  
449 environmental stressors such as traffic noise, light and human presence. To monitor chicks and  
450 prevent their escape after hatching, nests were fenced individually with fishing nets in Chatt-  
451 Essalem site until we sampled chick blood. According to our observation, we did not notice that  
452 fences affected parents or chicks' behaviors. Nevertheless, we cannot exclude the high level of  
453 genotoxicity or oxidative stress could have been partially caused by a lack of freedom even if  
454 chicks had spaces, could hide and find shade. Regarding the high level of genotoxicity and  
455 oxidative stress also detected in chicks from Sfax, where chicks were not fenced, we concluded  
456 that fences we designed had a limited impact on chick physiological status. This field work  
457 pointed Monastir salina as a reference site as chicks had the "best health status", whilst in Sfax  
458 salina and Chatt-Essalem previously described as highly polluted sites, chicks had altered  
459 physiological status. Therefore, we consider that the level of genotoxicity and oxidative stress  
460 detected in chicks from these sites was related to local pollution exposure.

461  
462 The measured oxidative damages observed here are not restricted to lipids. Lipid peroxidation,  
463 as a process of chain reactions alter membrane structure, permeability and fluidity (Almroth *et*  
464 *al.*, 2005; Lushchak and Bagnyukova, 2006) but most of end products promoted by lipid  
465 deterioration have also genotoxic and mutagenic potential. Chicks from contaminated sites  
466 suffer from genotoxic effects triggered by free radicals or other genotoxicants such as PAH  
467 known to be highly reactive and own mutagenic and cancerogenic properties (Lushchak and  
468 Bagnyukova, 2006; Valavanidis *et al.*, 2006). Indeed, previous studies highlighted moderate to  
469 high hydrocarbons contamination of surface coastal waters in the gulf of Gabes compared to



470 other Mediterranean regions (PAH concentrations were 15.48-56.79 ng.L<sup>-1</sup> and 8.94-197.83  
471 ng.L<sup>-1</sup> in Gabes and Sfax coasts respectively; [Fourati et al., 2017](#)). The probability to detect  
472 micronucleated erythrocytes was more than three times lower in the reference site than in  
473 contaminated sites demonstrating that chicks were exposed to genotoxicants. This difference is  
474 similar to that found in purple heron nestlings from polluted and reference sites (three- to six-  
475 fold difference) in the Ebro basin in Spain and which exhibited also reduced blood antioxidant  
476 defenses ([Quirós et al., 2008](#)).

477 It is worthy to report that, in addition to micronuclei, chicks revealed other analogous of  
478 abnormal nuclear structures as shown in Figure 2. Although their frequency was not determined  
479 in this study, erythrocyte nuclear abnormalities (ENA) has already been successfully performed  
480 in genotoxic surveys with fish and amphibians but only more recently with birds ([Santos et al.,  
481 2017](#)). Therefore, the ENA assay could presumably be an additional tool for assessing  
482 genotoxic damage in *sterna hirundo*.

483 Thus, the overall physiological status of chicks from contaminated areas might expose them to  
484 further defects at different level of organization. Despite the high level of contamination around  
485 Chatt-Essalem, this site is still an attractive area for nesting for years which could be suitable  
486 in the short term but could lead to long term impacts on progeny survival and reproduction.

487  
488 In fish, the exposure of the brown bullhead *Ameiurus nebulosus* to carcinogens in contaminated  
489 hydrosystems induced age-selective mortality and population structure impairment (lack of the  
490 oldest age classes in contaminated sites; [Baumann et al., 1990](#)). Further delayed biological  
491 impacts, through trans-generational effects or early life-stage exposure, leading to decreases in  
492 juvenile survival and population growth rates, have also been highlighted in other fish species  
493 *Oncorhynchus gorboscha*, *Pimephales promelas* and *Chondrostoma nasus* ([Heintz et al., 2000](#);  
494 [Heintz, 2007](#); [White et al., 1999](#); [Santos et al., 2013a,b](#); [Devaux et al., 2011, 2015](#)). Hence,  
495 chicks may ultimately pay the outcome of oxidative and genotoxic damages observed in this  
496 study in terms of impaired growth, developmental abnormalities, reproduction or survival  
497 ([Koivula and Eeva, 2010](#)). As a first evidence of potential long term-effects, we outlined in the  
498 present work a significant correlation between an increase in physiological redox homeostasis  
499 disturbances and a decrease in chick mass three weeks post-hatching known to lead to a  
500 decrease in juvenile survival. Indeed, it is hypothesized that heavier individuals survive better  
501 than lighter ones because lightweight birds are more vulnerable to diseases, predation and  
502 parasites during their early life ([Van der Jeugd and Larsson, 1998](#)). Previous studies have found  
503 that offspring of higher body mass prior to or at fledging may have higher survival rates than

504 lighter ones (e.g Naef-Daenzer *et al.*, 2001 and Monrós *et al.*, 2002 for great tit; Monticelli and  
505 Ramos, 2012 for roseate tern; Mougín *et al.*, 2000 for Cory's Shearwater *Calonectris diomedea*;  
506 Gaston, 1997 for ancient Murrelet; Van der Jeugd and Larsson, 1998 for barnacle geese; Fear  
507 and Bristol, 2013 for sooty tern). In a four-year study on free-living great tits, a strong positive  
508 relationship between pre-fledging body mass and fledging success and recruitment probability  
509 was reported while pre-fledging resistance to oxidative stress significantly predicted fledging  
510 success (Losdat *et al.*, 2012). In another study, Noguera *et al.* (2011) demonstrated that chicks  
511 of European shags *Phalacrocorax aristotelis* suffering of elevated oxidative damage have  
512 decreased recruitment success probability in the next years. In spite of these evidences, there is  
513 still a dearth of studies that relate changes in demographic components such as juvenile survival  
514 and adult fertility to markers of oxidative stress and genotoxicity (Costantini *et al.*, 2015).

515

## 516 **5. Conclusion**

517 This work investigates ecotoxicological risks of contamination on populations of costal bird  
518 breeding in one of the main polluted areas in the Mediterranean region, the Gulf of Gabes. Our  
519 results highlighted that wild chicks from polluted sites suffer of genotoxicity and oxidative  
520 perturbations subsequent to excessive ROS generation and impoverished antioxidant defenses.  
521 The selected battery of biomarkers was effective to monitor biological impacts of contamination  
522 in *Sterna hirundo* in different locations discriminating chicks from sites under high chemical  
523 pressures. It is noteworthy to mention that such physiological impacts might not be costly only  
524 to exposed individuals but also to future generations. Physiological disruptions may potentially  
525 give rise to further disorders which could affect demographic traits and lead to long-term  
526 changes in population dynamics. As a first line of evidence, this work underlined a relationship  
527 between an increase of chick redox homeostasis disturbances and a decrease in body mass three  
528 weeks post-hatching, well-known to impact juvenile survival. Further analyses will be  
529 implemented to assess the relationship between chick redox status, chick body condition and  
530 the level of morphological abnormalities. Finally, this work highlighted that it is of major  
531 interest to incorporate more ecotoxicological approaches to population dynamic research  
532 programs and vice-versa to better investigate the impact of chemical pressures on aquatic  
533 wildlife.

534

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**Table 1.** Total surface and percentage of anthropogenic pressures in the surrounding 10 km of the three nesting sites of common tern in 2015. *Differences in total surface investigated is linked to variations in sea surface areas around investigated sites (see Figure S1).*

Site	Monastir	Sfax	Chatt-Essalem
Total surface (Km <sup>2</sup> )	228.43	170.23	146.77
Natural habitat and agricultural zone (%)	88.35	84.34	77.17
Urban zone (%)	11.39	13.16	18.59
Industrial zone (%)	0.26	2.5	4.24
	100	100	100
Distance from industrial zone (Km)	4.7	1.8	0

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766 **Table 2.** Early nests clutch size distribution of common tern among monitored areas in 2015

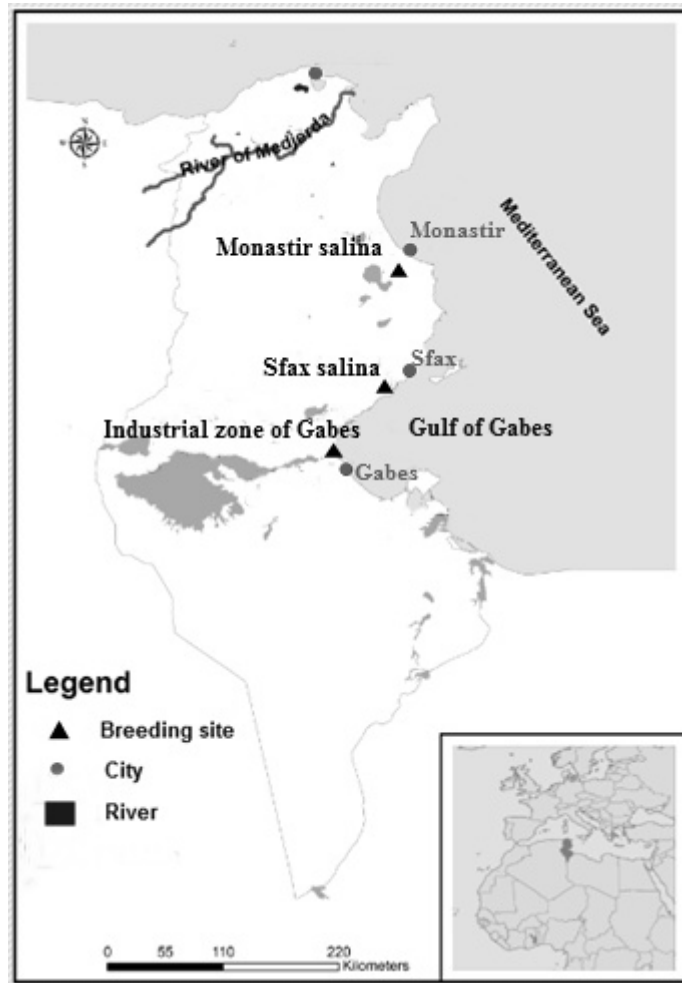
	Monastir salina	Sfax salina	Chatt-Essalem
1 egg	6	33	26
2 eggs	12	85	47
3 eggs	105	403	372
Total	123	521	445
767 Mean	2.80 ± 0.05	2.71 ± 0.03	2.78 ± 0.03

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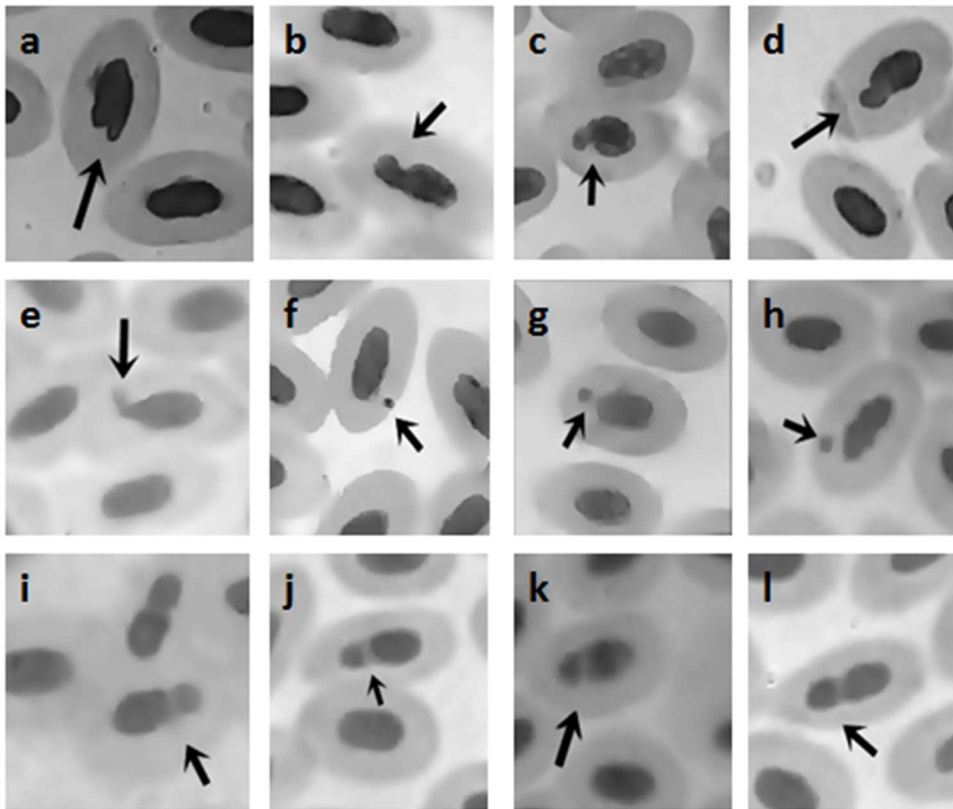
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**Figure 1.** Geographical location of selected sites in Tunisia for studying pollution effects on *Sterna*  
*hirundo* in 2015.

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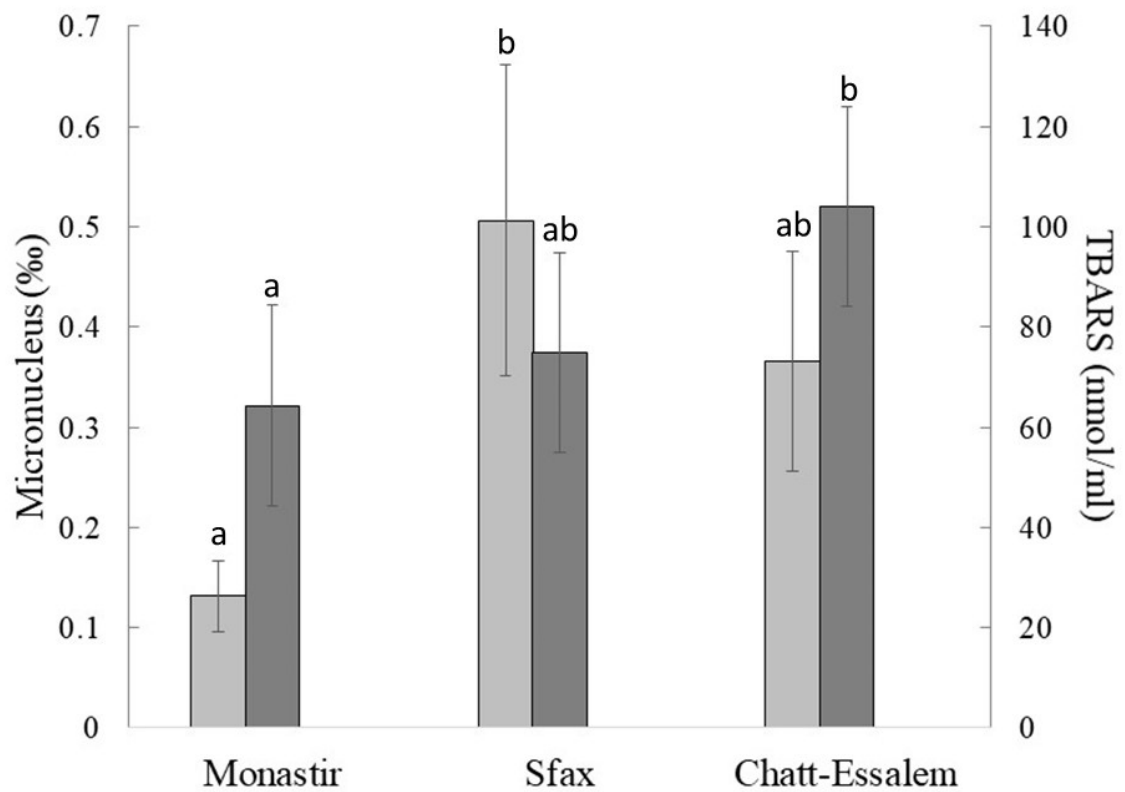
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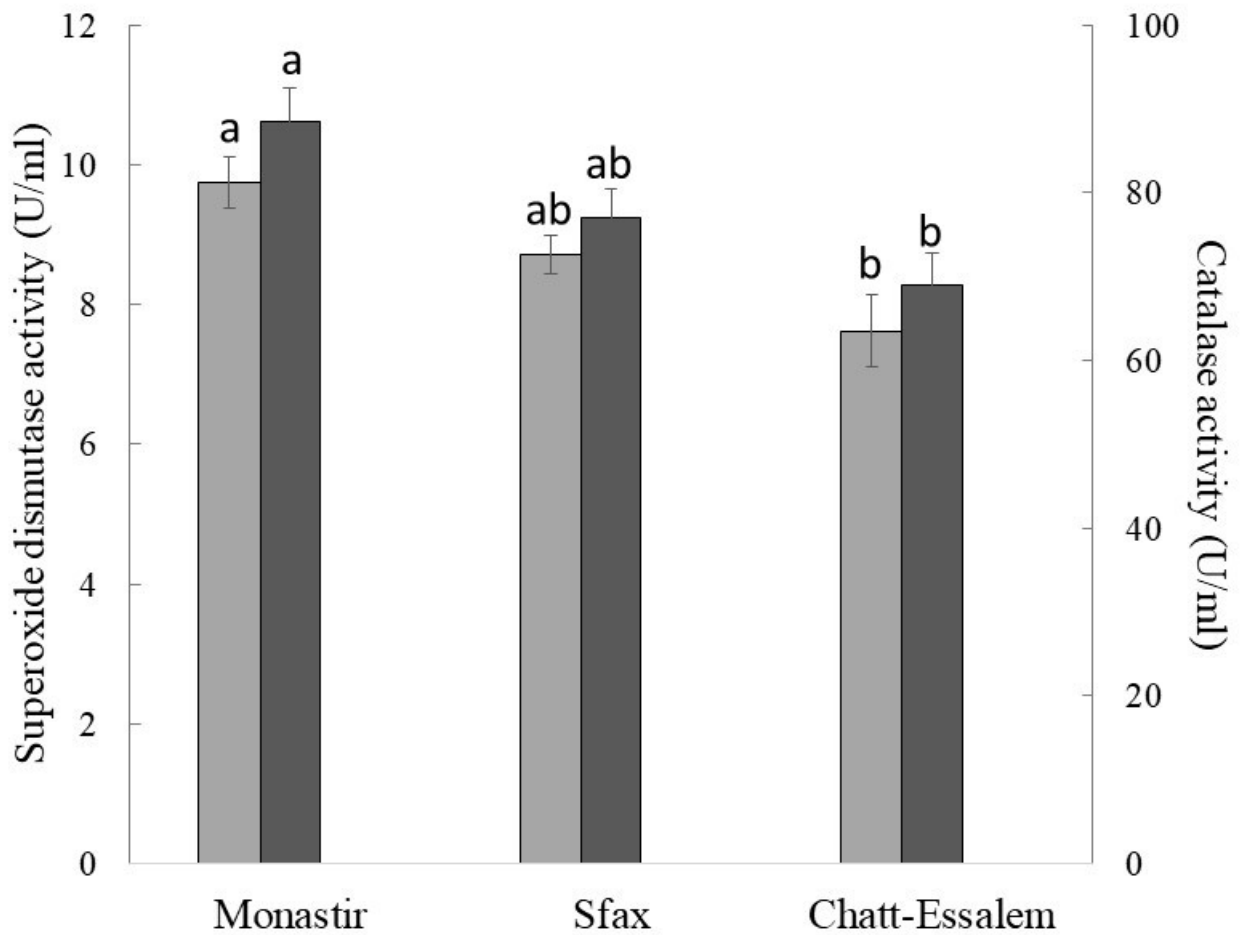
**Figure 2.** Erythrocyte nuclear abnormalities observed in *Sterna hirundo* chicks in 2015: a-b: notched nuclei, c-d: nuclear buds, e: nuclear tail, f-g: micronuclei, h: polymorphic nuclei+MN, i-l: segmented nuclei

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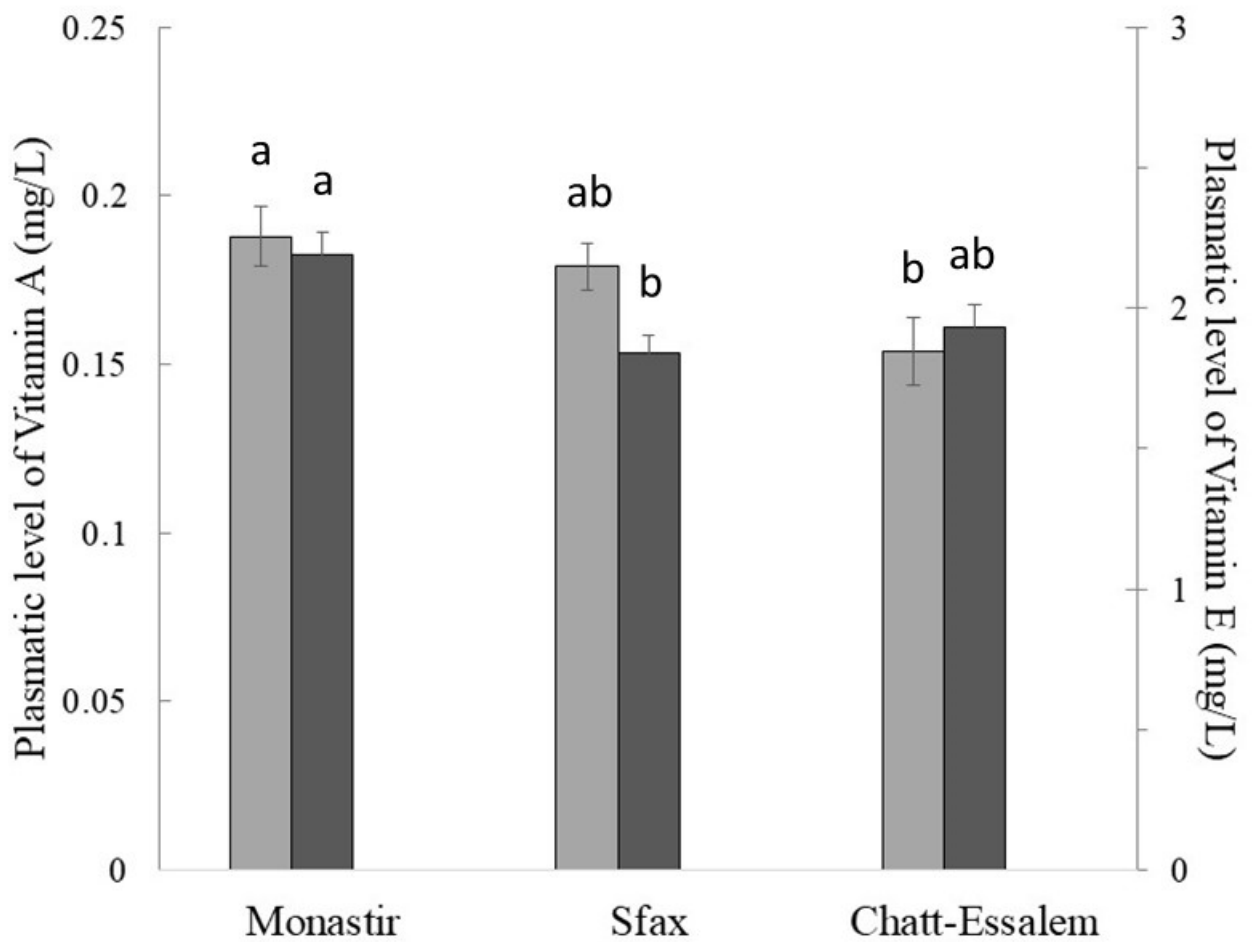
**Figure 3.** Micronucleus frequencies (%) and TBARS levels (nmol/ml) in blood of common tern chicks from the three nesting sites in 2015. Values are expressed as the mean ( $\pm$ SE). For each biomarker, different letters (a-b) denote significant differences of means ( $p < 0.05$ ) between sites according to pair-wise non-parametric tests.



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**Figure 4.** Superoxide Dismutase (SOD; U/ml) and Catalase (CAT; U/ml) levels in blood of common tern chicks from the three nesting sites in 2015. Values are expressed as the mean ( $\pm$ SE). For each biomarker, different letters (a-b) denote significant differences of means ( $p < 0.05$ ) between sites according to pair-wise non-parametric tests.

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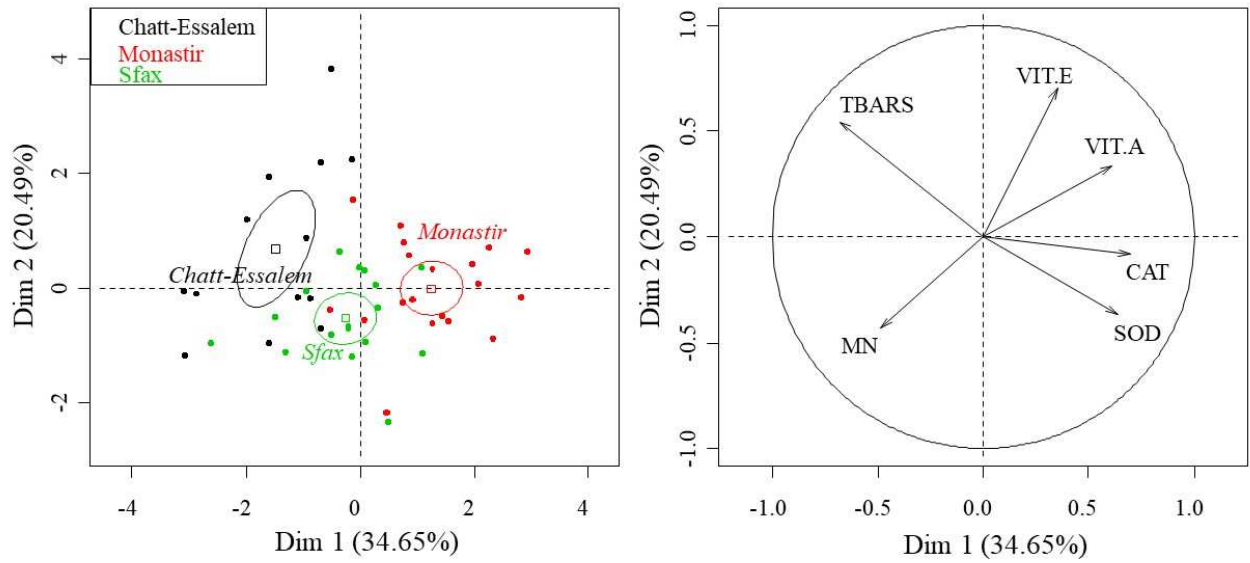
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**Figure 5.** Vitamin A (mg/L) and Vitamin E (mg/L) levels in blood of common tern chicks from the three nesting sites in 2015. Values are expressed as the mean ( $\pm$ SE). For each biomarker, different letters (a-b) denote significant differences of means ( $p < 0.05$ ) between sites according to pair-wise non-parametric tests.



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809 **Figure 6.** Principal component analysis performed on biomarker responses in *Sterna hirundo* blood

810 samples of chicks from the three nesting sites (Monastir, Sfax and Chatt-Essalem) in 2015.

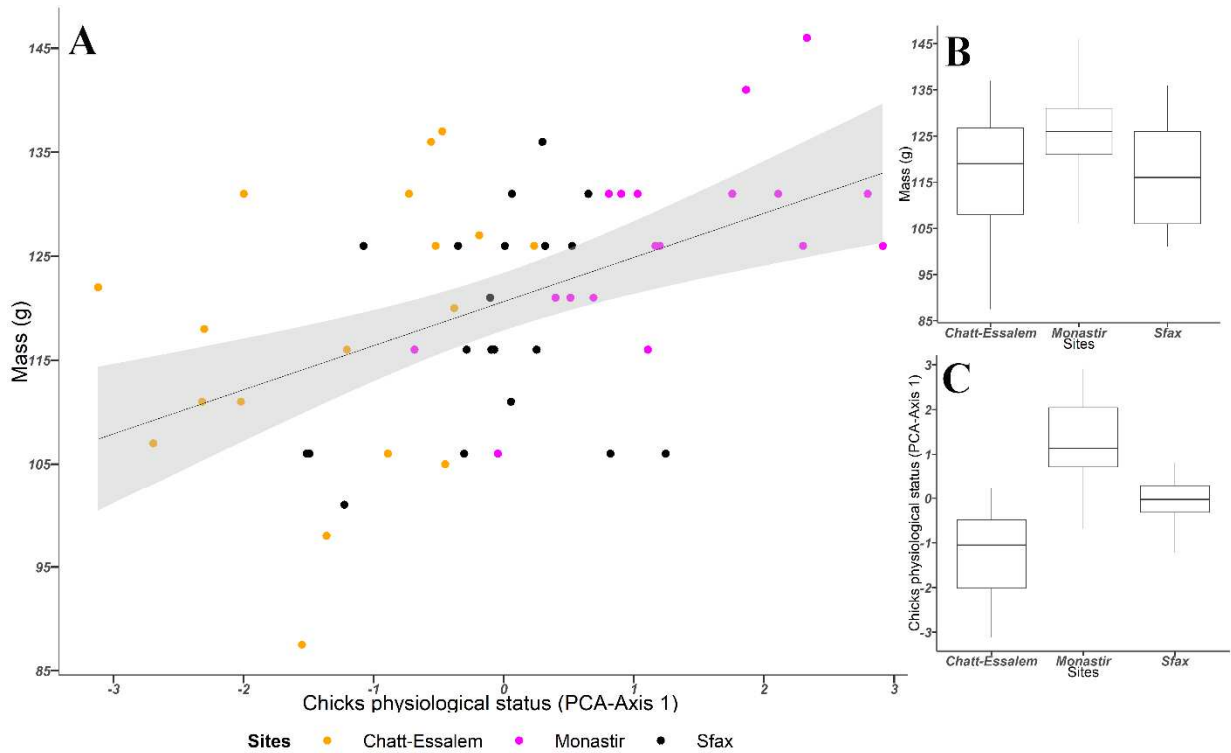
811 *CAT*: Catalase activity (U/ml); *MN*: number of micronucleus (‰); *SOD*: Superoxide dismutase

812 activity (U/ml); *TBARS*: level of lipid peroxidation in equivalent TBARS (nmol/ml); *Vit.A*: level of

813 plasmatic vitamin A (mg/L); *Vit.E*: level of plasmatic vitamin E (mg/L).

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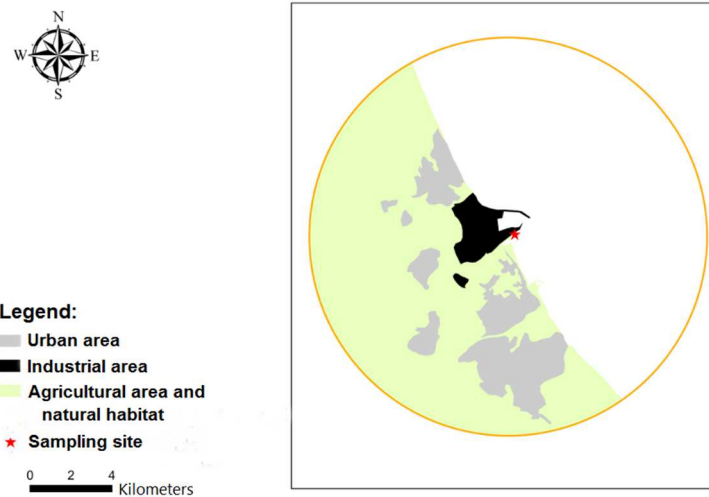
817 **Figure 7.** Relationship between chicks body mass three weeks post-hatching and their physiological  
 818 status (A) representing chick redox homeostasis disturbances and genotoxicity (PCA-Axis 1: highest  
 819 values represent chicks with a “good physiological status”). Solid line corresponds to the linear model  
 820 with 0.95 confidence intervals of this slope in grey ( $R^2 = 0.20$ ;  $p < 0.01$ ). Body mass (B) and  
 821 physiological status values (PCA-Axis 1) (C) of common tern chicks from the three nesting sites in  
 822 2015.

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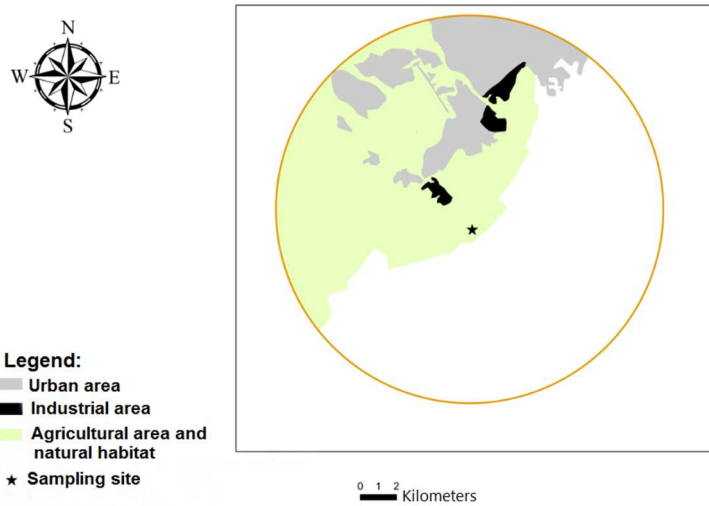
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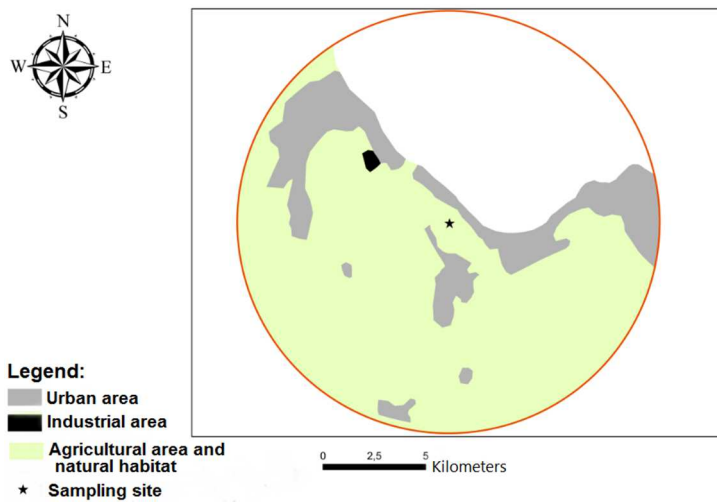
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831 **Supplementary material S1:** Landuse map in 10 Km around nesting sites (Chatt-Essalem, Sfax and

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Monastir) of common tern in Tunisia in 2015.

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