Biochemical and Cellular Biomarkers in Brown Trout (Salmo trutta f. fario) in Response to the Antidepressants Citalopram and Venlafaxine

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Research

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Abstract

Background: During the last decades, a worldwide increase in the number of cases of depression accompanied by rising prescription rates of antidepressants was recorded. In Germany, the two most prescribed antidepressants are the selective serotonin reuptake inhibitor (SSRI) citalopram and the serotonin and noradrenalin reuptake inhibitor (SNRI) venlafaxine, taking about 30% of the market share. Both antidepressants are found frequently in surface waters and have the potential of adversely affecting aquatic organisms. Most studies dealing with antidepressants address apical endpoints and behaviour, however, only few studies investigate biochemical biomarkers and histopathological alterations.

We conducted citalopram and venlafaxine exposure experiments with brown trout eggs in the eyed ova stage for five months, as well as with juvenile brown trout for four weeks. Exposure concentrations ranged from environmentally relevant 1 µg/L up to 1 mg/L. In this study, we investigated the effects of the antidepressants on β-esterase activity (neurotoxicity), stress protein level (proteotoxicity) and superoxide dismutase activity (oxidative stress). Additionally, we assessed the health status of the liver by means of histopathological analyses.

Results: We were able to show that both antidepressants did neither induce proteotoxic nor neurotoxic effects in brown trout. But for the antidepressant venlafaxine, the biochemical biomarker for oxidative stress (superoxide dismutase activity) was significantly increased in larvae exposed to at least 10 µg/L venlafaxine at 7°C. With regard to liver histopathology, fish exposed to higher citalopram concentrations were in a worse health condition than control fish irrespective of their life stage. Also the energy storage of fish exposed to 1 mg/L citalopram was reduced.

Conclusion: Thus, we here report citalopram-dependent histopathological alterations in brown trout liver, and the induction of oxidative stress by venlafaxine.

Background

Depression is one of the most common mental illnesses worldwide and has undergone a continuous increase during the last decades (34, 70). Worldwide, the number of cases of depression showed an increase of about 50% since 1990 (34). Associated with an increasing number of cases of depression, the number of prescribed antidepressants is constantly rising. In Germany, prescription rates of antidepressants increased about 40% between 2009 and 2018 (59). Selective serotonin reuptake inhibitors (SSRI) and serotonin and noradrenalin reuptake inhibitors (SNRI) constitute the majority of antidepressants. Two of the most prescribed antidepressants worldwide are citalopram (SSRI) and venlafaxine (SNRI). They amount to about 30% of the market share in Germany (59). This leads to a total consumption of 4.5 t (262.3 million defined daily doses (DDD)) of citalopram and 20.5 t (205.4 million DDD) of venlafaxine in 2018 in Germany only (59). Both antidepressants affect a similar biochemical pathway by inhibiting the reuptake of neurotransmitters like serotonin and noradrenalin from the synaptic cleft. They are known to be metabolised in liver and kidney of the human body. However, 12% of citalopram and 4.7% of venlafaxine are excreted unmetabolised via the urine and, therefore, enter wastewater treatment plants (22, 37). There, 18–27% of the citalopram is degraded, while the degradation of venlafaxine is even less efficient, amounting to 1–19% (19, 32). Therefore, both
antidepressants are found frequently in surface waters around the world (16, 17, 20, 21, 31, 47, 57). Citalopram surface water concentrations commonly range from 4 ng/L to 219 ng/L (47, 57) but, occasionally, also concentrations of up to 8 µg/L have been found in Indian lakes (17). Concentrations of citalopram in wastewater treatment plant effluents are in the range of 44 ng/L to 431 ng/L (21, 47, 62), but up to 76 µg/L citalopram was found downstream of an Indian wastewater treatment plant, and up to 840 µg/L in the effluent of an Indian drug manufacturer (17, 33). Venlafaxine concentrations in surface waters are higher than those for citalopram and range between 8 and 690 ng/L in wastewater-dominated streams (16, 20, 21).

Moreover, up to 2.5 µg/L venlafaxine was found in wastewater effluents (32). Being detected in such high concentrations, the effects of antidepressants, particularly of venlafaxine and citalopram, on aquatic life are important to being investigated. For both antidepressants, most studies were conducted regarding behavioural and apical endpoints in several fish species (7, 61). Behavioural changes in fish following exposure to venlafaxine or citalopram include changes in circadian rhythm (43), decreased escape behaviour (50), reduced attacks on food (26), changes in swimming behaviour (28, 40) and increased curiosity (27).

With regard to biochemical or structural biomarkers, however, knowledge about possible effects of citalopram and venlafaxine in aquatic wildlife is scarce. While venlafaxine did not affect the activity of the enzymes citrate synthase, catalase and pyruvate kinase in zebrafish, effects on superoxide dismutase (SOD) and acetylcholine esterase activities and the heat shock response were detected in fish exposed to different antidepressants (41, 71, 73). In respect to possible histopathological consequences in fish different results have been reported. Fathead minnow exposed to either venlafaxine, fluoxetine or sertraline did not reveal any histological effects in liver (58), while zebrafish and *Chichlosoma dimerus* exposed to fluoxetine, sertraline, paroxetine or mianserine showed histopathological alterations in liver, spleen and head kidney (48, 54). Thus, no consistent picture of possible effects of these pharmaceuticals on suborganismic effects in aquatic species can be drawn. In addition, data on the interplay of antidepressants and the confounding factor temperature are lacking up to now, and potential variation in the sensitivity of different life stages of fish has not been investigated so far. Consequently, we conducted experiments with brown trout (*Salmo trutta f. fario*), a feral fish species at different life stages at two temperatures. Eggs in the eyed-ova stage and juvenile fish were exposed to environmentally relevant and explicitly higher concentrations of the antidepressants citalopram and venlafaxine with focus on neurotoxicity (b-esterase activity), oxidative stress (SOD activity), proteotoxicity (stress protein level) and liver histopathology (overall health condition).

B-esterases, including acetylcholinesterase and carboxylesterases among others, are enzymes that are inhibited by paraxon (2, 30). Acetylcholinesterase (AChE) is an enzyme present in the synaptic cleft of cholinergic synapses, hydrolysing the neurotransmitter acetylcholine into its subcomponents acetic acid and choline (52). AChE is inhibited by various substances, such as carbamates or organophosphates and, thus, their inhibition can be seen as an adequate biomarker for neurotoxicity (71). Carboxylesterases (CbE) are known to detoxify insecticides like organophosphates, carbamates or pyrethroids and protectively counteract organophosphate-induced acetylcholinesterase inhibition (8, 56). It has already been shown, that antidepressants like citalopram, fluoxetine or sertraline affect acetylcholinesterase activity in aquatic organisms (45, 71, 72).

Superoxide dismutase (SOD) is an enzyme of the first line of the antioxidant defence system that catalyses the dismutation of O$_2$− radicals to hydrogen peroxide (H$_2$O$_2$). This compound can subsequently be
decomposed to water and oxygen, catalysed by the enzyme catalase (53, 71). Reactive oxygen species (ROS) are formed in several biological reactions, however, can also be induced by xenobiotics (51). These ROS, including superoxide radicals (O$_2^-$) are known to react with proteins, DNA and lipids resulting in oxidative stress for the organism (69). Therefore, an increased SOD activity is accompanied by an increased amount of ROS and thus represents an appropriate biomarker for oxidative stress (66). An increased SOD activity has been shown in amitriptyline-exposed zebrafish (73), sertraline-exposed goldfish (71), venlafaxine-exposed meagre (41) and citalopram-exposed daphnids (72).

Stress proteins or heat shock proteins with 70 kDa molecular weight (Hsp70) are chaperones, that bind to unfold, malfold or newly synthesised proteins to assist correct folding to the native protein (15, 18). These Hsp70s are induced by an increased portion of malfolded proteins in the cytoplasm and their induction is therefore regarded as a suitable biomarker for proteotoxicity (29). Maulvault, et al. (41) have shown an elevated heat shock protein response and ubiquitin level in meagre exposed to venlafaxine.

The liver is the main metabolic organ of vertebrates and is responsible for many vital functions like the metabolism of proteins, lipids and carbohydrates, as well as the detoxification of xenobiotics (55). Therefore, histopathological investigation of the liver is a powerful tool to assess sublethal effects of xenobiotics in fish (4, 25, 64). In zebrafish exposed to the antidepressants sertraline, paroxetine or mianserine a reduced proliferation of liver hepatocytes has been detected (48). Changes in liver-associated macrophage centres were shown in fluoxetine-exposed *Chichlosoma dimerus* (54).

The present study has been designed to complement existing data on citalopram and venlafaxine toxicity exerted on freshwater organisms, particularly in respect to biochemical and cytological effects of these two compounds. Behavioural and apical endpoints that, furthermore, have been investigated in our experiments have been previously published (74, 75).

**Material And Methods**

**Test organisms**

Juvenile brown trout and brown trout eggs were purchased from a commercial trout farm in Southern Germany (Forellenzucht Lohmühle, Alpirsbach-Ehlenbogen, Germany). This commercial fish breeder is listed as disease free (category I) according to the EC Council Directive (14). The eggs were obtained in the eyed ova stage and directly transferred into the experiments. Experiments with brown trout larvae took place in two climate chambers set on 7 °C or 11 °C, respectively, with a 10:14 hour light-dark-cycle. Prior to the experiments, juvenile brown trout were kept in an acclimation tank for at least one week. Experiments with juvenile brown trout were carried out in a climate chamber at 7 °C and the same photoperiod. All animal experiments were approved by the animal welfare committee of the Regional Council of Tübingen, Germany (ZO 2/16).

**Test substances**
Citalopram hydrobromide (C_{20}H_{21}F_{2}N_{2}O · HBr, CAS: 59729-32-7) and venlafaxine hydrochloride (C_{17}H_{27}NO_{2} · HCl, CAS: 99300-78-4) were obtained by Sigma Aldrich (Steinheim, Germany). Citalopram and venlafaxine stock solutions with concentrations of 1 and 100 mg/L were prepared with double distilled water and diluted to the respective test concentrations with filtered tap water (iron filter, active charcoal filter, particle filter). The concentrations refer to the respective free base substance (citalopram: C_{20}H_{21}F_{2}N_{2}O; venlafaxine: C_{17}H_{27}NO_{2}).

Exposure conditions

In both experiments, real chemical concentrations only differed slightly from nominal concentrations by 16% for venlafaxine and 19% for citalopram on the average, and limnochemical analyses revealed optimal test conditions with respect to temperature, oxygen content, pH and conductivity. Detailed data can be taken from (74, 75).

Experiments with brown trout larvae

As described previously (74, 75), the eyed-ova staged eggs of brown trout were exposed in a semi-static randomised three-block design at 7 °C and 11 °C. The respective nominal exposure concentrations of either citalopram or venlafaxine were 0, 1, 10, 100, 1000 µg/L. Fish were exposed in 25 L aquaria containing 10 L of the respective test solution, half of which was renewed twice a week. From the total yolk sac consumption until the end of the experiment fish were fed daily 3% of body weight with commercial trout feed (first 0.5 mm, then 0.8 mm, Inico Plus, Biomar, Brande, Denmark). Fish were exposed until eight respectively seven weeks after yolk sac consumption in the citalopram respectively venlafaxine experiment. At the end of the experiments, fish were euthanised with an overdose of the anaesthetic MS222 (1 g/L tricaine methanesulphonate buffered with NaHCO_{3}), followed by a cervical spine cut. Subsequently, fish were dissected. Livers were used for histopathology, heads for analyses of the b-esterase activity and the dorsal body part containing muscle and kidney for testing superoxide dismutase activity (only for the venlafaxine experiment). Livers were directly fixed in 2% glutaraldehyde (GA) dissolved from 25% GA with cacodylic buffer (0.1 M, pH 7.6) and stored at 4 °C until further processing. Samples for superoxide dismutase activity analysis were rinsed in PBS buffer (pH 7.4), and, along with the samples taken for b-esterase activity, frozen in liquid nitrogen and stored at -80 °C.

Experiments with juvenile brown trout

As described in previous publications (74, 75), juvenile brown trout were exposed to 0, 1, 10, 100 and 1000 µg/L of either citalopram or venlafaxine at 7° C in a semi-static randomised block design with three replicates. The fish were exposed in 25 L tanks containing 15 L of the respective concentrations. Twice a week, half of the respective test solution was replaced by freshly prepared test medium. Trout were fed 3% of body weight daily (0.8 mm, Inico Plus, Biomar, Brande, Denmark). After approximately four weeks of exposure (citalopram experiment: 28 days, venlafaxine experiment: 25 days), fish were euthanised with an overdose of MS222 (1 g/L buffered with NaHCO_{3}) followed by a cervical spine cut. Samples were taken from the liver for
histopathological analyses, the head was used for analyses of b-esterase activity, the dorsal part of the body containing mainly muscle and kidney was taken for superoxide dismutase testing (only venlafaxine experiment), and the gills were used for stress protein analysis. Liver samples were directly fixed in 2% glutardialdehyde (GA) dissolved from 25% GA with cacodylic buffer (0.1 M, pH 7.6) and stored at 4 °C. Samples for superoxide dismutase activity were rinsed in PBS buffer (pH 7.4) and, along with samples for the other biochemical biomarkers, frozen in liquid nitrogen and stored at -80 °C.

Histopathology

After at least one week of fixation, samples were washed three times with cacodylic buffer (0.1 M, pH 7.6), dehydrated in an ascending ethanol series, followed by 2-propanolol as an intermedium and infiltrated with paraffin wax in a tissue processor (TP 1020, Leica, Wetzlar, Germany). Afterwards, 3 µm sections were cut with a sledge microtome (SM 2000 R, Leica, Wetzlar, Germany). One of the microscope slides were stained with haematoxylin-eosin as an overview staining, and every other slide with alcian blue in combination with periodic acid-Schiff reagent (alcian blue-PAS) as a stain for glycogen and mucopolysaccharides. Slides were visually analysed using a light microscope (Axioskop 2, Zeiss, Oberkochen, Germany). A first evaluation of all samples was conducted to get an overview and to qualitatively assess occurring pathologies. In a second round, samples were re-labelled in a randomised order to obtain an observer-blinded assessment. The second evaluation was semi-quantitative, in which samples were classified into five different categories (1 - control, 2 - slight reaction, 3 - moderate reaction, 4 - strong reaction, 5 - destruction) according to the criteria described by Triebskorn, et al. (65). Additionally, the glycogen content of the samples was classified into three categories (1 - low, 2 - medium, 3 - high glycogen content). After the observer-blind evaluation, the respective class was assigned to the corresponding treatment.

Biochemical biomarkers

Superoxide dismutase activity

Superoxide dismutase (SOD) activity was measured in venlafaxine-exposed fish only, by using the superoxide dismutase assay kit by Cayman Chemical (Item No. 706002, Cayman Chemical Company, Ann Arbor, Michigan, USA). Tissues were homogenised in HEPES buffer (20 mM HEPES, 1 mM EGTA, 210 mM mannitol, 70 mM sucrose, pH 7.2, tissue(buffer ratio 1:5 w/v) with a pestle, and subsequently centrifuged (1500 × g, 5 min, 4 °C). After centrifugation, the supernatant was taken, mixed with Tris-HCl buffer (50 mM, pH 8) (supernatant/buffer ratio 1:30 w/v) and stored at -80 °C. Before performing the assay, the radical detector solution was freshly prepared from 19.95 mL assay buffer (50 mM Tris-HCl, 0.1 mM DTPA, 0.1 mM Hypoxanthine) and 50 µL tetrazolium salt solution (Item No. 706004, Cayman Chemical Company, Ann Arbor, Michigan, USA). Additionally, the SOD-standard curve (0, 0.005, 0.01, 0.02, 0.03, 0.04, 0.05 SOD activity (U/mL)) was freshly prepared containing bovine erythrocyte SOD (Item No. 706005, Cayman Chemical Company, Ann Arbor, Michigan, USA). The assay was performed in 96-well plates with each well being filled with 200 µL radical detector. In the first wells, 10 µL of the standard and in the remaining wells, 10 µL sample was pipetted in duplicates. Subsequently, 50 µL xanthine oxidase (Item No. 706006, Cayman Chemical
Company, Ann Arbor, Michigan, USA) was mixed with 1.95 mL assay buffer. 20 µL of this solution was given in each well, the plate was sealed with adhesive foil and incubated for 30 min at room temperature. Subsequently, the plate was measured photometrically at 450 nm and the SOD-activity was calculated according to the following equation:

\[
\left( \frac{\text{sample linearised rate} - \text{y-intercept}}{\text{slope}} \right) \times \left( \frac{0.23 \text{ ml}}{0.01 \text{ ml}} \right) \times \text{sample dilution}
\]

Results are expressed as unit (U)/mL. One unit is defined as the amount of enzyme needed to exhibit 50% of the superoxide radical.

**B-esterase activity**

For measuring the b-esterase activity, fish head samples were homogenised manually in Tris-buffer (10 mM Tris, 10 mM NaCl, pH 7.3) mixed with protease inhibitors (aprotinin, leupeptin, pepstatin, antipain, trypsin). After homogenisation, samples were centrifuged (5000 × g, 10 min, 4 °C), the supernatant was taken and 10% glycerol was added before storage at -20 °C. The protein content of the samples was determined photometrically at 650 nm according to the method of Lowry, et al. (36), modified by Markwell, et al. (39). Bovine serum albumin was used as a standard. The determination of the acetylcholinesterase (AChE) activity was conducted according to Ellman, et al. (13), modified by Rault, et al. (52) and measured for five minutes at 405 nm. The activity of the carboxylesterases (CbE) was analysed using the substrates 4-nitrophenylacetate (NPA) and 4-nitrophenylvalerate (NPV). Determination was conducted according to Carr and Chambers (8) and Chanda, et al. (9), modified by Sanchez-Hernandez, et al. (56). The activity was also measured for five minutes at 405 nm. All samples were analysed in triplicates. The activity of AChE and CbE is referred to the protein content and expressed as milliunits (mU)/mg protein. One unit is defined as one micromole of substrate hydrolysed per minute.

**Stress protein level**

The quantification of 70 kDa stress proteins (Hsp70) was conducted in gill samples of the juvenile fish. Samples for larvae were not analysed in this assay due to the low weight of the samples and the high base loads of stress proteins in developing brown trout (24). Gills were homogenised with 98% of extraction buffer and 2% protease inhibitor (tissue/buffer ratio 1:4 w/v) according to the protocol of Dieterich, et al. (10). The total protein content was quantified according to Bradford (6). For the quantification of Hsp70, a total amount of 40 µg total protein was analysed. The proteins were separated by sodium dodecyl sulphate polyacrylamide gel electrophoresis (SDS-PAGE) and subsequently blotted on a nitrocellulose membrane in a semi-dry chamber according to Dieterich, et al. (10). Subsequently, the protein bands on the membrane were immunostained with a monoclonal α-Hsp70 IgG (mouse anti-human Hsp70, Dianova, Hamburg, Germany) and secondary peroxidase-coupled α-IgG (goat anti-mouse IgG conjugated to peroxidase, Jackson ImmunoResearch, West Grove, Pennsylvania, USA). Subsequently, membranes were stained with 4-chloro-1-naphthol until the protein bands became visible. Finally, the optical volume (= band area × average grey scale
value) of the bands was quantified by planimetry and densitometry and referred to an internal standard (brown trout total body homogenate). Results are expressed relative to the standard.

**Statistics**

Statistical analyses were performed with SAS JMP 14 software. Data for histological evaluation, either pathological classes or glycogen classes, were analysed with the likelihood-ratio-\(\chi^2\)-test. Whenever significant differences occurred, single comparisons between control and each treatment were also performed with the likelihood-ratio-\(\chi^2\)-test. To correct for multiple comparisons, the \(\alpha\)-level was adjusted according to the method developed by Benjamini and Hochberg (3). Data for biochemical biomarkers were analysed using a nested ANOVA with a subsequent post-hoc Dunnett’s test. If necessary, data were transformed to fulfil model assumptions. If no normal distribution could be achieved by transformation, data were analysed using a Kruskal-Wallis-Test with subsequent post-hoc Steel with control. If no homogeneity of variance was given, data were analysed using a Welch ANOVA with subsequent post-hoc Dunnett’s test. The \(\alpha\)-level was always set to 0.05.

**CRED**

Criteria for reporting and evaluating ecotoxicity data (CRED) aim to improve the evaluation, transparency, reproducibility and consistency of reliability of ecotoxicological studies. For the experiment with citalopram CRED were previously published and provided by Ziegler, et al. (74) and, for the experiments with venlafaxine, CRED are provided in Ziegler, et al. (75).

**Results**

**Histopathology**

Generally, the health status of the liver was negatively affected by an exposure in the experimental setup.

In the citalopram experiments, livers of the control and the 1 and 10 µg/L-exposed animals were characterized by large hepatocytes with a bright cytoplasm (Figure 1 A). In contrast, brown trout exposed to 100 and 1000 µg/L citalopram had darker cells and regularly showed hyperplasia of hepatocytes (Figure 1 C). In livers of fish exposed to 1000 µg/L citalopram, inflammations occurred more frequently (Figure 1 B). Additionally, livers of fish exposed to the highest treatment showed reduced glycogen contents compared to the control fish (Figure 1 D, E). In juvenile fish treated with 100 or 1000 µg/L citalopram, vacuolisation was more pronounced than in the controls. In all treatments, dilated capillaries, necrosis and macrophages occurred (Figure 1 C, F). When comparing the overall health status of the larvae, those animals exposed at 11 °C generally were in a worse condition than fish exposed at 7 °C. Livers of larvae exposed to 1000 µg/L citalopram at 11 °C and those exposed to 100 µg/L citalopram at 7 °C, as well as juvenile brown trout exposed to 1000 µg/L citalopram were in a significantly worse condition than livers of control fish (Figure 2 A-C) (brown trout larvae 11 °C: likelihood-ratio-\(\chi^2\)-test: \(df=16, \chi^2=30.986, p=0.0135\), pairwise comparison [control|1000 µg/L] \(p=0.0036\); brown trout larvae 7 °C: likelihood-ratio-\(\chi^2\)-test: \(df=16, \chi^2=40.467, p=0.0007\),...
pairwise comparison [control|1000 µg/L] \( p=0.0047 \); juvenile brown trout: likelihood-ratio-\( \chi^2 \)-test: \( df=16 \), \( \chi^2=30.706 \), \( p=0.0147 \), pairwise comparison [control|1000 µg/L] \( p=0.0095 \)). Furthermore, brown trout larvae treated with 1000 µg/L citalopram had a significantly decreased glycogen content when exposed at 11 °C (Figure 2 D) (brown trout larvae 11 °C: likelihood-ratio-\( \chi^2 \)-test: \( df=8 \), \( \chi^2=48.951 \), \( p<0.0001 \), pairwise comparison [control|1000 µg/L] \( p=0.0008 \)). Both, brown trout larvae exposed to 1000 µg/L at 7 °C and juvenile brown trout exposed to 1000 µg/L had, tendentially a lower glycogen content than the control (Figure 2 E, F). However, this effect was not significant (brown trout larvae 7 °C: likelihood-ratio-\( \chi^2 \)-test: \( df=8 \), \( \chi^2=19.807 \), \( p=0.0111 \), pairwise comparison revealed no difference to control; juvenile brown trout: likelihood-ratio-\( \chi^2 \)-test: \( df=8 \), \( \chi^2=8.804 \), \( p=0.3591 \)). Generally, the glycogen content appeared to be lower in juvenile brown trout than in brown trout larvae. With regard to the liver condition however, no distinct differences between larvae and juveniles were observed.

After exposure to venlafaxine as well as in control fish larvae, livers were characterized by large hepatocytes with a bright cytoplasm (Figure 1 A). Throughout all treatment groups at 11 °C, inflammations and macrophage infiltration occurred more frequently (Figure 1 B) compared to larvae exposed at 7 °C (and the respective controls). In fish exposed at 11 °C, necrosis and fibrosis was more abundant than in fish exposed at 7 °C, however, no difference between the control and venlafaxine- treated groups were detected. The liver conditions of venlafaxine-exposed juvenile brown trout ranged between category 1 and 5. In most cases, the cytoplasm of the hepatocytes did not appear bright (Figure 1 C) and the cells were medium-sized. In livers of juvenile fish exposed to 10 and 1000 µg/L, inflammations were observed more frequently than in the other treatments (Figure 1 B). Dilated capillaries, intercellular spaces, vacuolisation, irregular shaped nuclei and necrosis occurred equally throughout all treatments (Figure 1 C, F). The semi-quantitative assessment revealed no difference in the health status between controls and venlafaxine-exposed fish (brown trout larvae 11 °C: likelihood-ratio-\( \chi^2 \)-test: \( df=16 \), \( \chi^2=10.701 \), \( p=0.8275 \); brown trout larvae 7 °C: likelihood-ratio-\( \chi^2 \)-test: \( df=16 \), \( \chi^2=15.494 \), \( p=0.4888 \); juvenile brown trout: likelihood-ratio-\( \chi^2 \)-test: \( df=16 \), \( \chi^2=19.575 \), \( p=0.24 \)). Likewise, the glycogen content of fish did not differ significantly from the respective controls at both temperatures (brown trout larvae 11 °C: likelihood-ratio-\( \chi^2 \)-test: \( df=8 \), \( \chi^2=9.941 \), \( p=0.2692 \); brown trout larvae 7 °C: likelihood-ratio-\( \chi^2 \)-test: \( df=8 \), \( \chi^2=6.98 \), \( p=0.5388 \); juvenile brown trout: likelihood-ratio-\( \chi^2 \)-test: \( df=8 \), \( \chi^2=9.911 \), \( p=0.2713 \)). However, livers of venlafaxine-exposed brown trout larvae showed a lower glycogen content than the livers from fish in all other experiments.

**Biochemical biomarkers**
### Table 1

Biochemical biomarkers in brown trout larvae exposed to citalopram.

Results are given as arithmetic means and standard deviations. AChE= acetylcholinesterase, CbE= carboxylesterase, NPA= 4-nitrophenylacetate, NPV= 4-nitrophenylvalerate.

| concentration citalopram (µg/L) | 11 °C | 7 °C |
|---------------------------------|-------|------|
| AChE activity (mU/mg Protein)   |       |      |
| 0                               | 45.4  | 43.8 |
| ±                               | ± 8.4 | ± 9.1|
| 1                               | 43.8  | 47.0 |
| ±                               | ± 9.1 | ± 10.4|
| 10                              | 47.8  | 42.8 |
| ±                               | ± 10.4| ± 10.4|
| 100                             | 42.8  | 47.8 |
| ±                               | ± 8.6 | ± 8.6|
| 1000                            | 43.3  | 45.3 |
| ±                               | ± 10.8| ± 10.8|
| 0                               | 41.7  | 44.7 |
| ±                               | ± 9.1 | ± 7.9|
| 1                               | 44.7  | 43.3 |
| ±                               | ± 8.4 | ± 6.7|

| CbE activity (NPA substrate) (mU/mg Protein) |       |      |
| 0                               | 112.3 | 112.3|
| ±                               | ± 26.7| ± 19.1|
| 1                               | 121.5 | 121.5|
| ±                               | ± 28.1| ± 28.1|
| 10                              | 120.9 | 120.9|
| ±                               | ± 21.8| ± 21.8|
| 100                             | 123.0 | 123.0|
| ±                               | ± 26.7| ± 26.7|
| 1000                            | 125.0 | 125.0|
| ±                               | ± 12.9| ± 12.9|
| 0                               | 94.6  | 94.6 |
| ±                               | ± 15.3| ± 15.3|
| 1                               | 98.8  | 98.8 |
| ±                               | ± 20.0| ± 20.0|
| 10                              | 91.7  | 91.7 |
| ±                               | ± 15.3| ± 15.3|
| 100                             | 96.5  | 96.5 |
| ±                               | ± 21.4| ± 21.4|
| 1000                            | 96.2  | 96.2 |
| ±                               | ± 16.0| ± 16.0|

| CbE activity (NPV substrate) (mU/mg Protein) |       |      |
| 0                               | 82.1  | 82.1 |
| ±                               | ± 25.0| ± 21.5|
| 1                               | 95.8  | 95.8 |
| ±                               | ± 28.4| ± 25.0|
| 10                              | 97.6  | 97.6 |
| ±                               | ± 28.0| ± 28.0|
| 100                             | 96.4  | 96.4 |
| ±                               | ± 28.3| ± 28.3|
| 1000                            | 94.6  | 94.6 |
| ±                               | ± 26.5| ± 26.5|
| 0                               | 71.2  | 71.2 |
| ±                               | ± 16.4| ± 16.4|
| 1                               | 66.0  | 66.0 |
| ±                               | ± 23.5| ± 23.5|
| 10                               | 73.4  | 73.4 |
| ±                               | ± 19.7| ± 19.7|

### Table 2

Biochemical biomarkers in brown trout larvae exposed to venlafaxine.

Results are given as arithmetic means and standard deviations. AChE= acetylcholinesterase, CbE= carboxylesterase, NPA= 4-nitrophenylacetate, NPV= 4-nitrophenylvalerate, SOD= superoxide dismutase; asterisks represent significant differences to the respective control (*p ≤ 0.05; **p ≤ 0.01; ***p ≤ 0.001).

| concentration venlafaxine (µg/L) | 11 °C | 7 °C |
|---------------------------------|-------|------|
| AChE activity (mU/mg Protein)   |       |      |
| 0                               | 63.0  | 63.0 |
| ±                               | ± 8.0 | ± 11.1|
| 1                               | 64.4  | 59.9 |
| ±                               | ± 16.5| ± 16.5|
| 10                              | 59.9  | 61.6 |
| ±                               | ± 16.4| ± 16.4|
| 100                             | 57.1  | 57.1 |
| ±                               | ± 14.5| ± 14.5|
| 1000                            | 70.8  | 70.8 |
| ±                               | ± 14.8| ± 14.8|
| 0                               | 71.0  | 71.0 |
| ±                               | ± 11.0| ± 11.0|
| 1                               | 71.5  | 71.5 |
| ±                               | ± 15.1| ± 15.1|
| 10                               | 69.3  | 69.3 |
| ±                               | ± 15.9| ± 15.9|
| 100                             | 64.1  | 64.1 |
| ±                               | ± 12.0| ± 12.0|

| CbE activity (NPA substrate) (mU/mg Protein) |       |      |
| 0                               | 141.8 | 141.8|
| ±                               | ± 23.3| ± 26.6|
| 1                               | 146.5 | 146.5|
| ±                               | ± 27.9| ± 27.9|
| 10                              | 143.2 | 143.2|
| ±                               | ± 29.6| ± 29.6|
| 100                             | 130.2 | 130.2|
| ±                               | ± 22.9| ± 22.9|
| 1000                            | 129.9 | 129.9|
| ±                               | ± 22.4| ± 22.4|
| 0                               | 142.3 | 142.3|
| ±                               | ± 22.9| ± 22.9|
| 1                               | 145.5 | 145.5|
| ±                               | ± 19.5| ± 19.5|
| 10                               | 139.5 | 139.5|
| ±                               | ± 16.0| ± 16.0|
| 100                             | 135.5 | 135.5|
| ±                               | ± 23.9| ± 23.9|

| CbE activity (NPV substrate) (mU/mg Protein) |       |      |
| 0                               | 104.2 | 104.2|
| ±                               | ± 28.9| ± 28.0|
| 1                               | 108.2 | 108.2|
| ±                               | ± 32.8| ± 32.8|
| 10                              | 105.1 | 105.1|
| ±                               | ± 36.9| ± 36.9|
| 100                             | 94.5  | 94.5 |
| ±                               | ± 22.9| ± 22.9|
| 1000                            | 93.0  | 93.0 |
| ±                               | ± 23.7| ± 23.7|
| 0                               | 92.2  | 92.2 |
| ±                               | ± 20.1| ± 20.1|
| 1                               | 92.8  | 92.8 |
| ±                               | ± 19.4| ± 19.4|
| 10                               | 90.1  | 90.1 |
| ±                               | ± 13.7| ± 13.7|

| SOD activity (U/mL) |       |      |
| 0                               | 166.0 | 166.0|
| ±                               | ± 29.3| ± 31.7|
| 1                               | 149.3 | 149.3|
| ±                               | ± 40.4| ± 40.4|
| 10                              | 148.2 | 148.2|
| ±                               | ± 51.6| ± 51.6|
| 100                             | 158.8 | 158.8|
| ±                               | ± 29.3| ± 29.3|
| 1000                            | 161.1 | 161.1|
| ±                               | ± 37.5| ± 37.5|
| 0                               | 133.9 | 133.9|
| ±                               | ± 37.1| ± 37.1|
| 1                               | 140.2 | 140.2|
| ±                               | ± 36.0| ± 36.0|
| 10                               | 158.2 | 158.2|
| ±                               | ± 36.1| ± 36.1|
| 100                             | 165.6 | 165.6|
| ±                               | ± 34.9| ± 34.9|
| 1000                            | 155.7 | 155.7|
Table 3
Biochemical biomarkers in juvenile brown trout exposed to citalopram and venlafaxine.

Results are given as arithmetic means and standard deviations. AChE= acetylcholinesterase, CbE= carboxylesterase, NPA= 4-nitrophenylacetate, NPV= 4-nitrophenylvalerate, Hsp70= 70 kDa stress protein family, SOD= superoxide dismutase, n.a.= not assessed.

| concentration (µg/L) | citalopram | venlafaxine |
|---------------------|------------|-------------|
|                     | 0          | 1           | 10          | 100         | 1000 | 0           | 1           | 10          | 100         | 1000         |
| AChE activity (mU/mg Protein) | 48.6 ± 13.8 | 54.6 ± 14.6 | 48.0 ± 9.6  | 48.6 ± 9.0  | 58.3 ± 16.0 | 70.6 ± 13.4 | 73.6 ± 12.5 | 73.1 ± 10.9 | 71.2 ± 11.6 | 69.6 ± 9.0  |
| CbE activity (NPA substrate) (mU/mg Protein) | 66.6 ± 16.6 | 64.9 ± 17.7 | 63.4 ± 16.8 | 65.7 ± 17.7 | 61.6 ± 17.4 | 104.7 ± 20.2 | 111.4 ± 26.2 | 114.0 ± 19.0 | 113.1 ± 18.1 | 107.5 ± 19.6 |
| CbE activity (NPV substrate) (mU/mg Protein) | 55.6 ± 23.3 | 50.1 ± 25.6 | 50.1 ± 24.7 | 51.2 ± 26.4 | 42.7 ± 24.7 | 94.7 ± 29.2 | 102.8 ± 32.4 | 106.1 ± 26.0 | 106.8 ± 24.6 | 97.3 ± 22.9  |
| SOD activity (U/mL) | n.a. | n.a. | n.a. | n.a. | 88.4 ± 31.4 | 100.4 ± 38.9 | 91.6 ± 32.5 | 100.4 ± 31.1 | 107.9 ± 35.2 |
| Hsp70 level (relative greyvalue) | 2.10 ± 0.35 | 2.21 ± 0.58 | 1.95 ± 0.43 | 2.02 ± 0.44 | 2.02 ± 0.41 | 2.06 ± 0.24 | 2.02 ± 0.24 | 2.06 ± 0.23 | 2.13 ± 0.31 | 1.90 ± 0.22 |

Superoxide dismutase activity

Regarding the activity of the superoxide dismutase (SOD) in brown trout larvae exposed at 11 °C to venlafaxine, no differences between controls and chemical-treated fish were observed (brown trout larvae 11 °C: nested ANOVA: df=4,10, F=1.5387, p=1946). However, at 7 °C, the SOD activity of venlafaxine-exposed larvae was increased up to 24 % in relation to the control. Brown trout larvae exposed to 10 and 100 µg/L venlafaxine at 7 °C showed a significantly increased SOD activity compared to the respective controls, and for larvae exposed to 1000 µg/L the same trend became obvious (brown trout larvae 7 °C: nested ANOVA: df=4,10, F=3.7377, p=0.0064, post-hoc Dunnett’s Test [control 10 µg/L] p=0.0442, [control 100 µg/L] p=0.0048, [control 1000 µg/L] p=0.082). Comparing the SOD-activity between the two temperatures, the activity in control larvae exposed at 11 °C (165.96 ± 29.32 U/ml) was remarkably higher (24 %) than in larvae exposed at 7 °C (133.86 ± 37.49). Regarding the SOD activity of juvenile brown trout, the activity of fish treated with venlafaxine was on average higher than the SOD activity of control fish. The activity of the SOD in juvenile fish exposed to the highest treatment was increased by 22 % compared to the respective controls, however, the difference was not statistically significant (juvenile brown trout: nested ANOVA: df=4,10, F=1.5804, p=0.1834).
**B-esterase activity**

The b-esterase activity did not differ between antidepressant-treated fish and the respective controls in all experiments. Thus, the b-esterase activity was neither influenced by citalopram nor venlafaxine and, furthermore, did not depend on the life stage or temperature (statistical details in the supplement).

**Stress protein level**

Hsp70 levels of juvenile brown trout were neither influenced by citalopram nor venlafaxine (juvenile brown trout + citalopram: nested ANOVA: df=4,10, F= 1.7073, p=0.1522; juvenile brown trout + venlafaxine: nested ANOVA: df=4,10, F=3.2671, p=0.0138, post-hoc Dunnett's test revealed no difference to control).

**Discussion**

Histopathological analyses made evident that the health status of the liver was negatively affected by citalopram in fish exposed to 100 µg/L or 1000 µg/L of the pharmaceutical. The most prominent symptoms were hyperplasia of the hepatocytes and inflammations, which occurred in larvae exposed at both temperatures, as well as in juvenile brown trout. The impaired health status of the fish liver is plausibly related to the accumulation of citalopram in the tissue of exposed fish (74) which might be responsible for an increased biotransformation activity and metabolism rate. High metabolism rates in the liver are likely reflected by a reduction of glycogen in the fish liver and hyperplasia of hepatocytes (63). Additionally, oxidative stress could also have contributed to the depicted damages (1, 76). According to previous studies, antidepressants in general were shown to have the potential for inducing negative effects in liver tissue. Although Schultz, et al. (58) did not observe any liver alterations in response to either fluoxetine, sertraline or venlafaxine in fathead minnow, Nowakowska, et al. (48), however, reported a decrease in hepatocyte proliferation in zebrafish exposed to 10 µg/L of the SSRIs sertraline, paroxetine and the antidepressant mianserin. Rats exposed to the SSRI fluoxetine revealed histopathological liver injury, with symptoms like e.g. increased numbers of Kupffer and inflammatory cells as well as focal necrosis (49, 76). Likewise, rats exposed to 20 mg/kg citalopram showed increased inflammation in their livers, higher numbers of Kupffer cells and liver lymphocytes as well as an increased activity of liver enzymes (1, 44). Additionally, isolated rat hepatocytes exposed to 500 µM citalopram showed cell death, oxidative stress and mitochondrial toxicity (1). Also in humans, exposure to citalopram resulted in acute liver damage and increased levels of liver enzymes, indicating hepatitis (23, 46). The observed reduction of glycogen in brown trout exposed to 1000 µg/L very likely went along with the observed reduced food intake. A reduced food uptake may directly cause a decrease in glycogen storage in the liver (11, 67) and might be responsible for the lower growth rate in brown trout exposed to 1000 µg/L citalopram, compared to the controls (74).

Venlafaxine exposure did not cause severe liver damage in brown trout. Only in juveniles exposed to 10 and 1000 µg/L venlafaxine, slightly more inflammations became evident than in control fish. The lacking cellular reactions in the fish livers might be due to the fact, that tissue concentrations of venlafaxine were up to 500-fold lower than those for citalopram in citalopram-exposed fish (74, 75), resulting in lower metabolic and biotransformation activity. Similarly, Schultz, et al. (58) did not find any evidence of venlafaxine-dependent
histopathological alterations in fathead minnow livers. Furthermore, venlafaxine had no effect on the glycogen storage in the liver of exposed brown trout. Moreover, it has to be mentioned that also in control fish, slight histopathologies became obvious which might be due to the artificial exposure conditions in the aquaria which, however, were kept as appropriate as possible during the experiments. As previously stated by (60), this might be a general physiological response of the fish to the laboratory exposure. Furthermore, the liver status was generally impacted by temperature. Livers of larvae exposed at 11 °C were in a worse condition with increased inflammations and necrosis than livers of larvae exposed at 7 °C. The diminished health condition might be related to the higher metabolism of fish exposed at 11 °C. In a study with Japanese flounder also a reduced liver health status could be shown in relation to higher temperature (35). Additionally, data for SOD-activity suggested a higher oxidative stress when exposed to the higher temperature with SOD values being 24% higher at 11 °C than at 7 °C. This difference in SOD activity might be influenced by the higher temperature of 11 °C, but differences due to different exposure time cannot be excluded. Additionally, larvae exposed at 11 °C showed a higher mortality rate compared to larvae exposed at 7 °C substantiating the poorer health condition at 11 °C (75). The higher temperature might have increased the overall SOD-activity which could possibly resulted in the worse health status and the elevated mortality. In several studies it has become evident, that oxidative stress responses are in general temperature sensitive (5, 38, 68). Possibly, the relatively high basic level of SOD in controls at 11 °C prevented a further induction by venlafaxine, which became obvious in fish exposed at 7 °C, with more pronounced responses in larvae than in juveniles. In larvae, this effect appeared in individuals exposed to at least 10 µg/L, with a maximum increase of 24% while, in juveniles, the maximum SOD-activity was 22% higher than in the control, however not significant. Along this line, effects of different antidepressants on oxidative stress were previously published. Elevated oxidative stress indicated by changes in catalase, peroxidase, glutathione-S-transferase (GST) and SOD-activity have been shown in meagre exposed to venlafaxine (41), zebrafish exposed to amitriptyline (73), goldfish exposed to sertraline (71) and daphnids exposed to citalopram (72). In a multi stressor experiment temperature effects on the anti-oxidant defence have been shown and potential deleterious effects of venlafaxine on the anti-oxidant defence mechanism of zebrafish were indicated (42). However, in one study, fluoxetine-exposed Pomatoschistus microps did not show a change in SOD activity (12). In general, both the vast majority of literature results and our own data indicate that venlafaxine and other antidepressants can induce oxidative stress in fish.

In both life stages of brown trout, neither citalopram nor venlafaxine did alter the AChE or CbE activity, thus, no neurotoxic effect of venlafaxine or citalopram on brown trout was detected. Nevertheless, effects of antidepressants on AChE have been reported before: decreased AChE activity could be detected in a marine clam species exposed to low µg/L fluoxetine (45), Pomatoschistus microps exposed to 1 mg/L fluoxetine (12) and Daphnia magna exposed to 1 mg/L citalopram (72). In contrast, goldfish exposed to low µg/L sertraline showed an increased AChE activity, which could not be detected at higher concentrations (71). Concerning the impact of antidepressants on carboxylesterases, to our knowledge, no data are published for fish so far.

Likewise, neither citalopram nor venlafaxine exposure induced a response of the Hsp70 system, thus, no proteotoxic effects became evident in juvenile brown trout. However, Maulvault et al. (2019) observed an increased heat shock response in meagre exposed to 20 µg/L venlafaxine. As Hsp70 is a rather unspecific
biomarker, which is induced by various types of stressors, differences might be attributed to different exposure conditions or to species-specificity.

Generally, temperature plays an important role in our scenarios, because it can have an impact on many physiological responses in fish (35, 38, 42, 68). We could see that fish exposed at 11 °C revealed an overall higher SOD-activity, however, differences between larvae exposed at 11 °C and 7 °C cannot solely be attributed to temperature effects, because of different exposure times. Furthermore, it could be shown that effects in larvae were more pronounced than in juveniles, however, these differences can also not solely be accredited to different sensitivities of the life stages but can also result from longer exposure times of larvae. A comparison of the presented biomarker data with previously published results obtained for endpoints like behaviour and growth impairment from the same experiments (74, 75) make evident, that the effect concentrations for behavioural and growth effects are in the same range as those for the biomarker responses shown here. The lowest observed effect concentration (LOEC) in citalopram-exposed brown trout was 100 µg/L reflected by histopathological as well as behavioural alterations. Similarly, in venlafaxine-exposed brown trout, behavioural and growth endpoints show similar LOECs of 10 µg/L as the biochemical biomarker SOD-activity. This demonstrates that citalopram and venlafaxine can induce effects on cellular and biochemical markers in equivalent concentrations as it affects growth and behavioural endpoints. As a result, the no observed effect concentrations (NOEC) in these experiments are just an order of magnitude above environmentally relevant concentrations and thus reasonably call for increased vigilance towards these pharmaceuticals in the environment.

Conclusion

Combining the results for histopathology, neurotoxicity, proteotoxicity and oxidative stress with previously published data, it became evident that effects and side-effects of citalopram and venlafaxine in brown trout are similar to those known for humans. LOECs were at 10 µg/L for venlafaxine and at 100 µg/L for citalopram, matching LOECs previously recorded for behavioural and growth endpoints. Although the NOECs for citalopram and venlafaxine are at least a decimal power higher than environmentally relevant concentrations, in consideration of safety factors and additive effects both antidepressants have necessarily to be considered in environmental risk assessment.

Declarations

Ethics approval

All animals were approved by the animal welfare committee of the Regional Council of Tübingen, Germany (ZO 2/16).

Consent for publication

Not applicable
Availability of data and materials

The datasets generated and/or analysed during the current study are available in the effectnet-seek repository, https://effectnet-seek.bioquant.uni-heidelberg.de/investigations/9

Competing interests

The authors declare that they have no competing interests.

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Authors’ contributions

Michael Ziegler conceived, designed and performed the experiments, analysed the data and prepared the figures and tables and wrote the first draft of the paper and edited according to the coauthors. Helene Eckstein, Shannon Ottmann, Lukas Reinelt and Sabine Stepinski analysed part of the Data and approved the final draft. Heinz-R. Köhler and Rita Triebskorn conceived and designed the experiments and contributed reagents/materials/analysis tools as well as reviewed the first draft of the paper and approved the final draft.

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