

2 **Linking *in vitro* effects and detected organic micropollutants in surface**
3 **water using mixture toxicity modeling**

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27 **Abstract**

28 Surface water can contain countless organic micropollutants, and targeted chemical analysis alone
29 may only detect a small fraction of the chemicals present. Consequently, bioanalytical tools can be
30 applied complementary to chemical analysis to detect the effect of complex chemical mixtures. In
31 this study, bioassays indicative of activation of the aryl hydrocarbon receptor (AhR), activation of
32 the pregnane X receptor (PXR), activation of the estrogen receptor (ER), adaptive stress responses
33 to oxidative stress (Nrf2), genotoxicity (p53) and inflammation (NF- κ B) and fish embryo toxicity
34 were applied along with chemical analysis to water extracts from the Danube River. Mixture
35 toxicity modeling was applied to determine the contribution of detected chemicals to the biological
36 effect. Effect concentrations for between 0 to 13 detected chemicals could be found in the literature
37 for the different bioassays. Detected chemicals explained less than 0.2% of the biological effect in
38 the PXR activation, adaptive stress response and fish embryo toxicity assays, while five chemicals
39 explained up to 80% of ER activation and three chemicals explained up to 71% of AhR activation.
40 This study highlights the importance of fingerprinting the effects of detected chemicals.

41

42 **Keywords:** bioassay; chemical analysis; bioanalytical equivalent concentration; mixture effect
43 modeling; surface water, concentration addition

44 **Introduction**

45 Human-impacted rivers can contain a complex mixture of micropollutants, such as pharmaceuticals,
46 pesticides and industrial compounds, as well as their transformation products.^{1,2} The sources of
47 these contaminants can include both point sources, such as wastewater effluent discharge, and
48 diffuse sources, such as run-off from urban and agricultural areas.³ Given the diversity of
49 micropollutants in water, targeted chemical analysis alone is insufficient to detect all chemicals
50 present in the aquatic environment. Bioanalytical tools can be applied complementary to chemical
51 analysis as they can provide information about the biological effect of chemicals present in a
52 sample and reveal the presence of active compounds not detected by targeted analysis.

53

54 *In vitro* bioassays based on various cellular response pathways, including induction of xenobiotic
55 metabolism, receptor-mediated effects, adaptive stress responses and cytotoxicity, have been
56 applied to detect the presence of micropollutants in water samples.⁴⁻⁶ While activation of these
57 endpoints does not necessarily translate into higher level effects, biological response at the cellular
58 level is a key step in the adverse outcome pathway.⁷ Further, bioassays indicative of xenobiotic
59 metabolism and repair and defense mechanisms can be applied as sensitive tools to detect the
60 presence of micropollutants, as effects in these endpoints often occur at lower concentrations than
61 those causing cell death or damage.

62

63 Bioanalytical tools also have the advantage that they can take into account mixture effects among
64 chemicals, rather than focusing on individual chemicals. The mixture effects that occur among
65 chemicals can be categorized as concentration addition or independent action for chemicals acting
66 according to the same or a different mode of action, respectively, both of which assume no
67 interaction among the mixture components, or synergism or antagonism, where the mixture
68 components can interact.⁸ For environmental samples, such as surface water, containing many
69 chemicals at low concentrations, synergism is rare and instead concentration addition has been

70 suggested as a conservative approach to evaluate mixture toxicity of multicomponent mixtures not
71 only for receptor-mediated effects,⁹ but also adaptive stress responses¹⁰ and cytotoxicity.¹¹ Mixtures
72 that act in a concentration additive manner can be described using the bioanalytical equivalent
73 concentration (BEQ) concept, which represents the concentration of a reference compound that
74 elicits an equivalent response in a particular assay as the sample and can be determined from both
75 bioassays and chemical analysis. By comparing the BEQ from bioanalysis (BEQ_{bio}) and BEQ from
76 chemical analysis (BEQ_{chem}) it is possible to determine the contribution of detected chemicals to the
77 biological effect.¹² This approach has been applied to a wide range of water types including surface
78 water,^{13,14} wastewater,^{6,14,15} recycled water¹⁶ and swimming pool water.¹⁷

79

80 In this study, a suite of bioanalytical tools was applied to water samples from the human-impacted
81 Danube River. The BEQ concept was utilized as a simple mixture effect prediction model to
82 determine the contribution of detected chemicals to the biological effect. The battery of bioassays
83 follows previous recommendations on the selection of sensitive indicator bioassays that cover
84 endpoints related to different stages of cellular toxicity pathways including induction of xenobiotic
85 metabolism and receptor-mediated effects representing important molecular initiating events, as
86 well as adaptive stress responses and cytotoxicity or other apical endpoints.⁴ Bioassays indicative of
87 specific modes of action, such as estrogenic activity, have previously shown that a small number of
88 chemicals often explain a high proportion of the biological effect in wastewater,^{14,16,18} but less is
89 known about the explanatory power of known chemicals in other endpoints.

90

91 Activation of the ligand-dependent transcription factor aryl hydrocarbon receptor (AhR) was
92 assessed using the CAFLUX assay.¹⁹ While most applications focus on dioxin-like compounds,
93 which are unlikely to be found in the water phase due to their hydrophobicity, around 16% of the
94 320 environmental compounds examined by Martin et al.²⁰ were found to induce AhR-dependent
95 gene expression. Activation of the pregnane X receptor (PXR), which is an important factor in

96 xenobiotic metabolism regulation, was assessed using the HG5LN-hPXR assay and 73% of
97 chemicals studied in Martin et al.²⁰ activated PXR. Activation of the estrogen receptor (ER) was
98 assessed using the reporter gene MELN assay. A suite of bioassays indicative of adaptive stress
99 responses reacting to oxidative stress (*ARE-bla*), genotoxicity (*p53RE-bla*) and inflammation (NF-
100 κ B-*bla*) were also included. Adaptive stress response pathways are activated to restore the cell to
101 homeostasis after damage.²¹ The oxidative stress response is mediated by Nrf2 and the antioxidant
102 response element,²² and 26% of the 1859 chemicals in the US EPA ToxCast database were active in
103 the *ARE-bla* assay.²³ The p53 response is activated after DNA damage, leading to either repair or
104 apoptosis,²⁴ and activation of p53 can indicate the presence of genotoxic carcinogens.²⁵
105 Approximately 15% of chemicals in the ToxCast database were active in the *p53RE-bla* assay.²³
106 The NF- κ B pathway is an important driver of the inflammatory response and can target cytochrome
107 P450s, cytokines and apoptosis regulators,²¹ and 3% of chemicals in the ToxCast database induced
108 a response in the NF- κ B-*bla* assay.²³ Finally, the fish embryo toxicity (FET) test using zebrafish
109 was applied complementary to the cell-based bioassays as it can provide information about the
110 organism-level response. Apical endpoints in this test include embryo coagulation and lack of heart-
111 beat²⁶ and a recently published database containing 641 chemicals showed that 74% of reviewed
112 chemicals caused mortality in the FET test.²⁷

113

114 The current study aimed to assess what fraction of the biological effects of the cellular toxicity
115 pathway can be explained by the quantified chemicals with available effect data. We used samples
116 from a large water body with high dilutions and low levels of micropollutants stemming from very
117 diverse sources to test the hypothesis that in these water types there is typically not a dominant
118 chemical or chemical group but instead the effects are largely driven by the mixture effects of many
119 chemicals. Large volume solid phase extraction (LVSPE) water extracts from the Danube River
120 were analyzed in the bioassays introduced above to determine BEQ_{bio}. The effect analysis was
121 complemented with targeted chemical analysis of 272 water relevant chemicals, including

122 pesticides, pharmaceuticals, artificial sweeteners, steroidal hormones and industrial compounds.
123 The target list is by no means comprehensive, but is based on previous targeted and non-targeted
124 analysis of the Danube River.²⁸ Effect concentrations for the individual detected chemicals were
125 collected from the literature to determine BEQ_{chem} . By comparing BEQ_{bio} and BEQ_{chem} it was
126 possible to determine the extent to which the detected chemicals contributed to the mixture effect in
127 each bioassay.

128

129 **Experimental**

130 *Sampling*

131 Sampling occurred during the 3rd Joint Danube Survey (JDS3)²⁹ between August and September
132 2013 using LVSPE (Maxx GmbH, Rangendingen, Germany).³⁰ The sampling locations, which
133 included both the Danube River and its tributaries, are shown in Table S1 and Figure S1 of the
134 Supporting Information (SI), along with detailed information about sample enrichment and
135 extraction. Briefly, up to 500 L of water was passed through a stainless steel chamber containing
136 neutral sorbent Chromabond® HR-X, anionic exchanger Chromabond® HR-XAW and cationic
137 exchanger Chromabond® HR-XCW (Macherey-Nagel, Dueren, Germany). After extraction, each
138 solid phase was freeze-dried, then extracted with solvents and the eluates were combined. The
139 sample aliquots were reduced to dryness via rotary and nitrogen evaporation prior to shipping, then
140 re-suspended in either DMSO or methanol, depending on the assay.

141

142 *Chemical Analysis*

143 Target screening analysis of the JDS sample extracts for 264 chemicals was performed by liquid
144 chromatography-high resolution mass spectrometry (LC-HRMS) using an Agilent 1200 LC coupled
145 to a Thermo LTQ Orbitrap XL. Analysis was run in both positive and negative mode electrospray
146 ionization. For further details see Hug et al.³¹ Eight steroidal hormones and industrial phenolic
147 compounds were analyzed by LC-MS/MS using an Agilent 1260 LC coupled to a ABSciex QTrap

148 6500 instrument operated in negative mode electrospray ionization. Further details are provided in
149 Section S2 and Table S2. A full list of analyzed chemicals is provided in Table S3, along with
150 method detection limits (MDL) for each chemical in units of nanogram per liter. For mixture
151 modeling the detected chemical concentration was converted to molar units.

152

153 *Bioanalysis*

154 Information about the studied bioassays and the derivation of effect concentrations can be found in
155 Table 1 and Section S3. The data was expressed in units of relative enrichment factor (REF), which
156 takes into account sample enrichment by LVSPE and dilution in the assays. The data was expressed
157 as concentration causing 10% effect (EC₁₀), effect concentration causing an induction ratio of 1.5
158 (EC_{IR1.5}) or concentration causing 50% mortality (LC₅₀). Linear concentration-effect curves were
159 used to determine EC₁₀ and EC_{IR1.5}, while LC₅₀ was evaluated using log-logistic concentration-
160 effect curves.⁴ Cytotoxicity was assessed in parallel for the AhR, ER, oxidative stress response, p53
161 response and NF-κB assays and cell viability EC₁₀ values were derived from log-logistic
162 concentration-effect curves.³²

163

164 *Bioanalytical Equivalent Concentrations*

165 The LC and EC values were converted to BEQ_{bio} using Equation 1, with the LC₅₀ or EC₁₀ or EC_{IR1.5}
166 value of the reference compound (ref) and the matching LC₅₀ or EC₁₀ or EC_{IR1.5} value of the extract
167 only.

168

$$\text{BEQ}_{\text{bio}} = \frac{\text{LC}_{50}(\text{ref})}{\text{LC}_{50}(\text{extract})} \text{ or } \frac{\text{EC}_{10}(\text{ref})}{\text{EC}_{10}(\text{extract})} \text{ or } \frac{\text{EC}_{\text{IR1.5}}(\text{ref})}{\text{EC}_{\text{IR1.5}}(\text{extract})}$$

169

170

(1)

171

172 The BEQ concept has been typically applied to log-logistic concentration-effect curves; however,
173 for many environmental samples, linear concentration-effect curves may be more suitable for data
174 evaluation. This is because environmental samples may only induce low effects and to obtain the
175 50% effect concentration one would have to either use an unfeasibly high enrichment factor or
176 extrapolate the data. For linear concentration-effect curves to remain valid they should reach no
177 more than 20 to 30% effect or have an induction ratio (IR) no greater than 5 to ensure that they
178 remain in the linear range of the curve. Linear concentration-effect curves have previously been
179 shown to be a robust data evaluation method for environmental samples, individual chemicals and
180 chemical mixtures.¹⁰ It must be stressed that the BEQ concept is only valid if the slopes of the
181 sample and reference compound are parallel in log-logistic concentration-effect curves.³³ However,
182 parallel slopes are not a requirement for linear concentration-effect curves with a common intercept
183 at the effect axis as the BEQ is the ratio between concentrations at a given effect level and is
184 therefore independent of the effect level. The EC value from a linear concentration-effect curve was
185 calculated using Equation 2 using the example of EC₁₀, but the same equation is applicable for
186 EC₂₀, for example, with 20% used instead of 10% or EC_{IR1.5} with an IR of 1.5 as the effect
187 benchmark. BEQ can then be calculated using Equation 3 and while this example is for EC₁₀, this
188 ratio is constant across the entire linear concentration-effect curve range.

189

$$EC_{10} = \frac{10\%}{\text{slope}}$$

190

(2)

191

$$BEQ_{\text{bio}} = \frac{EC_{10}(\text{ref})}{EC_{10}(\text{extract})} = \frac{10\%}{\text{slope}(\text{ref})} \cdot \frac{\text{slope}(\text{extract})}{10\%} = \frac{\text{slope}(\text{extract})}{\text{slope}(\text{ref})}$$

192

(3)

193

194 To calculate BEQ_{chem} it was first necessary to determine the relative effect potency (REP_i) of the
195 detected chemicals (i). As the EC values for the detected chemicals were generally provided as EC_{50}
196 values in the literature, it was necessary to use EC_{50} values derived from log-logistic concentration-
197 effect curves for the AhR, PXR and ER assays. REP_i was calculated using Equation 4, with the
198 LC_{50} or EC_{50} or $EC_{IR1.5}$ value of the reference compound and the matching LC_{50} or EC_{50} or $EC_{IR1.5}$
199 value of detected chemical i.

$$REP_i = \frac{LC_{50} (ref)}{LC_{50} (i)} \text{ or } \frac{EC_{50} (ref)}{EC_{50} (i)} \text{ or } \frac{EC_{IR1.5} (ref)}{EC_{IR1.5} (i)}$$

201
202 (4)

203
204 All LC values for the detected chemicals in the FET test were collected from Scholz et al.²⁷, while
205 the EC values were collected from the peer-reviewed literature (AhR, PXR, ER assays) or the
206 ToxCast database (oxidative stress response, p53 response, NF- κ B assays),²³ which includes over
207 1800 compounds in over 800 different assays. All ToxCast data was re-evaluated to determine
208 $EC_{IR1.5}$ using linear concentration-effect curves. As each chemical in the ToxCast database was run
209 multiple times, it was possible to determine the mean $EC_{IR1.5}$ value and the associated standard
210 deviation. BEQ_{chem} was calculated for each JDS sample using REP_i and the detected concentration
211 (M) (Equation 5). The variability associated with BEQ_{chem} for the chemicals present in the ToxCast
212 database was assessed using error propagation. EC and LC values collected from the literature
213 generally did not include standard deviation, so it was not possible to determine the variability
214 associated with BEQ_{chem} for the AhR, PXR, ER and FET assays.

215

$$BEQ_{chem} = \sum_{i=1}^n REP_i \cdot C_i$$

216 (5)

217 **Results and Discussion**

218 *Chemical Analysis*

219 Of the 272 analyzed chemicals, 94 were detected at least once in the 22 JDS samples. The number
220 of chemicals detected at each site ranged from 20 to 64. The sum of the molar concentration and
221 number of detected chemicals at each site are shown in Figure 1A, with the concentrations in pM
222 for each of the detected chemicals at the different sampling sites shown in Table S4. The most
223 frequently detected chemicals were the artificial sweetener acesulfame, the industrial compounds
224 triphenylphosphine oxide and 2-benzothiazolesulfonic acid and the antimicrobial sulfamethoxazole,
225 which were present at detectable levels at all studied sites. In all but one tributary other common
226 wastewater micropollutants, including carbamazepine and its transformation products, the corrosion
227 inhibitors benzotriazole and methylbenzotriazole, the artificial sweeteners cyclamate and sucralose
228 and several herbicides and transformation products (metolachlor, isoproturon, atrazine, and
229 terbuthylazine-2-hydroxy), were detected. The antidiabetic pharmaceutical metformin was found at
230 the highest concentrations, with concentrations up to 7.6 nM. Overall, chemical contamination was
231 relatively low, with none of the detected chemicals exceeding the Water Framework Directive
232 environmental quality standards.³⁴

233

234 *Bioanalysis*

235 The EC and LC values for the different JDS water samples are shown in Figure 1B and Table S5,
236 with the concentration-effect curves for all assays shown in Figure S2. The assays indicative of
237 activation of ER, activation of PXR, activation of AhR and NF- κ B response tended to be the most
238 responsive, followed by the oxidative stress response. The p53 response occurred at higher effect
239 concentrations. The least responsive assay was the FET test, which required a REF of 100 to 500
240 for 50% mortality, or a REF of 50 to 300 for 10% mortality.

241

242 While most samples did have a response in the assays, the effects were relatively low, with the EC
243 values for the oxidative stress and AhR assays similar to previously benchmarked EC values for
244 surface water.³⁵ Kittinger et al.³⁶ also only detected minimal effects in Danube River samples when
245 assessing genotoxicity. Further, ER activation, when expressed as BEQ_{bio} (0.02-1.1 pM), was lower
246 than generally observed in wastewater effluent,³⁷ due to dilution in the river, though one
247 contaminated site, JDS 41 (BEQ_{bio} 4.7 pM), was identified by this assay.

248

249 JDS 64 had the lowest sum chemical concentration, and this corresponded to no effect at the
250 maximum REF for the oxidative stress response, p53 response and NF-κB assays, and only minimal
251 effects at high concentrations in the other assays. Cell viability was assessed in parallel for most
252 cell-based assays and in most cases there was negligible cytotoxicity in the studied concentration
253 range. However, cytotoxicity did mask other endpoints manifestation at high REFs in some samples
254 for the AhR (JDS 41 and 63), ER (JDS 35, 55, 57, 59, 63 and 67), oxidative stress response (JDS 55
255 and 67), p53 response (JDS 41, 55, 57 and 63), and NF-κB (JDS 36 and 41) assays. Hence, it was
256 not possible to derive EC values for induction for these particular samples, but EC₁₀ values for
257 cytotoxicity were calculated and are included in Table S5. JDS 41, which had the highest effect in
258 the ER activation and oxidative stress response assays and was cytotoxic in several other assays,
259 was the most polluted site with the highest amount of total detected chemical concentration.
260 Overall, there was no significant relationship between effect and sum detected chemicals at each
261 site for the different assays.

262

263 For mixture modeling, the EC and LC values were converted to BEQ_{bio} using the respective
264 reference compounds for each assay (Table 2, Table S5). While the EC or LC values give an
265 indication of the sensitivity of the assay, BEQ_{bio} converts the effect into the concentration of
266 reference compound that would elicit the same response as the sample mixture. Further, the BEQ
267 concept simplifies mixture toxicity modeling.^{10,11}

268

269 *Bioanalytical equivalent concentration from chemical analysis*

270 Prior to calculating BEQ_{chem} , the published literature and ToxCast database were searched for EC or
271 LC values for the detected chemicals. For each assay, between 0 and 13 literature EC or LC values
272 could be found for the 94 detected chemicals (Table S6). Using the literature EC or LC values for
273 each chemical and the EC or LC value of the assay reference compound, the REP_i was calculated
274 using Equation 4 (Table 3).

275

276 LC_{50} values at 48 h exposure for the FET test were collected from Scholz et al.²⁷, with REP_i
277 calculated using the mean 3,4-dichloroaniline LC_{50} value from the same study. Thirteen EC values
278 were collected from the literature for the PXR assay, while six EC values were available for the ER
279 assay. Although no EC values were available for the detected chemicals in the AhR CAFLUX
280 assay, EC values were available for three of the detected chemicals (terbutylazine, carbaryl,
281 daidzein) in the mouse AhR CALUX assay. While these assays focus on the same endpoint
282 (reporter gene expression), they utilize different animal cell lines (rat hepatoma versus mouse
283 hepatoma) and previous work has shown species-specific differences in responsiveness to some
284 AhR ligands.³⁸ To account for differences in sensitivity between the mouse and rat AhR models,
285 TCDD EC values were also collected from each study to calculate REP_i , rather than using the
286 TCDD EC value from the current study.

287

288 EC values for the oxidative stress response, p53 response and NF- κ B assays were collected from
289 the ToxCast database.²³ A total of 486, 278 and 62 chemicals in the ToxCast database were active
290 in the oxidative stress response, p53 response and NF- κ B assays, respectively (Figure 2). Of the 94
291 chemicals detected in the JDS samples, 49 of these were also included in the ToxCast database.
292 However, many of the detected compounds were not active in the assays, with 13 compounds active
293 in the oxidative stress response assay, 4 compounds active in the p53 response assay and none

294 active in the NF- κ B assay. $EC_{IR1.5}$ values for the detected chemicals in the oxidative stress response
295 and p53 response assays were calculated from raw emission data available in the ToxCast MySQL
296 database. To derive REP_i for the detected chemicals, experimental $EC_{IR1.5}$ values for reference
297 compounds tBHQ and mitomycin were used.

298

299 Using REP_i and the detected chemical concentration, BEQ_{chem} was calculated using Equation 5 for
300 each water sample (Table 2). BEQ_{chem} could not be calculated for some samples for the AhR and
301 ER bioassays as none of the chemicals with literature EC values were detected in the samples.
302 Further, it was not possible to derive BEQ_{chem} values for the NF- κ B assay as none of the detected
303 chemicals were active, despite the NF- κ B assay being one of the more responsive for the JDS
304 samples (Figure 1B). Only 3% of the 1859 chemicals in the ToxCast database were active in the
305 NF- κ B assay, compared to 26 and 15% of chemicals in oxidative stress response and p53 response
306 assays, respectively (Figure 2). The NF- κ B assay has been used for water quality monitoring in only
307 one study⁴ and it is still unclear what types of water-based pollutants induce a response in this
308 assay.

309

310 *What percent of biological effect can be explained by chemical analysis?*

311 The comparison between BEQ_{bio} and BEQ_{chem} for each assay is shown in Table 2, while the
312 contribution of the individual detected chemicals to the biological effect is shown in Figure 3. For
313 some JDS samples it was not possible to determine the contribution of detected chemicals to the
314 biological effect, and this was attributed to either cytotoxicity masking manifestation of other
315 endpoints, no effect at the maximum REF or the active chemicals being below the MDL.

316

317 The BEQ_{chem} for AhR activation was calculated using only three chemicals, but they explained
318 between 3 and 71% of the biological effect (Figure 3A). The effect was mostly driven by the
319 phytoestrogen daidzein, which has previously been shown to be a weak AhR activator in mouse

320 cells, but not in human cells,³⁹ and the herbicide terbuthylazine. The insecticide carbaryl only
321 explained 0.5% of the effect in JDS 67. Similarly, the BEQ_{chem} for activation of ER could explain
322 up to 80% of the effect, with the hormone estrone and the phytoestrogen genistein contributing
323 significantly. Estrogenic effects in wastewater are often explained by the presence of potent natural
324 and synthetic estrogenic hormones, such as 17 β -estradiol and 17 α -ethinylestradiol,^{14,18} but these
325 compounds were below the detection limit in the current study. Previous studies have also attributed
326 ER activation in river water to genistein.⁴⁰

327

328 In contrast, the detected chemicals could explain less than 0.2% of the biological effect in the
329 adaptive stress response assays, PXR assay and the FET test (Figure 3). A number of studies have
330 also shown similarly low contributions of detected chemicals to the oxidative stress response in a
331 range of water types including wastewater and pool water.^{10,17,41} It has also been demonstrated that
332 detected chemicals in surface water and wastewater can only explain a small fraction of PXR
333 activity.¹⁵ However, for this receptor, the use of the concentration addition model may be limited as
334 it has been recently demonstrated that, due to a large ligand-binding pocket, PXR can stably bind
335 binary mixtures of certain weakly active chemicals, which leads to synergistic activation of target
336 genes.⁴² The BEQ_{chem} vs BEQ_{bio} comparison for water samples has not been conducted previously
337 for the p53 response or FET assays, thus it was not possible to compare our results with the
338 literature. The herbicide metolachlor mostly contributed to the effect in the PXR assay. Genistein
339 dominated the contribution of quantified chemicals to the biological effect for the oxidative stress
340 response. While carbaryl and the disinfectant chlorophene were detected in the JDS samples and are
341 active in the oxidative stress response assay, they occurred only in samples where cytotoxicity
342 masked induction and could not be used to explain the biological effect. Genistein and the industrial
343 compound 2,4-dinitrophenol mostly contributed to the p53 response in samples collected from
344 Austria to Serbia (JDS 8 to 39), while the fungicide carbendazim dominated the effect further
345 downstream. Finally, 2,4-dinitrophenol and genistein together contributed up to 0.08% of the effect

346 explained by quantified chemicals in the FET assay, though antimicrobial triclosan alone could
347 explain up to 0.15% of the biological effect in JDS 59 (Figure 3).

348

349 The small contribution of detected chemicals to the biological response in the adaptive stress
350 response and PXR assays is not surprising since many compounds can activate these endpoints, as
351 discussed earlier. Further, 471 out of 641 or 74%, of reviewed compounds in Scholz et al.²⁷ had a
352 response in the FET test. Consequently, many compounds are active in these assays and while
353 comparability may have been improved with more literature EC and LC values for the detected
354 compounds, it is unlikely to have a significant influence on the comparison. To illustrate this point,
355 if we assume that 74% of chemicals should have an effect in the FET test, then 70 of the 94
356 detected chemicals could be active in this assay. However, published LC values were only available
357 for 12 of the detected chemicals. If we simply extrapolate the effect explained by the detected
358 chemicals with available LC₅₀ values in each sample to all 70 detected chemicals, without
359 considering differences in potencies, we can still only explain up to 1.6% of the effect. This
360 example does not take into consideration differences in mode of action or chemical potency, but
361 simply aims to illustrate the potential for many compounds to contribute to effect in apical
362 endpoints.

363

364 *Limitations and outlook*

365 There are some limitations associated with the current study. Primarily, improved understanding of
366 the contribution of the detected chemicals to the biological effect is hampered by the lack of REP_i
367 values for detected chemicals. Out of the 94 detected chemicals in the JDS samples, between 0 and
368 13 corresponding EC or LC values could be found for the different assays. While the US EPA
369 ToxCast program provides EC values for a large number of compounds, many of these are not
370 typical water pollutants and only 52% of chemical detected in the JDS samples were present in the
371 ToxCast database. Many of the detected chemicals in the ToxCast database were not active in the

372 adaptive stress response assays, but such information is not readily available for the other studied
373 assays. However, this information is important as it makes a difference to the effect balance if a
374 chemical's contribution is zero or if it is unknown. Consequently, fingerprinting the biological and
375 toxicological effect(s) of commonly detected water pollutants is recommended to help fill in the
376 knowledge gap. Further, the available literature data stems from a number of different sources and it
377 is possible that the experimental protocols for the same assay may differ slightly, leading to
378 potential differences in sensitivity or reproducibility. This limitation could be overcome by
379 improved standardization of bioassays.

380

381 A specific limitation associated with the AhR assay is that the EC values available in the literature
382 are based on the mouse AhR model, while BEQ_{bio} is based on the rat AhR model. Hence, potential
383 differences in species sensitivity may be a source of variability for the comparison of BEQ_{bio} and
384 BEQ_{chem} . A further limitation with using literature EC and LC values is that the error associated
385 with the value is often not provided. This was the case for the AhR, PXR, ER and FET assays and
386 consequently it was not possible to calculate the error associated with the BEQ_{chem} values. It was
387 possible to calculate the error associated with BEQ_{chem} for the oxidative stress response and p53
388 response assays as the $EC_{IR1.5}$ values used to derive REP_i were re-evaluated from a series of
389 replicate experiments from the ToxCast MySQL database (standard deviations associated with
390 BEQ_{chem} are provided in Table S7).

391

392 This study demonstrated the applicability of the BEQ concept to assess the contribution of detected
393 chemicals to the biological effect of chemical mixtures present in the Danube River. As the detected
394 chemicals could not explain a significant proportion of the effect, particularly in the adaptive stress
395 response, PXR and FET assays, this supports the application of bioanalytical tools complementary
396 to chemical analysis for water quality monitoring. Further, as targeted chemical analysis was
397 applied, we cannot exclude the fact that we may not have targeted the most relevant chemicals.

398 Consequently, further identification using tools such as effect-directed analysis may provide
399 improved understanding about chemical stressors in the Danube River.

400

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414

415 **Supporting Information:** Further information on the LVSPE sampling, analyzed chemicals and
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417

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596

Table 1: Overview of bioassays used in the current study

Endpoint	Assay	Method reference	Positive reference compound	Maximum REF	Data evaluation method	EC or LC value
Activation of AhR	CAFLUX	Nagy et al. ¹⁹	2,3,7,8-Tetrachlorodibenzo- <i>p</i> -dioxin (TCDD)	500	Linear concentration-effect curve	EC ₁₀
Activation of PXR	HG5LN-hPXR	Lemaire et al. ⁴³ ; Creusot et al. ¹⁵	SR 12813*	500	Linear concentration-effect curve	EC ₁₀
Activation of ER	MELN	Balaguer et al. ⁴⁴ ; Kinani et al. ⁴⁵	17 β -Estradiol	500	Linear concentration-effect curve	EC ₁₀
Oxidative stress response	ARE- <i>bla</i>	Invitrogen ⁴⁶	tert-Butylhydroquinone (tBHQ)	500	Linear concentration-effect curve	EC _{IRI,5}
p53 response	p53RE- <i>bla</i>	Neale et al. ⁴⁷	Mitomycin	500	Linear concentration-effect curve	EC _{IRI,5}
NF- κ B response	NF- κ B- <i>bla</i>	Jin et al. ⁴⁸	Tumor necrosis factor alpha (TNF α)	250	Linear concentration-effect curve	EC _{IRI,5}
Mortality	Fish embryo toxicity (FET)	OECD ²⁶	3,4-Dichloroaniline	1000	Log-logistic concentration-effect curve	LC ₅₀

598 *Tetraethyl 2-(3,5-di-tert-butyl-4-hydroxyphenyl)ethenyl-1,1-bisphosphonate

599 **Table 2:** BEQ_{bio} and BEQ_{chem} values for each sample in the different bioassays with the percentage of effect that can be explained by the detected

600 chemicals

ID	Activation of AhR			Activation of PXR			Activation of ER			Oxidative stress response			p53 response			Fish embryo toxicity		
	BEQ _{bio} (M)	BEQ _{chem} (M)	% effect	BEQ _{bio} (M)	BEQ _{chem} (M)	% effect	BEQ _{bio} (M)	BEQ _{chem} (M)	% effect	BEQ _{bio} (M)	BEQ _{chem} (M)	% effect	BEQ _{bio} (M)	BEQ _{chem} (M)	% effect	BEQ _{bio} (M)	BEQ _{chem} (M)	% effect
8	4.40 ×10 ⁻¹⁴	1.94 ×10 ⁻¹⁵	4.4%	3.43 ×10 ⁻⁹	6.51 ×10 ⁻¹³	0.02%	9.61 ×10 ⁻¹⁴	1.79 ×10 ⁻¹⁴	19%	3.11 ×10 ⁻⁸	1.10 ×10 ⁻¹¹	0.04%	1.51 ×10 ⁻¹⁰	4.00 ×10 ⁻¹⁴	0.03%	6.74 ×10 ⁻⁸	2.66 ×10 ⁻¹¹	0.04%
22	3.28 ×10 ⁻¹⁴	<MDL	-	4.98 ×10 ⁻¹⁰	3.65 ×10 ⁻¹³	0.07%	3.14 ×10 ⁻¹³	1.45 ×10 ⁻¹⁴	4.6%	2.75 ×10 ⁻⁸	1.37 ×10 ⁻¹²	0.01%	2.03 ×10 ⁻¹⁰	4.53 ×10 ⁻¹⁵	0.002%	5.88 ×10 ⁻⁸	6.48 ×10 ⁻¹²	0.01%
27	2.65 ×10 ⁻¹⁴	1.44 ×10 ⁻¹⁵	5.4%	2.40 ×10 ⁻⁹	4.94 ×10 ⁻¹³	0.02%	5.67 ×10 ⁻¹³	1.04 ×10 ⁻¹⁴	1.8%	3.59 ×10 ⁻⁸	1.77 ×10 ⁻¹¹	0.05%	2.46 ×10 ⁻¹⁰	6.39 ×10 ⁻¹⁴	0.03%	5.07 ×10 ⁻⁸	3.57 ×10 ⁻¹¹	0.07%
29	5.37 ×10 ⁻¹⁴	<MDL	-	2.13 ×10 ⁻⁹	4.31 ×10 ⁻¹³	0.02%	7.44 ×10 ⁻¹³	3.96 ×10 ⁻¹⁴	5.3%	3.40 ×10 ⁻⁸	1.45 ×10 ⁻¹¹	0.04%	4.04 ×10 ⁻¹⁰	7.74 ×10 ⁻¹⁴	0.02%	4.96 ×10 ⁻⁸	5.04 ×10 ⁻¹¹	0.1%
30	1.01 ×10 ⁻¹⁴	<MDL	-	3.45 ×10 ⁻¹⁰	3.42 ×10 ⁻¹³	0.10%	1.03 ×10 ⁻¹²	4.32 ×10 ⁻¹⁴	4.2%	2.05 ×10 ⁻⁸	2.39 ×10 ⁻¹¹	0.1%	2.59 ×10 ⁻¹⁰	3.88 ×10 ⁻¹⁴	0.01%	9.20 ×10 ⁻⁸	2.32 ×10 ⁻¹¹	0.03%
32	3.50 ×10 ⁻¹⁴	6.38 ×10 ⁻¹⁵	18%	<3.17× 10 ⁻¹¹	4.74 ×10 ⁻¹³	-	1.82 ×10 ⁻¹³	1.11 ×10 ⁻¹³	61%	5.93 ×10 ⁻⁸	1.27 ×10 ⁻¹⁰	0.2%	3.47 ×10 ⁻¹⁰	2.54 ×10 ⁻¹³	0.07%	2.11 ×10 ⁻⁷	1.31 ×10 ⁻¹⁰	0.06%
33	4.05 ×10 ⁻¹⁴	1.34 ×10 ⁻¹⁵	3.3%	7.72 ×10 ⁻¹⁰	4.82 ×10 ⁻¹³	0.06%	1.98 ×10 ⁻¹³	6.00 ×10 ⁻¹⁴	30%	3.62 ×10 ⁻⁸	5.19 ×10 ⁻¹¹	0.1%	1.89 ×10 ⁻¹⁰	1.31 ×10 ⁻¹³	0.07%	7.43 ×10 ⁻⁸	6.90 ×10 ⁻¹¹	0.09%
35	3.36 ×10 ⁻¹⁴	8.55 ×10 ⁻¹⁵	25%	4.61 ×10 ⁻⁹	6.45 ×10 ⁻¹³	0.01%	Cytotox	3.07 ×10 ⁻¹⁴	-	9.58 ×10 ⁻⁸	1.31 ×10 ⁻¹¹	0.01%	4.04 ×10 ⁻¹⁰	3.69 ×10 ⁻¹⁴	0.01%	7.08 ×10 ⁻⁸	2.47 ×10 ⁻¹¹	0.03%
36	2.94 ×10 ⁻¹⁴	<MDL	-	5.24 ×10 ⁻¹⁰	3.57 ×10 ⁻¹³	0.07%	3.47 ×10 ⁻¹³	2.37 ×10 ⁻¹⁴	6.8%	3.60 ×10 ⁻⁸	1.69 ×10 ⁻¹¹	0.05%	1.87 ×10 ⁻¹⁰	4.75 ×10 ⁻¹⁴	0.03%	3.54 ×10 ⁻⁸	2.57 ×10 ⁻¹¹	0.07%
37	<1.25 ×10 ⁻¹⁴	<MDL	-	1.53 ×10 ⁻⁹	3.19 ×10 ⁻¹³	0.02%	2.25 ×10 ⁻¹³	1.68 ×10 ⁻¹⁴	7.4%	2.79 ×10 ⁻⁸	2.28 ×10 ⁻¹²	0.01%	5.58 ×10 ⁻¹⁰	2.45 ×10 ⁻¹⁴	0.004%	4.80 ×10 ⁻⁸	1.37 ×10 ⁻¹¹	0.03%
39	2.02 ×10 ⁻¹⁴	1.02 ×10 ⁻¹⁴	50%	2.84 ×10 ⁻⁹	6.90 ×10 ⁻¹³	0.02%	1.15 ×10 ⁻¹³	4.63 ×10 ⁻¹⁴	40%	2.63 ×10 ⁻⁸	2.03 ×10 ⁻¹¹	0.08%	2.63 ×10 ⁻¹⁰	6.67 ×10 ⁻¹⁴	0.03%	4.90 ×10 ⁻⁸	3.99 ×10 ⁻¹¹	0.08%

41	Cytotox	1.91 ×10 ⁻¹⁴	-	1.91 ×10 ⁻⁹	4.84 ×10 ⁻¹³	0.03%	4.56 ×10 ⁻¹²	1.43 ×10 ⁻¹³	3.1%	1.58 ×10 ⁻⁷	1.25 ×10 ⁻¹¹	0.01%	Cytotox	1.18 ×10 ⁻¹³	-	7.22 ×10 ⁻⁸	1.48 ×10 ⁻¹¹	0.02%
44	1.85 ×10 ⁻¹⁴	59%	3.22 ×10 ⁻⁹	6.86 ×10 ⁻¹³	0.02%	4.47 ×10 ⁻¹³	3.06 ×10 ⁻¹⁴	6.8%	5.20 ×10 ⁻⁸	2.30 ×10 ⁻¹¹	0.04%	7.02 ×10 ⁻¹⁰	1.57 ×10 ⁻¹³	0.02%	5.93 ×10 ⁻⁸	3.58 ×10 ⁻¹¹	0.06%	
53	2.59 ×10 ⁻¹⁴	5.7%	1.28 ×10 ⁻⁹	4.75 ×10 ⁻¹³	0.04%	2.73 ×10 ⁻¹³	3.68 ×10 ⁻¹⁴	14%	5.69 ×10 ⁻⁸	3.67 ×10 ⁻¹²	0.01%	<1.01 ×10 ⁻¹⁰	2.91 ×10 ⁻¹⁴	-	4.98 ×10 ⁻⁸	1.83 ×10 ⁻¹¹	0.04%	
55	2.99 ×10 ⁻¹⁴	5.0%	6.08 ×10 ⁻¹⁰	5.16 ×10 ⁻¹³	0.08%	Cytotox	<MDL	-	Cytotox	2.82 ×10 ⁻¹²	-	Cytotox	1.36 ×10 ⁻¹³	-	5.32 ×10 ⁻⁸	1.85 ×10 ⁻¹¹	0.03%	
57	1.71 ×10 ⁻¹⁴	7.1%	1.88 ×10 ⁻⁹	3.99 ×10 ⁻¹³	0.02%	Cytotox	1.22 ×10 ⁻¹⁴	-	4.07 ×10 ⁻⁸	2.45 ×10 ⁻¹²	0.01%	Cytotox	1.06 ×10 ⁻¹³	-	5.10 ×10 ⁻⁸	1.58 ×10 ⁻¹¹	0.03%	
59	2.96 ×10 ⁻¹⁴	69%	2.21 ×10 ⁻⁹	6.94 ×10 ⁻¹³	0.03%	Cytotox	4.53 ×10 ⁻¹⁴	-	5.85 ×10 ⁻⁸	2.38 ×10 ⁻¹¹	0.04%	2.45 ×10 ⁻¹⁰	1.63 ×10 ⁻¹³	0.07%	4.94 ×10 ⁻⁸	9.91 ×10 ⁻¹¹	0.20%	
60	1.69 ×10 ⁻¹⁴	71%	<3.17 ×10 ⁻¹¹	4.42 ×10 ⁻¹³	-	3.77 ×10 ⁻¹³	1.19 ×10 ⁻¹⁵	0.31%	5.50 ×10 ⁻⁸	2.89 ×10 ⁻¹²	0.01%	2.23 ×10 ⁻¹⁰	1.04 ×10 ⁻¹³	0.05%	4.52 ×10 ⁻⁸	7.85 ×10 ⁻¹²	0.02%	
63	Cytotox	-	2.41 ×10 ⁻⁹	1.15 ×10 ⁻¹²	0.05%	Cytotox	4.14 ×10 ⁻¹⁴	-	8.11 ×10 ⁻⁸	8.74 ×10 ⁻¹²	0.01%	Cytotox	2.16 ×10 ⁻¹³	-	1.35 ×10 ⁻⁷	5.18 ×10 ⁻¹¹	0.04%	
64	5.87 ×10 ⁻¹⁵	-	1.25 ×10 ⁻¹⁰	7.21 ×10 ⁻¹⁴	0.06%	1.69 ×10 ⁻¹⁴	1.36 ×10 ⁻¹⁴	80%	<4.99 ×10 ⁻⁹	1.99 ×10 ⁻¹²	-	<1.01 ×10 ⁻¹⁰	1.88 ×10 ⁻¹⁴	-	4.49 ×10 ⁻⁸	1.80 ×10 ⁻¹¹	0.04%	
65	3.33 ×10 ⁻¹⁴	22%	9.97 ×10 ⁻¹⁰	6.28 ×10 ⁻¹³	0.06%	6.15 ×10 ⁻¹³	4.76 ×10 ⁻¹⁴	7.7%	6.53 ×10 ⁻⁸	4.20 ×10 ⁻¹²	0.01%	3.69 ×10 ⁻¹⁰	1.29 ×10 ⁻¹³	0.03%	5.85 ×10 ⁻⁸	1.78 ×10 ⁻¹¹	0.03%	
67	1.14 ×10 ⁻¹⁴	19%	8.43 ×10 ⁻¹⁰	5.74 ×10 ⁻¹³	0.07%	Cytotox	3.12 ×10 ⁻¹⁴	-	Cytotox	2.91 ×10 ⁻¹¹	-	4.52 ×10 ⁻¹⁰	1.42 ×10 ⁻¹³	0.03%	9.00 ×10 ⁻⁸	2.49 ×10 ⁻¹¹	0.03%	

601 MDL: method detection limit; cytotox: induction was masked by cytotoxicity

Table 3: The relative effect potencies (REP_i) for the detected chemicals in the different assays.

Assay	Activation of Ahr			Activation of PXR			Activation of ER			Oxidative stress response			p53 response			Fish embryo toxicity		
	No.	Chemical	REP _i	Chemical	REP _i	Chemical	Chemical	REP _i	Chemical ^k	REP _i	Chemical ^k	Chemical ^k	REP _i	Chemical ^k	REP _i	Chemical ^k	Chemical ^k	REP _i
	3*		3.33 × 10 ⁻⁶	Bezafibrate ^d	6.21 × 10 ⁻⁴	Benzophenone-3 ^g	7.04 × 10 ⁻⁷	2,4-Dinitrophenol	0.06	2,4-Dinitrophenol	1.09 × 10 ⁻³	2,4-Dinitrophenol	1.09 × 10 ⁻³	2,4-Dinitrophenol	1.09 × 10 ⁻³	2,4-Dinitrophenol	1.09 × 10 ⁻³	0.49
	2,3,7,8-		5.92 × 10 ⁻⁴	Bisphenol A ^d	4.63 × 10 ⁻³	Bisphenol A ^e	4.49 × 10 ⁻⁵	2-Phenylphenol	0.08	Carbendazim	7.35 × 10 ⁻³	Atrazine	7.35 × 10 ⁻³	Atrazine	7.35 × 10 ⁻³	Atrazine	7.35 × 10 ⁻³	0.04
			1.36 × 10 ⁻⁴	Carbamazepine ^d	4.61 × 10 ⁻⁴	Bisphenol S ^h	1.18 × 10 ⁻⁶	Bisphenol A	0.07	Diclofenac	1.02 × 10 ⁻³	Caffeine	1.02 × 10 ⁻³	Caffeine	1.02 × 10 ⁻³	Caffeine	1.02 × 10 ⁻³	3.41 × 10 ⁻³
				Diclofenac ^d	1.02 × 10 ⁻³	Daidzein ⁱ	6.47 × 10 ⁻⁵	Carbaryl	1.33	Genistein	1.26 × 10 ⁻³	Carbamazepine	1.26 × 10 ⁻³	Carbamazepine	1.26 × 10 ⁻³	Carbamazepine	1.26 × 10 ⁻³	0.05
				Diuron ^e	2.61 × 10 ⁻³	Estrone ^j	0.02	Chlorophene	0.11	Carbaryl	0.18	Carbaryl	0.18	Carbaryl	0.18	Carbaryl	0.18	0.18
				Erythromycin ^e	2.92 × 10 ⁻⁵	Genistein ⁱ	5.35 × 10 ⁻⁴	Daidzein	0.07	Diclofenac	0.45	Chlorotoluron	0.45	Chlorotoluron	0.45	Chlorotoluron	0.45	0.45
				Estrone ^e	3.92 × 10 ⁻³			Diclofenac	0.06	Diclofenac	0.13	Diclofenac	0.13	Diclofenac	0.13	Diclofenac	0.13	0.13
				Isoprotruron ^e	1.67 × 10 ⁻³			Genistein	0.74	Genistein	0.62	Genistein	0.62	Genistein	0.62	Genistein	0.62	0.62
				Ketoprofen ^d	3.59			Metolachlor	0.35	Metolachlor	1.22	Metoprolol	1.22	Metoprolol	1.22	Metoprolol	1.22	1.22

	$\times 10^{-4}$			$\times 10^{-3}$
Metolachlor ^f	0.10	Perfluorohexanoic acid	0.03	p-Nitrophenol
Prometryn ^d	8.71	Tri(butoxyethyl) phosphate	0.14	Salicylic acid
Terbuthylazine ^d	3.76	Triclosan	0.18	Triclosan
Triclosan ^d	0.02	Triethyl citrate	0.04	Triclosan

603 *REP; data from AhR CALUX using mouse hepatoma cell line

604 †Tetraethyl 2-(3,5-di-tert-butyl-4-hydroxyphenyl)ethenyl-1,1-bisphosphonate

605 ^aDenison et al.⁴⁹; ^bLong et al.⁵⁰; ^cGhisari et al.⁵¹; ^dCreusot et al.¹⁵; ^eCreusot⁵²; ^fLemaire et al.⁴³; ^gMolina-Molina et al.⁵³; ^hMolina-Molina et al.⁵⁴; ⁱcurrent study; ^jPillon
 606 et al.⁵⁵; ^kUS EPA ToxCast database²³; ^lScholz et al.²⁷

607 **List of Figures**

608

609 **Figure 1:** A) The sum molar concentration of chemicals detected at each JDS site (black bars),
610 along with the number of chemicals detected at each site (red circles) and B) LC or EC values for
611 all samples in units of relative enrichment factor (REF).

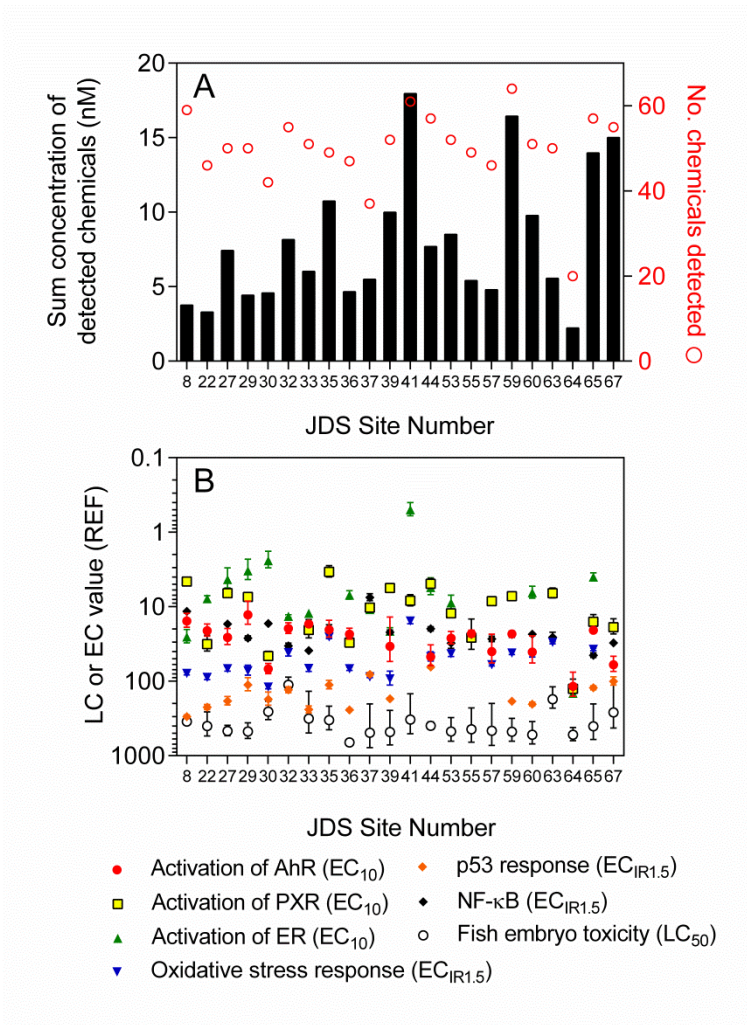
612

613 **Figure 2:** Overview of the active and inactive detected chemicals present in the ToxCast database
614 in the oxidative stress response (ARE, red), p53 response (blue) and NF- κ B response (green)
615 assays.

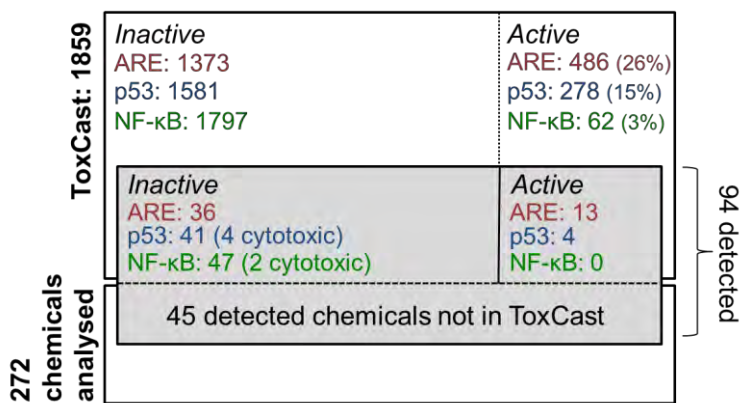
616

617 **Figure 3:** Percent of the biological effect explained by individual detected chemicals for A)
618 activation of AhR, B) activation of PXR, C) activation of ER, D) oxidative stress response, E) p53
619 response and F) fish embryo toxicity (FET)

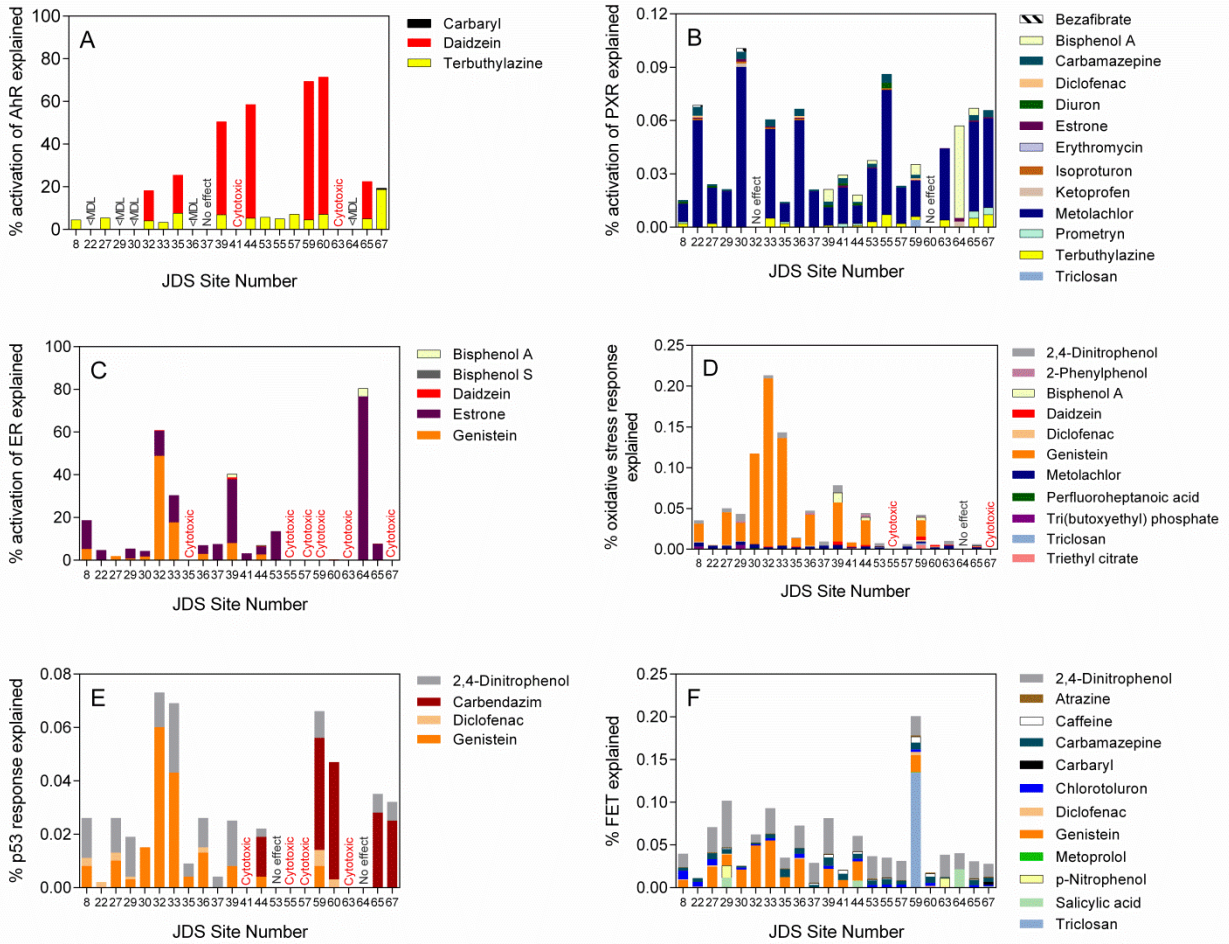
620

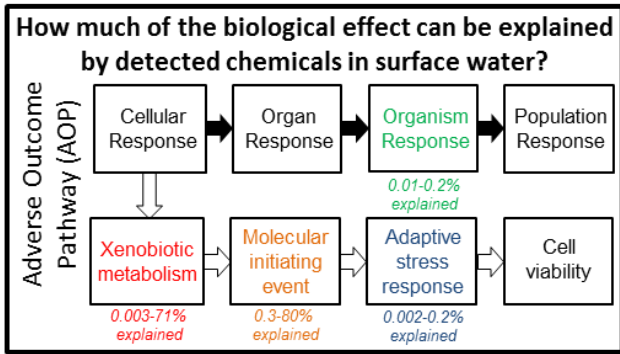


623 **Figure 2**



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