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1 ENVIRONMENTALLY RELEVANT MIXTURE OF PESTICIDES AFFECT MOBILITY AND DNA INTEGRITY  
2 OF EARLY LIFE STAGES OF RAINBOW TROUT (*ONCORHYNCHUS MYKISS*)

3

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## 12 Abstract

13 The aim of this study was to analyze the impact of three concentration of a pesticide mixture on the  
14 first development stages of rainbow trout (*Oncorhynchus mykiss*). The mixture was made up of three  
15 commonly used pesticides in viticulture: glyphosate (GLY), chlorpyrifos (CPF) and copper sulfate (Cu).  
16 Eyed stage embryos were exposed for 3 weeks to three concentrations of the pesticide mixture. Lethal  
17 and sub-lethal effects were assessed through a number of phenotypic and molecular endpoints  
18 including survival, hatching delay, hatching success, biometry, swimming activity, DNA damage (Comet  
19 assay), lipid peroxidation (TBARS), protein carbonyl content and gene expression. Ten target genes  
20 involved in antioxidant defenses, DNA repair, mitochondrial metabolism and apoptosis were analyzed  
21 using real-time RT-qPCR. No significant increase of mortality, half-hatch, growth defects, TBARS and  
22 protein carbonyl contents were observed whatever the pesticide mixture concentration. In contrast,  
23 DNA damage and swimming activity were significantly more elevated at the highest pesticide mixture  
24 concentration. Gene transcription were up-regulated for genes involved in detoxification (*gst* and  
25 *mt1*), DNA repair (*ogg1*), mitochondrial metabolism (*cox1* and *12S*), and cholinergic system (*ache*).  
26 This study highlighted both adaptive and deleterious molecular and behavioral responses of rainbow  
27 trout larvae following exposure to realistic mixture of pesticides.

28

29

30 **Keywords:** copper, glyphosate, chlorpyrifos, early life stages, rainbow trout, swimming behavior, DNA  
31 damage, development

## 32 1. Introduction

33 Increased concern around chemical contaminants in natural environments has led to a growth  
34 in research aimed at predicting the impacts of these contaminants on ecosystems.

35 Pesticides are the only kind of chemicals that are purposely released into the environment [1].  
36 Currently, more than 400 chemical compounds are used to treat crops against pests and weeds [2].  
37 Global pesticide usage is calculated at 4.6 million tons per year [1,3]. It has been estimated that 98 %  
38 of pesticides applied to agricultural crops do not reach their intended target, and could affect  
39 terrestrial and aquatic ecosystems. Studies have shown that 40 % of worldwide land mass poses a risk  
40 in terms of pesticide runoff into rivers and streams [4]. While most ecotoxicological studies focus on  
41 individual pesticides, several reports have also confirmed that pesticides are usually found in complex  
42 mixtures at low concentrations [2,5], and aquatic organisms are directly exposed to them [6–8].  
43 Exposure to pesticide mixtures may trigger additive effects when molecules have similar modes of  
44 action and affect the same molecular target, or by independent action when they have dissimilar  
45 modes of action affecting the same or different molecular targets, and therefore, synergism or  
46 antagonism may occur [9].

47 Because of its anti-microbial and anti-fungal properties, copper (Cu), also listed as a priority  
48 substance in the water Framework Directive (WFD), is widely used to protect vineyards from fungal  
49 diseases. Given its intensive use for both conventional and organic agriculture, Cu is a widely present  
50 pollutant in the environment, and transfer from soils to aquatic ecosystems is likely to occur [11,12].  
51 While it has an essential role in numerous cellular processes, excess Cu generates toxicity by  
52 producing reactive oxygen species (ROS) which may cause damages to lipids, proteins and nucleic  
53 acids, leading to cell death [13,14]. In a previous study, first stages of rainbow trout were found to be  
54 very sensitive to Cu [15]. Indeed, after a 3-week exposure to environmental concentrations of Cu (2  
55 and 20 µg/L), inhibitory effect on hatching and significant induction of malformations were observed.

56 In addition, several genes were down-regulated in Cu-exposed rainbow trout, especially those  
57 involved in detoxification (*gst*, *mt1* and *mt2*) and in cell cycle regulation (*p53*).

58 Glyphosate (GLY) is a broad-spectrum, non-selective and systemic herbicide used in numerous  
59 phytosanitary products. It acts as an inhibitor of the 5-enolpyruvyl-shikimate-3-phosphate synthase  
60 (EPSPS), an enzyme involved in the shikimic acid pathway that is common in all plants. Because of its  
61 extensive use, especially in genetically modified crops [16], high concentrations (ranging from 7.5 to  
62 700 µg/L) have been found in different streams and lakes near agricultural basins [17–19]. Several  
63 reports have highlighted the sub-lethal effects of GLY on fish in particular on DNA integrity [20–22],  
64 acetylcholinesterase (AChE) inhibition [23], swimming behavior alterations [24,25], and antioxidant  
65 enzyme activities [26,27]. We reported that following a 3-week embryo-larval exposure to glyphosate  
66 (0.1 and 1 mg/L), rainbow trout larvae exhibited reduced head size when compared to control larvae.  
67 Additionally, swimming behavior was also affected and exposed larvae had increased mobility  
68 compared to non-exposed larvae [28].

69 Chlorpyrifos (CPF) is an organophosphorus insecticide (OP) which has been widely used in the  
70 past by both industrial and private users. However, because of its high toxicity for humans and  
71 animals, the use of the product for domestic purposes has been restricted (US EPA), and there is a  
72 declining trend in its consumption [3]. Because it is on the list of priority substances for the WFD, its  
73 presence in surface and groundwater is closely monitored. While CPF has been detected  
74 predominantly in sediments from cultivated areas, low concentrations have also been observed in  
75 stream water, where maximum concentrations ranged between 0.06 and 0.45 µg/L in the USA and  
76 Argentina respectively [29,30]. As an OP, CPF affects the nervous system by inhibiting the AChE  
77 enzyme [31], disrupting the transmission of nerve impulses through synaptic terminals, impacting  
78 essential functions such as respiration and swimming behavior of fish [32,33]. In a recent study, low  
79 concentrations of CPF (0.3 and 3 µg/L) did not affect rainbow trout embryonic and larval viabilities.  
80 However, sub-lethal effects were observed on mobility of larvae which was reduced for those exposed

81 to 3 µg/L of CPF compared to control conditions. Low concentrations of CPF also down-regulated  
82 genes involved in steroid hormone pathways such as *er-b* and *cyp19a1* [34].

83           These three compounds (Cu, GLY and CPF), which have differing modes of action and  
84 functions, may be applied to crops simultaneously. Given the ease with which they are transported  
85 into water bodies through runoff, spray drift, or groundwater, it could be valuable to consider the  
86 toxicity of their combined effects, using realistic environmental concentrations, in aquatic organisms.  
87 To date, data on the effects of pesticide mixtures remain scarce, particularly relating to the early life  
88 stages of fish. To assess the sub-lethal effects of environmental concentrations of a pesticide mixture,  
89 a 3-week exposure was performed on rainbow trout embryos and several developmental and  
90 behavioral endpoints were recorded. The endpoints studied included survival, hatching delay,  
91 hatching success, morphological anomalies, swimming behavior, genotoxicity (measured by the comet  
92 assay), lipid peroxidation and protein carbonyl content, and gene transcription levels. The expression  
93 levels of ten genes were selected according to their biological functions in antioxidant defenses (*cat*,  
94 *sod*), detoxification (*mt1*, *gst*), mitochondrial metabolism (*cox1*, *12S*), cholinergic system (*ache*), DNA  
95 repair (*ogg1*, *rad*) or apoptosis (*bax*).

96

## 97 **Materials and methods**

### 98 **1.1. Test chemicals**

99 Chlorpyrifos-ethyl (CAS No. 2921-88-2) was purchased from ChemService (England). Copper sulfate  
100 ( $\text{CuSO}_4 \cdot \text{H}_2\text{O}$ , CAS No. 7758-99-8, 99.99 %) was purchased from Sigma Aldrich (Lyon, France).  
101 Glyphosate was purchase as a commercial formulation (available on the market) Roundup®GT Max.  
102 The active substance of Roundup®GT Max is 480 g/L of glyphosate acid, which is equivalent to 588 g/L  
103 of potassium salt of glyphosate.

104

### 105 **1.2. Exposure system**

106 Eyed-stage rainbow trout embryos (*Oncorhynchus mykiss*) were purchased from INRAE-PEIMA (Sizun,  
107 FR). Embryos were exposed for 3 weeks from stage 240 DD (Degree Days) to larvae stage 500 DD at  
108  $12 \pm 0.5$  °C. Stock solutions of copper (0.67, 2 and 20 mg/L) and glyphosate (33.3, 100 and 1000 mg/L)  
109 were prepared using osmosis water. Stock solutions of chlorpyrifos-ethyl (33.3, 100 and 1000 mg/L)  
110 were prepared using DMSO (dimethyl sulfoxide) as a solvent. Each condition (control-solvent, mixtures  
111 conditions) contained 0.0003 % DMSO.

112 Prior embryonic exposure, experimental apparatus units and tanks (1 L in polyethylene terephthalate)  
113 were saturated with chlorpyrifos diluted in distilled water for 2 weeks with the studied concentrations  
114 for each condition to ensure saturation of aquaria and avoid dramatic decrease of this compound  
115 during exposure. Test solutions were prepared by dilution of the stock solutions in a 5 L tank of spring  
116 water (dechlorinated) from Laqueuille (4.7 mg/L Ca, 1.8 mg/L Mg, 5.9 mg/L Na, 2.8 mg/L K, 40.3 mg/L  
117  $\text{HCO}_3^-$ , 0.2 mg/L  $\text{SO}_4^{2-}$ , 0.5 mg/L  $\text{NO}_3^-$ , 7.5 pH, <1.2 mg/L Cl<sup>-</sup>) oxygenated and were renewed every two  
118 days. The experimental conditions were design as follows: condition A (solvent control), condition B  
119 (0.1 µg/L of CPF + 0.67 µg/L of Cu + 33.3 µg/L of GLY), condition C (0.3 µg/L of CPF + 2 µg/L of Cu +  
120 100 µg/L of GLY) and condition D (3 µg/L of CPF, 20 µg/L of Cu + 1000 µg/L of GLY). Conditions C (3-

121 fold condition B) and D (10-fold condition C), which agree with the medium and highest  
122 concentrations in this study, correspond to the lowest and highest concentrations previously studied  
123 [15,28]. Each studied condition consisted of 3 replicates with 75 embryos in one L aquaria. A peristaltic  
124 pump (ISMATEC, ISM942) allowed maintaining a continuous flow rate (9 mL/min) of contaminated  
125 water from tanks into the incubation aquaria. Dissolved oxygen was measured each day with a fiber-  
126 optic mini-sensor Fibox 3 (PreSens Precision Sensor, Regensburg, Germany) and data was recorded  
127 with OxyView v6.02 software (PreSens Precision Sensor).

128

### 129 **1.3. Chemical analysis of pollutants in water**

130 Water samples were collected at  $T_0$  (at the moment of exposure),  $T_{24}$  and  $T_{48}$  (before water was  
131 renewed). Water samples were analyzed to determine Cu concentrations. Water sample of 40 mL for  
132 each condition were acidified with 5 % of nitric acid (Nitric acid 65 %, Fluka). Copper concentrations in  
133 water of condition “20  $\mu\text{g/L}$ ” and fish samples were analyzed by inductively Coupled Plasma Optic  
134 Emission Spectrometry (ICP-OES 720, Agilent Technologies), whereas copper concentrations in water  
135 of controls and “2  $\mu\text{g/L}$ ” conditions were analyzed using an atomic absorption spectrophotometer  
136 (Varian SpectrAA 240Z, Agilent Technologies, Santa Clara, USA). Detection limits (DL) for ICP-OES was  
137 2.26  $\mu\text{g/L Cu}^{2+}$  and for atomic absorption was 0.5  $\mu\text{g/L Cu}^{2+}$ . Glyphosate concentration and its main  
138 metabolite aminomethylphosphonic acid (AMPA) in water samples were analyzed using the method  
139 described by [35]. These samples were analyzed using an HPLC-ESI MS MS (Dionex Ultimate 3000,  
140 Thermo Fisher scientific – API 2000 ABSciex). Concentrations were determined with a calibration curve  
141 from 1 to 10  $\mu\text{g L}^{-1}$ . To ensure accuracy of each analysis, isotope-labeled surrogates was quantified,  
142 derivatization blanks were analyzed and quality controls were performed every 10 samples during  
143 analysis. Finally, data processing was performed with Analyst V1.6.2. CPF concentrations in water  
144 samples were measured by chromatographic methods using the Trace GC Ultra Gas Chromatograph  
145 (Thermo Fisher Scientific) equipped with an AS-3000 Autosampler (Thermo Fisher Scientific) and

146 coupled to a TSQ Quantum GC Triple Quadrupole (Thermo Fisher Scientific). The limit of quantification  
147 was 100 ng/L. The control of the device and the data processing were carried out by the XCalibur  
148 software.

149

#### 150 **1.4.Embryo-toxicity assay**

151 Embryonic and larval viability was recorded daily, and dead embryos were removed. Embryonic and  
152 larval viability denote the number of living individuals compared to the total number of embryos at  
153 the start of the experiment or total number of hatching larvae. Hatching rate is calculated by dividing  
154 the number of hatched embryos by the total number of embryos at the beginning of the experiment.  
155 Duration of development expressed in degree-days (DD) is the duration of embryonic development  
156 from fertilization to hatching. At the end of the experiment, 15 larvae were placed individually in Petri  
157 dish with ice water and few drops of carbonated water to sedate them. Photos of each larva were  
158 taken with stereomicroscope (MZ 7.5 Leica) coupled to a camera CCD (DFP420C Leica) and a cold light  
159 (Intralux® 4100, Volpi AG, Schlieren, Switzerland). From the photos, total body length, head length and  
160 yolk sac area were measured for each larva. Larvae were also observed for the presence of  
161 developmental anomalies including edemas, yolk-sac absorption, spinal malformations, craniofacial  
162 anomalies, presence of hemorrhages [15,36].

163

#### 164 **1.5.Swimming behavior analyses**

165 Swimming behavior analysis was carried out on 10 larvae per replicate at the end of the 3-week  
166 exposure. For 30 min, larvae were acclimated in the dark in 6-well microplates containing 3 mL of  
167 exposure solution at  $12.0 \pm 0.5^\circ\text{C}$ . After acclimation, the microplates were placed in the recording  
168 chamber (Daniovision Image Analysis System with Ethovision software version 12.0 Noldus) coupled to  
169 a thermoregulatory system set at  $12 \pm 0.5^\circ\text{C}$  (Pilot one®, Huber). The DanioVision recording chamber

170 includes a white light which can mimic a day/night cycle. For this, larvae were subjected to a light  
171 stress with a cycle duration of 30 min comprised of 10 minutes dark, 10 minutes light and 10 minutes  
172 dark. An infrared camera in the recording chamber recorded the motion of each larva where  
173 responses to light stimulation were analyzed. The Ethovision software recorded the average velocity  
174 every 30 seconds, average velocity, and total distance swam and time of mobility were determined for  
175 each light and dark phase.

176

### 177 **1.6.Comet assay with FPG enzyme**

178 Comet assay was performed on blood cells sampled by decapitation of 6 larvae per replicate using a  
179 heparinized pipette. Before decapitation, larvae were sedated using iced water and a few drops of  
180 carbonated water. Samples were immediately frozen in liquid nitrogen in microtubes with 200  $\mu$ L of  
181 cryoconservation solution (250 mM sucrose, 40 mM citrate trisodique, 5 % DMSO, pH adjusted to 7.6  
182 with nitric acid 1 M) until analysis. To improve the sensibility of the comet assay, slices were incubated  
183 with formamidopyrimidine glycosylase (Fpg) enzyme for 30 min at 37°C, as described by [37]. The  
184 comet assay was assessed following the protocol of [36] and [15]. After 20 min of DNA unwind in  
185 alkaline solution, electrophoresis was performed with a voltage of 25 V and 300 mA for 20 min. After  
186 nuclei separation, slides were stained with 20  $\mu$ g/mL of ethidium bromide solution, and comet lecture  
187 was carried out using an epifluorescence microscope (Olympus BX51) at x20 equipped with an  
188 Olympus U-RFL-T reflected fluorescence system lamp. 100 nuclei per slide were quantified using the  
189 Comet Assay IV software (Instrument Perspective Ltd). Results are expressed as a percentage of DNA  
190 tail.

191

### 192 **1.7.Biochemical analysis**

193 *Preparation of supernatant*

194 Pools of 2 larvae (approximately 250 mg) were homogenized in 250  $\mu$ L of phosphate buffer (0.1 M; pH  
195 7.5; 4°C) using an UltraTurrax® tissue homogenizer with a potter at 9000 g and 4°C. The supernatant  
196 S9 fraction was obtained after centrifugation at 9000 g for 25 min at 4°C. Each S9 fraction was split up  
197 in three tubes for total protein, TBARS and carbonyl protein analysis.

#### 198 *Total protein*

199 The protein content was determined according to the method described by [38] on the S9 fraction.  
200 Measurements were performed using bovine serum albumin as standard, and absorbance was  
201 recorded at 750 nm using a spectrophotometer microplate reader (Synergy HT, BioTek).

#### 202 *Lipid peroxidation (TBARS)*

203 Lipid peroxidation was performed as reported by [39], adjusted for a microplate reader. A volume of  
204 500  $\mu$ L of a solution containing 20 % of butylated hydroxytoluene (BHT) and 20 % of trichloroacetic  
205 acid (TCA) was added to 500  $\mu$ L of S9 fraction. The mixture was centrifuged at 9000 g for 10 min. Then,  
206 600  $\mu$ L of the supernatant was added to 120  $\mu$ L of HCl and 480  $\mu$ L of TRISbase (25mM)-TBA  
207 (thiobarbituric acid – 100mM) and heated at 80°C for 15 min. Afterwards, samples were cooled in iced  
208 water and mixed. The absorbance of the mixtures was measured using a UV-spectrophotometer  
209 (Synergy HT, BioTek) at 530 nm. Results were expressed as nmol of thiobarbituric acid reactive  
210 substance (TBARS) equivalents per mg of protein.

#### 211 *Protein carbonyl assay*

212 Protein carbonyl content was performed using the procedure of [40]. S9 fraction (500  $\mu$ L) was added  
213 to 50  $\mu$ L of a solution of streptomycin sulfate (11 %) - phosphate buffer (100 mM pH 7.4), mixed and  
214 incubated for 15 min at room temperature. Mixtures were centrifuged at 6000 g for 10 min and then  
215 split up into two tubes. The first one used as a control contained 200  $\mu$ L of supernatant and 800  $\mu$ L of  
216 HCl (2.5 M), and the other one used as a sample where 200  $\mu$ L of supernatant was added to 800  $\mu$ L of  
217 DNPH (2,4-dinitrophenylhydrazine 10 Mm), and then left incubated for 1 h. Proteins were precipitated

218 with 20 % TCA (trichloroacetic acid), and the formed pellets were washed 3-times by resuspension  
219 with 1 mL of ethanol-ethyl acetate (v:v). Pellets were solubilized with 500 µL of 6 M guanidine  
220 hydrochloride and centrifuged at 10000 g for 10 min. The measure of carbonyl content was performed  
221 with a UV-spectrophotometer (Biotek Synergy HT) at 370 nm. Results are expressed as nanomoles of  
222 DNPH incorporated/mg of protein, using the molar absorption coefficient of 22000 M<sup>-1</sup>cm<sup>-1</sup>.

223

## 224 **1.8. Gene expression**

225 At the end of the exposure, 6 larvae per replicate were sampled and kept individually in RNA later  
226 buffer (Qiagen). Samples were stored at - 80°C.

### 227 *RNA extraction*

228 Total RNAs were extracted in whole larvae using the kit “SV Total RNA Isolation system” (Promega)  
229 following the indications of the provider. Larvae were mixed and homogenized with the MP fastprep®-  
230 24 (Biorad, 6 m/s, 40 s) using ceramic beads (MP Biomedicals, Lysing Matrix D bulk).

### 231 *Retro-transcription of total RNA into cDNA*

232 The reverse transcription of total purified RNA was performed using the kit “GoScript Reverse  
233 Transcription System” (Promega). A mixture of 1 µL of oligo dT (1 µM), 1 µL of hexanucleotides (1 µM)  
234 and 10 µL of total purified RNA (1 µg) were mixed and heated for 5 min at 70°C followed by 5 min at  
235 4°C with a thermocycler (Eppendorf Mastercycler) to allow primer annealing. Afterwards, 1 µL of dNTP  
236 solution (10 mM), 4 µL of activity buffer, 1.5 µL of MgCl<sub>2</sub> (25 mM), 1 µL of reverse transcriptase (1  
237 U/µL) and 0.5 µL of RNAsine were added. Reverse transcription was then performed at 42°C for 1h.  
238 The cDNA samples were kept at -20°C until analysis by quantitative real-time PCR.

### 239 *Quantitative real-time PCR*

240 The studied primers were designed using the Primer3plus software (Table 1). Each primer-pair was  
 241 tested and showed an efficiency greater than 95%. Real-time qPCR was performed using GoTaq® qPCR  
 242 Master Mix kit (Promega). Each reaction mixture was made up of 1 µL of cDNA sample, 2 µL of specific  
 243 primer pair mix (200 µM each) and 17 µL of a mix consisted of Nuclease-Free Water and GoTaq® qPCR  
 244 Master containing SyberGreen fluorescent dye. Real-time PCR reactions were done in a Mx3000P®  
 245 qPCR system (Stratagene), and the amplification program was one cycle at 95°C for 10 min, then 45  
 246 amplification cycles at 95°C for 30 s, 60°C for 30 s and 72°C for 30 s. The specificity of the  
 247 amplifications was checked using the dissociation curve of the PCR products. The dissociation curve  
 248 was acquired by following the SYBR Green fluorescence level during a gradual heating of the PCR  
 249 products from 60 to 95°C.

250 For each gene, the cycle thresholds (Ct) were obtained from the software MxPro™ qPCR. Two  
 251 reference genes (*rpl7* and *ef1α*) were used, and level of gene transcription was normalized with the  
 252 mean of Ct value of reference genes according to the method of  $2^{-\Delta\Delta Ct}$  [41]. Induction (>2) or  
 253 repression (<0.5) factors were determined by the ratio of transcription levels of each condition and  
 254 the control. The list of studied genes and primer pairs are presented in Table 1.

255

256 *Table 1: Accession number and specific primer pairs for the Oncorhynchus mykiss used in our study*

Gene	Accession number	Primer (5' – 3')
<i>rpl7</i>	NM_001160672.2	GGTCGCTCTCACAGACAACA <sup>a</sup> TTATGTCCGTCCCTCTGGGT <sup>b</sup>
<i>cat</i>	FJ226382.1	CAGGTGTCTTTCTTGTTTCAG <sup>a</sup> GTCCAGGATGGGAAGTTGC <sup>b</sup>
<i>sod Cu/Zn</i>	NM_001124329.1	TGATTGGGGAGATCTCGGGT <sup>a</sup> CGGGTCCAGTGAGAGTCAAC <sup>b</sup>
<i>gst</i>	BT073173.1	ATTTTGGGACGGGCTGACA <sup>a</sup> CCTGGTGCTCTGCTCCAGT <sup>b</sup>
<i>cox1</i>	KP013084.1	TCGTTTGAGCCGTGCTAGTT <sup>a</sup> CTTCTGGGTGGCCGAAGAAT <sup>b</sup>

<i>12S</i>	KY798500.1	GCGCCAGCTTAAAACCCAAA <sup>a</sup> GCCCATTTCTTCCCACCTCA <sup>b</sup>
<i>ogg1</i>	XR_002474791.1	CTGATGGACAAGGCCAGTGT <sup>a</sup> GTAAGGACCCCATGGCTGTC <sup>b</sup>
<i>rad51</i>	XM_021612309.1	AGGCTGGAGGAGGACATCAT <sup>a</sup> GTATTTGAGGGTGGCAGCCT <sup>b</sup>
<i>bax</i>	BT074328.1	CAGAAAACCCAGGGAGGCAT <sup>a</sup> AGAACACATCCTGGGCACAG <sup>b</sup>
<i>mt1</i>	M18104.1	GTGGATCCTGCAAGTGCTCA <sup>a</sup> GTAATGCACCAGGCCTCACT <sup>b</sup>
<i>ache</i>	XM_021577686	AGGAGGGTTCTACAGCGGAT <sup>a</sup> TATCCTGGACCCACTGGAGG <sup>b</sup>

<sup>a</sup>Forward primer

<sup>b</sup>Reverse primer

## 257 1.9. Statistics

258 Each condition was carried out in 3 independent replicates. Results are presented as mean ± SD  
 259 (standard deviation). Normality of data distribution was verified by the Shapiro-Wilk test ( $p < 0.01$ ),  
 260 and the homogeneity of variances by the Levene test ( $p < 0.05$ ). When data followed a normal  
 261 distribution, a one-way ANOVA analysis was used ( $p < 0.05$ ) followed by the Tukey post-hoc test. If  
 262 normality was not met, the non-parametric test of Kruskal-Wallis ( $p < 0.05$ ) was used. All statistical  
 263 analysis was performed using R (<http://cran.r-projet.org/>).

264

## 265 2. Results

### 266 2.1. Condition of exposure

267 Concentrations of CPF (chlorpyrifos), GLY (glyphosate) and Cu were analyzed at each water change at  
 268  $T_0$ ,  $T_{24}$ , and  $T_{48}$  in the different treatments to estimate the losses of the compound (Table 2). To  
 269 analyze the complexes between Cu-GLY in solution, GLY was treated with a metal complex EDTA

270 (ethylene diamine tetra-acetic acid), and Cu was acidified with HNO<sub>3</sub>. In all conditions, the measured  
271 concentrations of CPF were inferior to the nominal concentrations at T<sub>0</sub>. For the condition B, all CPF  
272 concentrations were below the detection limit. For the two other conditions C and D, CPF  
273 concentration strongly declined after a few hours due likely to compound sorption to exposure unit. In  
274 the case of GLY, measure concentration was superior to the nominal concentrations at T<sub>0</sub>, and slight  
275 decreases were observed at T<sub>24</sub> and T<sub>48</sub>, but always higher than the nominal concentration. For Cu, the  
276 measured concentration was below the quantification limit for the B condition but was very close to  
277 the targeted concentrations for conditions C and D at T<sub>24</sub>. In contrast to the two other compounds, Cu  
278 concentration increased over time.

279

## 280 **2.2.Embryonic and larval survival**

281 Dissolved oxygen varied from 84 to 94.6 % throughout the embryonic and larval exposure. After 3-  
282 weeks exposure to the mixture of pesticides, no significant mortality was observed in embryos and  
283 larvae in any of the conditions studied (Table 3). Embryonic and larval survival reached more than  
284 95 % in all conditions. Hatchability was high, and more than 90 % of embryos succeeded to hatch. No  
285 significant differences between conditions were observed for the duration of embryonic development.

286

## 287 **2.3.Phenotypic effects**

288 Biometric measurements included total size, head size and yolk sac area. No significant alterations  
289 were observed on biometrics for larvae exposed to mixtures of CPF, Cu and GLY compared to non-  
290 exposed larvae (table 3). Similarly, no significant increases of developmental anomalies were noted at  
291 the end of the 3-weeks exposure on larvae. Percentages of abnormal larvae were  $15.0 \pm 2.8$ ,  
292  $23.7 \pm 3.6$ ,  $15.0 \pm 7.8$  and  $22.8 \pm 11.3$  % for control and conditions B, C and D, respectively (Table 3).

293 Table 2: Nominal and measured concentrations of chlorpyrifos (CPF), copper (Cu) and glyphosate (GLY) for each condition. Concentrations of samples were  
 294 analyzed at T0, T24 and T48.

	CPF ( $\mu\text{g/L}$ ) <sup>1</sup>		Cu <sup>2+</sup> ( $\mu\text{g/L}$ ) <sup>2</sup>		GLY ( $\mu\text{g/L}$ ) <sup>3</sup>				
	Nominal	Measured	Nominal	Measured	Nominal	Measured concentration			
	concentration	concentration	concentration	concentration	concentration				
<b>Condition B</b>	0.1	T <sub>0</sub>	0.67	Acidified (HNO <sub>3</sub> )	33.3	T <sub>0</sub>	No EDTA	EDTA	
		< 0.04		T <sub>0</sub>		< 1.1	T <sub>0</sub>	63.3	64.7
		T <sub>24</sub>		< 1.1		T <sub>24</sub>	67.5	49.2	
		T <sub>48</sub>	< 1.1	T <sub>48</sub>	55.4	42.2			
<b>Condition C</b>	0.3	T <sub>0</sub>	2.0	T <sub>0</sub>	100	T <sub>0</sub>	160.0	175.8	
		0.18		T <sub>0</sub>		2.03	T <sub>24</sub>	140.0	168.0
		T <sub>24</sub>		4.16		T <sub>24</sub>	154.0	137.25	
		T <sub>48</sub>	4.81	T <sub>48</sub>	154.0	137.25			
<b>Condition D</b>	3.0	T <sub>0</sub>	20.0	T <sub>0</sub>	1000	T <sub>0</sub>	1345.0	1785.0	
		1.63		T <sub>0</sub>		19.63	T <sub>24</sub>	1645.0	2022.5
		T <sub>24</sub>		26.32		T <sub>24</sub>	1775.0	1790.0	
		T <sub>48</sub>	26.65	T <sub>48</sub>	1775.0	1790.0			

295 <sup>1</sup>Limit of quantification: 0.13  $\mu\text{g/L}$ , limit of detection: 0.04  $\mu\text{g/L}$ .

296 <sup>2</sup>Limit of quantification: 1.1 µg/L, limit of detection: 0.1 µg/L.

297 <sup>3</sup>Limit of quantification: 30, 600 and 6000 ng/L

298

299 *Table 3: Viability and developmental anomalies of early life stages of rainbow trout exposed to a*  
 300 *mixture of pesticides. Values represent Mean  $\pm$  SD. No significant differences were observed, N = 3,*  
 301 *ANOVA.*

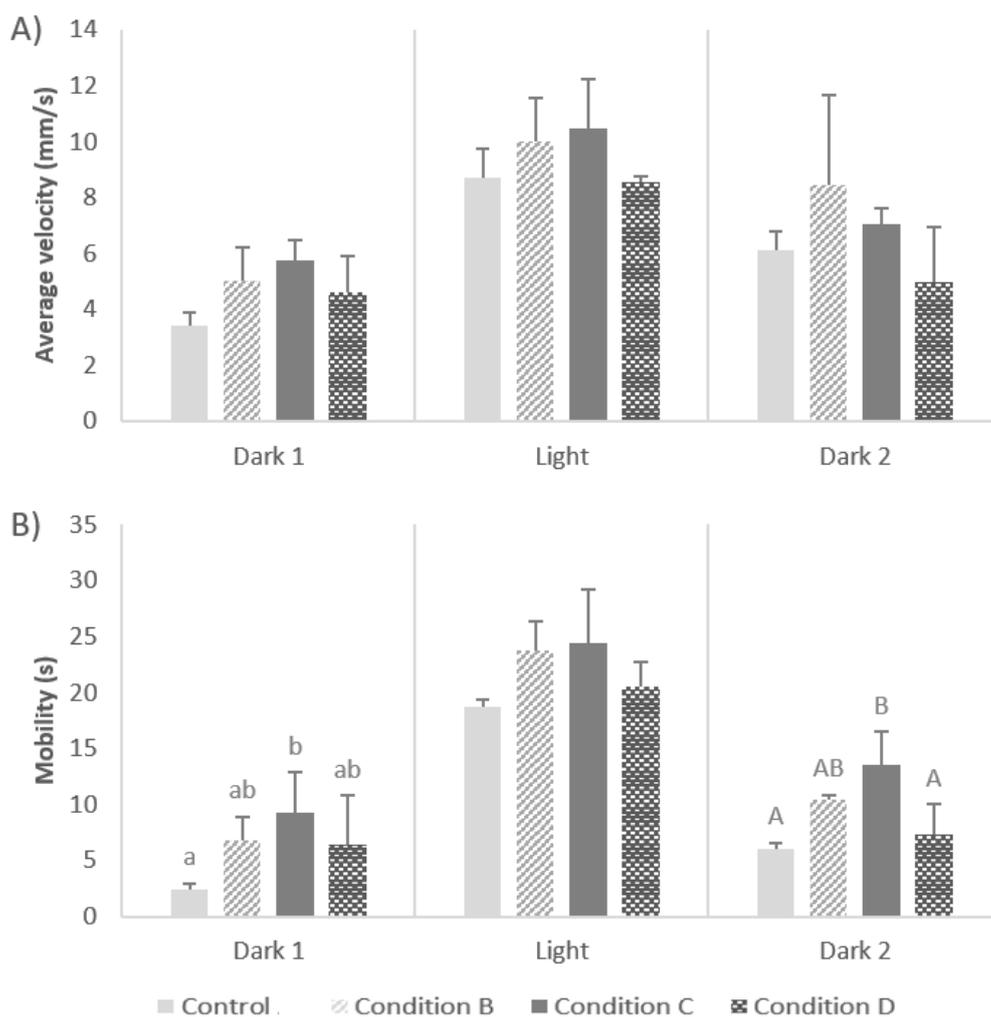
302

	Control (A)	Condition B	Condition C	Condition D
<b><i>Acute toxicity</i></b>				
Embryo viability (%)	98.7 $\pm$ 2.3	96.8 $\pm$ 3.4	96.4 $\pm$ 1.5	94.2 $\pm$ 4.1
Larval viability (%)	97.7 $\pm$ 2.1	94.5 $\pm$ 4.8	96.0 $\pm$ 3.2	95.2 $\pm$ 4.9
Cumulative viability (%)	96.4 $\pm$ 0.8	91.7 $\pm$ 7.3	92.6 $\pm$ 3.1	89.5 $\pm$ 3.5
Hatching rate (%)	95.6 $\pm$ 2.0	92.0 $\pm$ 4.6	92.0 $\pm$ 4.8	90.2 $\pm$ 4.1
<b><i>Sub-lethal toxicity</i></b>				
Duration of development (DD)	301.1 $\pm$ 2.2	294.9 $\pm$ 12.2	305.7 $\pm$ 1.1	309.9 $\pm$ 8.3
Total length (mm)	19.5 $\pm$ 0.3	19.6 $\pm$ 0.2	19.8 $\pm$ 0.2	19.1 $\pm$ 0.2
Head length (mm)	4.5 $\pm$ 0.0	4.5 $\pm$ 0.1	4.6 $\pm$ 0.0	4.3 $\pm$ 0.2
Ratio of head/body length (%)	23.2 $\pm$ 0.1	23.1 $\pm$ 0.2	23.1 $\pm$ 0.1	22.4 $\pm$ 0.7
Area of yolk sac (mm <sup>2</sup> )	10.8 $\pm$ 0.2	10.1 $\pm$ 0.4	10.4 $\pm$ 0.5	10.5 $\pm$ 0.1
<b>Abnormalities (%)</b>				
Total	15.0 $\pm$ 2.8	23.7 $\pm$ 3.6	15.0 $\pm$ 7.8	22.8 $\pm$ 11.3
Oedemas	7.5 $\pm$ 8.4	12.7 $\pm$ 6.2	9.3 $\pm$ 5.6	11.9 $\pm$ 6.5
Spinal	7.5 $\pm$ 8.4	16.6 $\pm$ 11.2	12.9 $\pm$ 11.3	16.5 $\pm$ 14.8
Craniofacial	1.8 $\pm$ 3.2	5.4 $\pm$ 5.3	11.3 $\pm$ 4.5	14.0 $\pm$ 4.5
Haemorrhages	7.6 $\pm$ 6.6	0.0 $\pm$ 0.0	9.3 $\pm$ 5.6	11.9 $\pm$ 6.5

303

304 **2.4.Swimming behavior**

305 Average velocity (mm/s) and mobility (s) of rainbow trout larvae subjected to light stress are  
 306 presented in Figure 1. Average velocity did not show any significant difference between conditions,  
 307 only a slight increase between conditions A and C ( $p$ -value = 0.052) was observed at the basal period  
 308 (dark 1, Figure 1-A). On the other hand, larvae exposed to condition C displayed a significant increase  
 309 in their mobility compared to control larvae ( $p < 0.05$ ) in both dark periods (dark 1 and 2, Figure 1-B).  
 310 In contrast, no significant differences were observed when larvae were subjected to light stress, due  
 311 to their high variability.

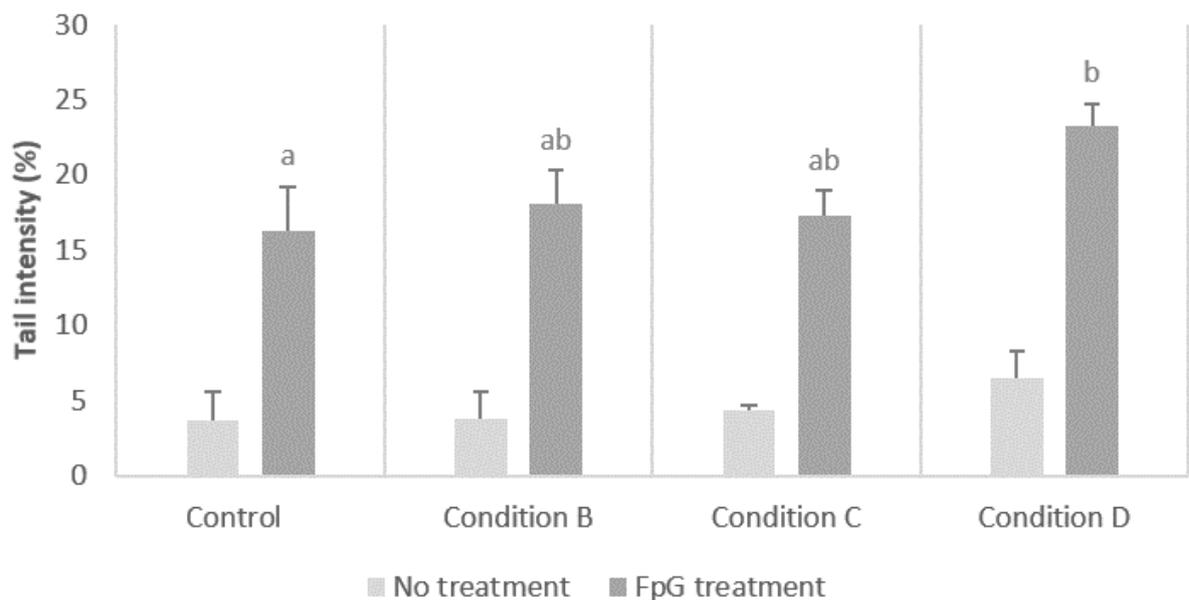


312  
 313 *Figure 1: Velocity average (mm/s) (A) and cumulative time of mobility (s) (B) of rainbow trout larvae*  
 314 *following exposure to mixture of pesticides. Different letters indicate significant differences between*  
 315 *conditions. Values represent Mean  $\pm$  SD (N = 3, ANOVA,  $p < 0.05$ ).*

316

## 317 2.5.DNA damage

318 Results of comet assay from blood cells are presented in Figure 2 with and without treatment by Fpg  
319 enzyme prior the alkaline unwinding of DNA. Fpg treatment was used to increase the sensitivity of the  
320 test. No significant differences in DNA damage were detected between conditions with the classical  
321 comet assay and tail DNA intensity ranged from 3.6 to 6.4 %. When nuclei were treated with Fpg  
322 enzyme, a 3 to 4 times increase of DNA damage was observed in comparison to results without Fpg  
323 treatment for the same exposure condition. In addition, DNA damage was significantly increased in  
324 larvae from control ( $16.2 \pm 3.0$  %) to condition D ( $23.3 \pm 1.5$  %).



325

326 *Figure 2: DNA damage in blood cells from rainbow trout larvae after exposure to mixtures of pesticides*  
327 *(CPF, Cu and GLY), with- and without treatment with the Fpg enzyme. Different letters indicate*  
328 *significant differences between conditions. Absence of letters indicates no significant differences (Mean*  
329  *$\pm$  SD, N = 3, ANOVA,  $p < 0.05$ ).*

330

331 **2.6.Lipid peroxidation and protein carbonyls**

332 A significant reduction in TBARS levels was observed on larvae from condition D when compared to  
333 control (Figure S1A). However, no significant alterations in protein carbonyl contents were observed in  
334 larvae from all conditions when compared to non-exposed larvae (Figure S1B).

335

336 **2.7.Gene expression**

337 Gene expression levels from whole larvae of rainbow trout revealed several up-regulated genes  
338 following a 3-weeks-exposure to pesticide mixture exposure (Table 4). Mixtures of GLY, Cu and CPF  
339 resulted in a significant induction of genes involved in detoxification (*gst* and *mt1*), DNA repair (*ogg1*),  
340 mitochondrial metabolism (*cox1* and *12S*) and cholinergic system (*ache*). High variability on gene  
341 transcription was observed for genes involved in oxidative stress (*cat*, *sod*) and no significant  
342 differences were observed. Gene involved in apoptosis (*bax*) was neither regulated. Strong genetic  
343 induction for 5 out of 10 genes was observed in larvae from condition D, indicating the likelihood that  
344 a higher concentration in the mixtures can lead to more severe toxic effects.

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357 Table 4: Transcription levels of whole-larvae of rainbow trout after 3 weeks of exposure to mixture of  
 358 CPF, Cu and GLY. Data was expressed as induction (above 2) or repression (below 0.5) factors  
 359 compared to the control condition. Asterisks refer to significant differences compared to control  
 360 condition (N = 3, ANOVA,  $p < 0.05$ ). /: identical to control.

	Condition B	Condition C	Condition D
<i>cat</i>	/	/	/
<i>sod</i>	/	/	/
<i>gst</i>	2.5*	/	/
<i>cox1</i>	5.3*	3.8*	23.8*
<i>12s</i>	4.9*	5.2*	32.4*
<i>ogg1</i>	4.4*	6.3*	34.8*
<i>rad51</i>	/	/	/
<i>bax</i>	/	/	/
<i>mt1</i>	9.4*	/	12.7*
<i>ache</i>	4.5*	7.0*	36.7*

361

362

### 363 3. Discussion

364 Numerous reports have shown that most aquatic ecosystems located close to farmland are  
365 contaminated with diverse kinds of pesticides, at different concentrations, which can have an impact  
366 on aquatic biota [6,9,42]. Commercial pesticide formulations including copper (Cu), chlorpyrifos (CPF)  
367 and glyphosate (GLY) are widely used for crop treatments, and are frequently detected in wetlands  
368 and streams, usually transported by runoff [11,17,19,43,44]. The interaction between these molecules  
369 can lead to changes in their overall toxicity as a function of the synergistic and/or antagonistic effects  
370 [5]. Each studied compound has a different mode of action, and its interaction may imply an induction  
371 or an inhibition of a specific metabolic pathway [5].

372 Our results show no significant mortality of embryos and larvae of rainbow trout exposed to  
373 low, moderate and highly concentrated mixtures of Cu, CPF and GLY. In previous studies performed  
374 with individual pesticides, no mortality was observed in early life stages of rainbow trout exposed to  
375 GLY [28] and CPF [34]. However, significant embryonic mortality (about 10 %) and a high frequency of  
376 half-hatched embryos (25 %) have previously been observed for embryos exposed to 20 µg/L of Cu  
377 [15]. In this study, the addition of CPF and GLY seems to inhibit the toxicity of Cu for embryos and  
378 larvae of rainbow trout. This result could be explained by possible interactions between compounds.  
379 Indeed, GLY is known to be a strong chelator of heavy metals such as Cu [45,46]. The functional groups  
380 of glyphosate (amine, carboxylate and phosphate) can react with metal ions to form complexes,  
381 resulting in a decreased bioavailability of Cu [47,48]. Our findings are consistent with similar studies  
382 that have been done with freshwater cladocera (*Ceriodaphnia dubia*) and earthworms (*Eisenia fetida*).  
383 Indeed, neonates of *C. dubia* were exposed to seven heavy metals, both alone and in binary mixtures  
384 with Roundup®, and most of the metals displayed less than additive toxicity [48]. In the same study,  
385 LC<sub>50</sub>-48h for *C. dubia* was estimated at 10 µg/L of Cu<sup>2+</sup>, but in the presence of GLY, mortality was  
386 reduced to 95 %. GLY could also affect the bioavailability of metals: for instance, it favors uptake of Hg  
387 and decreases uptake of Ag [48]. Zhou [47] tested the interactions between Cu and GLY on the toxicity

388 of earthworm (*Eisenia fetida*). Acute toxicity of Cu for *E. fetida* was calculated at 0.11 mg/L (LC<sub>50</sub>-48h),  
389 but when it was combined with GLY at concentrations of 0, 2 and 10 mg/L, worm mortalities declined  
390 from 57%, to 3% and 0 % respectively. The authors observed that the free Cu<sup>2+</sup> was reduced with the  
391 presence of GLY preventing the accumulation of Cu in worms.

392 For the swimming behavior of rainbow trout larvae exposed to single pesticides, we observed  
393 that GLY (100 µg/L) induced hyperactivity of larvae, increasing their velocity and mobility under light  
394 stimulation [28]. Similar observations were obtained by [24,49,50] in different fish species. On the  
395 other hand, larvae exposed to CPF (3 µg/L) were significantly less mobile than non-exposed larvae  
396 [34]. Comparable results were observed by [51] and [52] usually related to a decreased of  
397 acetylcholinesterase (AChE) activity [53–55]. In this study, rainbow trout larvae exposed to the mixture  
398 of pesticides at low and medium concentrations were significantly more mobile than control larvae.  
399 Mobility of larvae exposed to high concentrations of pesticide did not significantly differ from controls,  
400 and no differences were observed after light stimulation. Bonifacio [56] exposed adult females of ten  
401 spotted livebearer fish (*Cnesterodon decemmaculatus*) for 6 weeks at concentrations of CPF (0.1 to  
402 1 µg/L) and GLY (0.2 to 2 mg/L). They observed that fish exposed only to CPF had reduced swimming  
403 activity, but no differences were observed for fish exposed to GLY and binary mixtures. They  
404 hypothesized that GLY could decrease the effect of CPF on swimming activity. In another study, the  
405 addition of Cu also diminished the impact of CPF on swimming behavior of adult zebrafish (*Danio*  
406 *rerio*) exposed to a mixture of Cu (6.3 to 40 µg/L) and CPF (35 to 220 µg/L) for 24 h [57]. Indeed,  
407 decreased swimming activity was observed in zebrafish exposed to the highest concentration of CPF,  
408 suggesting that the addition of Cu may block the biochemical and neurological impacts of CPF or  
409 modify its uptake by neuron. Responses in terms of both hyperactivity and hypoactivity were observed  
410 on females of *Jenynsia multidentata* exposed for 24 and 96h to a binary mixture of CPF (0.4 and  
411 4 µg/L) and cypermethrin (0.04 and 0.4 µg/L) [53]. Low mixture concentrations significantly increased  
412 their swimming activity in the upper area of the aquaria, and higher mixture concentrations reduced  
413 their swimming activity and caused them to favor the bottom of the tank [53]. Alterations of normal

414 behavior may also depend on the ratio and concentrations of pollutants that are present in a mixture.  
415 For example, Kienle [51] exposed zebrafish embryos for 11 days to mixtures of CPF, at 0.25 and  
416 1 mg/L, and nickel chloride (NiCl<sub>2</sub>), at 7.5 and 15 mg/L. When fish were exposed to individual  
417 compounds, opposing behaviors were observed, i.e., CPF increased their swimming activity while  
418 nickel decreased it. Interestingly, when nickel was combined with low concentrations of CPF, the  
419 swimming activity of larvae was also decreased; but combined with higher concentrations of CPF,  
420 larvae increased their activity. Kienle [51] argued that both compounds have different modes of  
421 action, and in a mixture, they may act independently of each other. We can therefore suppose that  
422 the hyperactivity observed in our larvae may be an effect caused mostly by the dominance of GLY, in  
423 accordance with our previous results [28]. However, since there was an absence of significant  
424 response to light change for larvae exposed to mixtures of pesticides, we can assume that the  
425 presence of Cu and CPF antagonized or reduced the effects of GLY on larvae. Furthermore, numerous  
426 studies have documented the relationship between AChE activity and behavioral changes in fish  
427 [54,58,59]. In our work, we observed that *ache* gene expression was up-regulated in all conditions  
428 compared to control, suggesting that alterations observed on behavior of larvae could be related to  
429 this induction. Indeed, several pesticides, such as chlorpyrifos and glyphosate, have the capacity to  
430 inhibit the AChE activity [23,60]. This inhibition may cause acetylcholine accumulation in synapses of  
431 cholinergic neurons, leading to deficiency of important functions as swimming and feeding [61,62].

432         When embryos of rainbow trout were exposed separately to Cu, GLY and CPF, no significant  
433 increase in DNA damage on blood cells was observed after 3-week exposure compared to non-  
434 exposed larvae [15,28,34]. Blood cells from larvae exposed individually to GLY and CPF were also  
435 treated with the Fpg enzyme (formamidopyrimidine DNA glycosylase), but no significant changes were  
436 observed. Fpg is a DNA base excision repair enzyme that makes it possible to detect and remove  
437 lesions related to abasic sites (apuric or apyrimidic), alkylation and oxidative damage (8-oxoguanine)  
438 induced by Reactive Oxygen Species (ROS) [37]. In this study, the addition of Fpg enzyme reveals  
439 additional DNA damage, in particular for larvae exposed to the highest mixture concentration

440 (condition D). DNA damage have previously been detected by the comet assay in various aquatic  
441 organisms exposed individually to Cu, GLY and CPF using environmental concentrations [20,21,63,64].  
442 Few studies have examined the genotoxicity effects of mixtures of pesticides on fish. DNA damage was  
443 only observed in hemocytes of the freshwater clam *Corbicula fluminea* exposed to an herbicide  
444 mixture of Roundup® (2 and 10 mg/L) and atrazine (2 and 10 µg/L) for 96 h, but no genotoxic effects  
445 were observed when clams were exposed to the herbicides separately using the same concentrations  
446 [65]. In another study, a mixture of endosulfan and CPF (0.94 to 1.88 µg/L) also caused significant DNA  
447 damage to erythrocytes of fingerlings of Nile tilapia *Oreochromis niloticus*, after a 70-days exposure  
448 [66]. The genotoxicity of mixtures of metals has also been studied using the micronucleus assay on  
449 erythrocytes of fish, *Synodontis clarias* and *Tilapia nilotica*, exposed to Cu and zinc (Zn) [67]. In this  
450 case, micronuclei frequency was significantly increased when fish were exposed to binary mixtures,  
451 compared with exposure to individual metals. No significant protein carbonyl content was observed in  
452 exposed larvae when compared to control group. However, TBARS levels from larvae exposed to  
453 condition D were significantly lower than control ones. Exposure to individual compounds showed a  
454 significant reduction in TBARS levels in larvae exposed to 0.1 mg/L of GLY [28], while Cu and CPF alone  
455 did not modify TBARS [15,34]. As we hypothesized previously, the absence of increased TBARS levels  
456 and protein carbonyls may be the result of an efficient antioxidant system, serving to defend against  
457 oxidative stress as observed by [27,68,69] in piava (*Leporinus obtusidens*), freshwater catfish (*Channa*  
458 *punctatus*) and *Anguilla anguilla*, respectively.

459 With regard to gene expression levels, 3-week exposure to all conditions was mainly  
460 associated with the up-regulation in the genes investigated, with similar patterns observed in each of  
461 the studied conditions. The exposure to the mixture of the three compounds led to the up-regulation  
462 of genes involved notably in detoxification, mitochondrial metabolism and DNA repair. Cu, GLY and  
463 CPF are well known to produce individually reactive oxygen species (ROS), and the enzymes catalase  
464 (CAT) and superoxide dismutases (SOD) protect the cells from oxidative damage caused by ROS. In our  
465 study, no significant regulation of *cat* and cytoplasmic *sodCu/Zn* genes were observed on exposed

466 larvae because of their elevated variability. However, many studies showed evidence of the capability  
467 of these compounds to induce antioxidant gene expression [62,70–72]. ROS neutralization could also  
468 be completed by mitochondrial *sodMn* but also *gpx* (Glutathione peroxidase) via glutathione  
469 oxidation, but these two genes were not investigated in our study. Genes involved in detoxification,  
470 *gst* and *mt1*, were induced, especially in larvae exposed to the lowest and the highest mixture  
471 concentrations. Expression of *gst* (glutathione s-transferase) is usually up-regulated when fish are  
472 exposed to xenobiotics. In our study, *gst* expression was significantly increased in larvae exposed to  
473 the lowest mixture concentration (condition B), and this could indicate that larvae were able to  
474 implement defense mechanisms against low concentrations of pesticides. GST proteins play an  
475 important role in detoxifying xenobiotic compounds by catalyzing the conjugation of reduced  
476 glutathione (GSH) on primary metabolites from phase I metabolism [73]. *Mt1* gene encodes the  
477 metallothionein proteins, usually induced by metals exposure such as copper, but it has also an  
478 antioxidant role [74]. Metallothionein proteins have the capacity to bind xenobiotic heavy metals  
479 through their thiol groups, providing protection against metal toxicity and oxidative stress [74]. In a  
480 previous study using Cu [15], rainbow trout larvae showed a down-regulation of *mt1*, *mt2* and *gst*  
481 genes after 3-weeks exposure. However, no significant changes of *gst* gene expression were observed  
482 on larvae exposed individually to GLY and CPF [28,34]. Therefore, we can presume that the induction  
483 of these genes is the consequence of an additive effect over these functions on larvae exposed to the  
484 mixture of pesticides. In our study, a significant induction of *cox1* (cytochrome c-oxidase subunit 1)  
485 and *12S* gene expression was observed in all conditions compared to control larvae. These up-  
486 regulations were even stronger in larvae exposed to the highest mixture concentrations (condition D).  
487 An induction of these genes could reveal an increased mitochondrial number per cell and a possible  
488 mitochondrial electron transport chain disruption from the lowest mixture concentration tested. The  
489 mitochondria is involved in the production of energy for cellular metabolism by synthesizing ATP  
490 (adenosine triphosphate). Since the mitochondria is a major source of ROS (superoxide and hydrogen  
491 peroxide) through the disruption of the electron-transport chain [75], the up-regulation of *cox1* could

492 denote an increased demand for energy to fight the effects of toxicants. Furthermore, the putative  
493 increase of mitochondria could also stimulate ROS production [75,76]. The observed induction of *cox1*  
494 and 12S genes could mean a high-energy demand, in the form of ATP, to defend cells against ROS  
495 production or to repair cellular damage of pesticides. This demand of energy may be even required for  
496 functions that we have not considered in this study. We also observed, in all conditions, an induction  
497 of DNA repair gene expression, *ogg1*, which is responsible for the removal of 8-oxoguanine as a result  
498 of ROS production. The over-expression of *ogg1* (X34.8) on larvae exposed to the highest mixture  
499 concentration is probably related to the significant induction of DNA damage that we observed on  
500 blood cells after treatment with Fpg-enzyme. Consequently, DNA lesions observed in blood cells were  
501 likely related to oxidative damage. The up-regulation of DNA repair (*ogg1*) and detoxification (*gst* and  
502 *mt1*) gene expression may indicate that defense mechanisms were effectively implemented to avoid  
503 the effects of ROS production in cells and DNA. This could relate to high demand for energy, with up-  
504 regulation observed for 12S and *cox1* gene. However, to validate this relationship, future research  
505 could focus on the expression of mitochondrial superoxide dismutase gene (*sodMn*), which transforms  
506 the superoxide anion into hydrogen peroxide, and the expression of glutathione peroxidase (*gpx*),  
507 which detoxifies the hydrogen peroxide into water.

508           It was not clear whether the observed gene regulation in larvae was a consequence of a toxic  
509 effect or an adaptation effect. However, since no significant toxic effects were observed on exposed  
510 larvae (e.g. mortality, lipid/protein oxidation), we considered that the significant gene up-regulation  
511 was mostly an adaptive effect triggered to deal with toxic exposure. At higher mixture concentrations,  
512 energy demand increased and the DNA repair, *mt1* and *ache* gene expression greatly increase. This  
513 indicates a defense mechanisms induction to prevent toxic effects resulting from increased exposure  
514 to chemicals. When comparing the pattern of gene expression from the present work with previous  
515 work done with individual exposure of Cu, GLY and CPF [15,28,34], we could consider that the joint  
516 action of the three pollutants had an additive or a synergistic effects since the observed effects of the  
517 mixture were stronger than the isolated substances.

518

#### 519 **4. Conclusions**

520 Sub-chronic exposure to a mixture of copper, glyphosate and chlorpyrifos on rainbow trout embryos  
521 can induce hyperactivity and DNA damage at high concentrations. In addition, several genes were up-  
522 regulated, especially those involved in detoxification, mitochondrial metabolism, cholinergic system  
523 and DNA repair. The up-regulation of *gst*, *ogg1*, *mt1*, *cox1* and 12S gene expression suggests that  
524 exposure to the mixture of Cu, GLY and CPF promotes cellular defense mechanisms, indicating an  
525 adaptive effect which could limit the occurrence of more severe effects. Our results give new  
526 information about the spectrum of effects and the mechanisms involved in cellular response of  
527 rainbow trout embryos exposed to environmentally relevant mixtures of pesticides.

528

529

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