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Authors
Kupsco, Allison
Schlenk, Daniel

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Mechanisms of Selenomethionine Developmental Toxicity and the Impacts of Combined Hypersaline Conditions on Japanese Medaka (Oryzias latipes)

Allison Kupsco* and Daniel Schlenk

Department of Environmental Sciences, University of California- Riverside, Riverside, California 92507, United States

ABSTRACT: Selenium (Se) is an essential micronutrient that can cause embryotoxicity at levels 7–30 times above essential concentrations. Exposure to hypersaline conditions and 50 μM selenomethionine (SeMet) decreased embryo hatch and depleted glutathione in Japanese medaka embryos without affecting Se accumulation. To better understand the impacts of nonchemical stressors on developmental toxicity of Se in fish, several adverse outcome pathways were evaluated in the Japanese medaka (Oryzias latipes). We treated medaka embryos at 12 h post fertilization with 50 μM SeMet for 12 hours in freshwater or in 13 ppt hypersalinity and evaluated the contributions of oxidative stress, the unfolded protein response and apoptosis to reduced hatch. Exposure to SeMet and hypersalinity decreased embryo hatch to 3.7% ± 1.95, and induced teratogenesis in 100% ± 0 of hatched embryos. In contrast, treatments of freshwater, saltwater, and SeMet in freshwater resulted in 89.8% ± 3.91–86.7% ± 3.87 hatch, and no significant increase in deformities. We found no significant differences in lipid peroxidation, indicating that oxidative stress may not be responsible for the observed toxicity in embryos at this time point (24 h). Although significant changes in apoptosis were not observed, we witnessed up to 100 fold increases in transcripts of the endoplasmic reticulum (ER) chaperone, immunoglobulin binding protein (BiP) and trends toward increasing downstream signals, activating transcription factor 4 (ATF4) and ATF6 indicating potential contributions of the unfolded protein response to the effects of SeMet and hypersaline conditions. These data indicate that multiple adverse outcome pathways may be responsible for the developmental toxicity of Se and salinity, and these pathways may be time dependent.

I. INTRODUCTION

Selenium (Se) is an essential micronutrient; levels only 7–30 times greater than required can be toxic.1 This is a concern in aquatic environments, where anthropogenic activities can release large quantities of Se, and include agricultural runoff of irrigation waters in arid regions,2 waste rock from coal, phosphate, and uranium mining,3–5 and combustion waste from coal burning power plants.6 Se usually enters the waterways in its inorganic forms of selenate (Se+6) or selenite (Se+4), which can be taken up by microbes and primary producers and converted into various organic forms, including the amino acid, selenomethionine (SeMet).7 Consumers, such as fish and birds, are exposed to Se primarily in the diet and SeMet has been shown to be the major form of Se in the fish diet.8 One concern for SeMet toxicity is its bioaccumulation potential. SeMet has been demonstrated to move through the food chain by trophic transfer to higher-level organisms.8,10 Following a Se poisoning event at Belews lake, NC, Lemly found Se to have bioaccumulated from 519 times in periphyton to 3975 times in the visceral tissues of fish.10 This is of particular concern for oviparous carnivores, for which maternal offloading may impair development or reproductive success through respective teratogenesis or embryo lethality.10

Because of the potential for biomagnification, traditional water quality measurements of Se concentrations may be ineffective. Recently, the U.S. Environmental Protection Agency has begun advocating tissue concentration measurements for Se monitoring.11 However, even these measurements may not provide an accurate picture of Se effects on an ecosystem, because fish encounter multiple stressors in their environments, which can alter Se toxicity. Recent evidence suggests that hypersalinity may compound Se toxicity.12 This is of particular importance in areas such as the San Joaquin River Valley, CA, and the San Francisco Bay Delta area, where many historically freshwater–waterways are in danger of salinization.13 These areas are often spawning grounds for protected species such as the endangered delta smelt (Hypomesus transpacificus) and threatened steelhead trout (Oncorhynchus mykiss).

The mechanisms behind Se induced teratogenesis and mortality remain unclear. Several studies point to oxidative...
stress as one mode of action for Se toxicity.\textsuperscript{14–17} However, oxidative stress is most likely only one factor influencing SeMet toxicity. The unfolded protein response (UPR) is a cellular and molecular response to perturbations in endoplasmic reticulum (ER) homeostasis (See Hetz (2012) for review\textsuperscript{18}). Oxidative molecular response to perturbations in endoplasmic reticulum toxicity. The unfolded protein response (UPR) is a cellular and oxidative stress is most likely only one factor in.

Table 1. Primers, Accession Numbers, and Concentrations used for qRT-PCR

| name       | fwd primer (5′-3′)                     | rev primer (5′-3′)                     | accession #     | conc.  |
|------------|----------------------------------------|----------------------------------------|-----------------|--------|
| EF1-a      | CTACATCAAGAGATCCGCTACA               | CGACAGGGACAGGTCACAATAC                 | NM_001104662.1  | 2.5 μM |
| CASP3A     | CCCATTCGACGTACACGCAA                   | AGCCGAGGACAGGCAATCA                    | NM_001104670.1  | 5 μM  |
| BAX        | GCTGCTGCTAAGGCTCTAC                   | CCAGATTGCTGACGCTGAA                    | NM_131562.2     | 2.5 μM |
| BiP        | GAGGATACTCTGGAGCTGAC                  | GTGACAGTGGCAGGGTTAC                    | NM_001278801.1  | 0.5 μM |
| ATF6       | CAAGCCACCTCCAGCTGATC                  | GCGGACTCTGCTGTTATG                     | NM_001278901.1  | 0.5 μM |
| ATF4       | CTGAGGGTGAGCTGCTATG                   | TGGAGGGACTGCTGCAAA                     | XM_004066069.1  | 2.5 μM |

II. MATERIALS AND METHODS

2.1. Chemicals and Reagents. Seleno-L-methionine (Purity 98%), 1-butanol, phosphoric acid, thiobarbituric acid and all other reagents were purchased from Sigma-Aldrich (St. Louis, MO). A Milli-Q water purification system (Millipore, Billerica, MA) was used to obtain deionized water. Ethanol (Fisher, Pittsburgh, PA) was of molecular biology grade.

2.2. Embryo Collection and Exposure. Japanese Medaka were cultured at the University of California- Riverside and housed in a 2:3 ratio of males to females in medium-hard water at 27 °C and a photoperiod of 14 h light and 10 h dark. Adults were fed twice daily a diet of live brine shrimp. Embryos were collected 0–1 h following fertilization. Viable embryos were determined based on oil droplet migration to the vegetal pole as outlined by Kirchen and West (1976).\textsuperscript{19} Nonviable embryos were discarded and viable embryos were placed into 60 × 15 mm Petri dishes containing either freshwater or a makeup of saltwater from the San Joaquin River Valley (20–30 embryos per replicate, and 5–10 replicates per group). Although, waterborne SeMet exposures do not represent the most likely environmental exposure (the primary exposures for SeMet are dietary or via maternal transfer), they are sufficient to study the mechanistic effects of SeMet on medaka embryos. San Joaquin River saltwater was prepared in the lab according to a recipe from Westlands Water District located about 10 km south of Mendota in the San Joaquin River Drainage Basin, CA.\textsuperscript{20} Salinity was measured with a conductivity meter, and corresponds to about 13‰ and 15.3 g/L suspended solids.

Following 12 h of equilibration in fresh or salt water, the replicates were divided into three groups. The first subset of embryos were frozen −80 °C to represent a time zero control (12hpf). Other embryos were treated with a 50 μM solution of Seleno-L-Methionine (Cat. No. S3132, Sigma-Aldrich) in freshwater or saltwater and exposed for 12 h, then collected and frozen at −80 °C for analysis (24hpf). SeMet concentrations were chosen based on previous research and were intended to represent the upper levels of bioaccumulation measured in embryos.\textsuperscript{12} Previous studies have also demonstrated uptake of SeMet into the embryo, indicating this system was an effective exposure method.\textsuperscript{12} The final subset of embryos were left in freshwater or saltwater for 24 h to compare to the SeMet treated 12hpf samples.

2.3. Modified Embryo Larval Toxicity Assay. The medaka embryo-larval toxicity assay was adapted from Farwell et al. (2006).\textsuperscript{21} Embryos were collected and treated as above with one replicate equal to 15–20 embryos per dish. Following 12 h of SeMet treatment, embryos were rinsed and transferred to new dishes containing freshwater or saltwater. Water was changed every other day and dishes were monitored for mortality with removal of dead embryos. Embryo hatch was monitored for 21 days post fertilization. At hatch, embryos were assessed for deformities and terminated immediately. Percent hatch and percent of hatched embryos with deformities were recorded.

2.4. Analysis of Gene Expression. Total mRNA was isolated from embryos using the Lipid Tissue RNeasy kit (Qiagen, Valencia, CA) following the manufacturers instructions. mRNA quantity and quality was measured using the ND-1000 (NanoDrop, Wilmington, DE). mRNA (1 μg) was converted to cDNA using the Reverse Transcription System (Promega Corporation, Madison, WI), according to the manufacturers instructions.

Primers were designed using IDTDNA PrimerQuest software and optimized using PCR Miner\textsuperscript{22} (Table 1). As no BAX gene for Japanese medaka has been annotated in the NCBI database, BLAST was used on the medaka genome (http://compbio.dfci.harvard.edu/cgi-bin/tgi/Blast/index.cgi) against BAX from zebrafish (Danio rerio) to develop primers.
Similarity between the genes had an E value of $1 \times 10^{-48}$. EF1α was run as a housekeeping gene. qPCR was performed with the iScript One-step RT-PCR kit with SYBR Green from Bio-Rad (Hercules, CA), omitting the reverse transcriptase, on a MyiQ5 Thermo cycler (Biorad). The samples were denatured and the polymerase activated at 95 °C for 5 min, then 40 cycles of 10s at 95 °C and 30s of 55 °C. Samples were subject to melting curve analysis from 65 to 85 °C in 0.5 °C increments with continuous fluorescence measurement. qPCR was analyzed according to Schmittgen and Livak23 and fold change was calculated against the 12hpf freshwater controls. All data was compared against the 12hpf controls in order represent how the gene expression changed over the 12h time period, and how the treatments affected this change. Rather than observing a discrete point in development, we feel it is necessary to understand how these treatments altered normal development.

2.5. Analysis of Oxidative Stress. Thiobarbituric Reactive Substances (TBARS) were measured to estimate malondialdehyde (MDA) formation in medaka embryos.24 Embryos (15–20) were weighed and homogenized in 1.15% KCl then centrifuged at 3000 rpm for 5 min at 4 °C. The supernatant was then used in the assay. Samples were run in duplicate on a Wallac Victor2 multilabel plate reader (PerkinElmer, Waltham, MA) with excitation at 535 nm and emission at 585 nm.

2.6. Statistical Analysis. Statistical significance was assessed using a student’s T-Test or 2-way ANOVA in the statistical program R. Statistical significance was determined at $p \leq 0.05$, unless otherwise noted. If overall significance was determined following two-way ANOVA, Tukey’s HSD test was performed posthoc. Data was checked for normality and homogeneity of variances. Any non-normal data was log transformed. For data that remained non-normal following log transformation, Kruskal–Wallis tests were performed with Dunn’s test posthoc.

III. RESULTS

3.1. Embryo-Larval Toxicity Assay. While there was no difference in hatch between saltwater and freshwater control embryos, 50 μM SeMet significantly decreased embryo hatch in saltwater treated embryos to 3.7% (Figure 1A). However, SeMet treatment in freshwater did not significantly decrease hatch. The median day to hatch was not significantly affected by SeMet treatment in freshwater or saltwater (Figure 1B). SeMet and hypersaline treatment also significantly increased the deformities in treated embryos (Figure 1C). All SeMet and hypersaline treated embryos had deformities upon hatch. The most common deformities observed were kyphosis, lordosis, craniofacial abnormalities, and yolk sac edema (Figure 1D).

3.2. Oxidative Stress and Apoptosis. There was no significant difference in amount of lipid peroxidation between any of the treatments and no difference between embryos at 12hpf and 24hpf (Figure 2). BAX transcript levels decreased significantly from 12hpf to 24hpf (0.2 fold, $p = 0.011$), while Caspase 3A levels remained constant (Figure 3A). There was no significant difference in BAX or CASP3A gene expression following treatment with hypersalinity or SeMet (Figure 3B and C).

3.4. Gene Expression of UPR Mediators. UPR gene expression changed from 12hpf to 24hpf in medaka embryos. ATF6 gene expression decreased significantly from 12hpf to 24hpf (0.3 fold, $p = 0.019$), while BiP and ATF4 expression increased significantly (2 fold, $p = 0.048$, and 4 fold, $p = 0.008$, respectively). BiP mRNA was increased in both saltwater and freshwater SeMet treatments up to 39 fold over 12 h controls and 107 fold over 12 h controls in the SeMet freshwater and
SeMet saltwater treatments, respectively. Though there was a trend toward an increase in ATF6 expression with SeMet treatment ($p = 0.119$), SeMet and hypersalinity did not significantly alter ATF6 expression. Similarly, trends in ATF4 expression indicated a potential difference between freshwater and saltwater treatment ($p = 0.07$), with SeMet decreasing ATF4 in freshwater, yet increasing it in saltwater.

**IV. DISCUSSION**

The mechanism of action of SeMet toxicity is not well understood, particularly in the presence of multiple stressors, which may confound regulatory monitoring. We observed a decrease in hatch following treatment with hypersalinity and SeMet, while surviving embryos had deformities. Previous work at this developmental stage and SeMet concentration reported a significant decrease in embryo hatch following 24 h of 50 $\mu$M SeMet treatment in freshwater.$^{12}$ While we did not observe this trend, our results were expected considering the duration of SeMet treatment was half as that previously studied. Overall, our results are consistent with a plethora of other data reporting SeMet’s lethal and teratogenic effects in the field, for example, ref 10.

Japanese medaka are a euryhaline species; adults are able to spawn and embryos hatch in full seawater.$^{25}$ We observed no significant difference in toxicity between freshwater and hypersaline controls, indicating that our results are not due to osmotic stress alone. Others have found salinity to potentiate SeMet toxicity in embryos.$^{12}$ The mechanism behind this remains to be elucidated, however, FMO may play a role. FMOs have been shown to oxygenate SeMet, which may contribute to its toxicity.$^{26,12}$ Several studies have found FMO activity can increase under hypersaline conditions.$^{12,27,28}$ This increased FMO activity may increase SeMet oxygenation, which in turn may increase its embryo toxicity.

In contrast, studies have shown that SeMet activation can occur following methioninase generation of methylselenol,$^{29}$ which can subsequently generate oxidative stress in rainbow trout embryos.$^{14}$ Methylselenol has also been implicated in induction of caspase-mediated apoptosis in cancer cell lines.$^{30}$ However, considering that neither oxidative stress nor apoptosis was induced in SeMet and hypersaline treated embryos and that FMO has been found to be induced by hypersaline conditions, we conclude that FMO activation may be a major contributor to the observed toxicity.

Of interest is that SeMet and hypersalinity did not generate oxidative stress as measured by lipid peroxidation after 12 h of treatment. However, several groups, including ours, have identified oxidative stress as one of the main modes of action of SeMet toxicity.$^{12-15}$ As lipid peroxidation is an end point for severe oxidative stress, TBARS is not sensitive to small changes in cellular redox.$^{31}$ Furthermore, in this study, TBARS was measured in whole embryos and did not consider localized...
effects. Thus, oxidative stress may still be occurring in SeMet and hypersaline treatments, yet it may not be detected by our assays or was not as high as that observed after 24 hpf. While our results do not eliminate oxidative stress as a mechanism of SeMet induced embryotoxicity, they indicate that other processes may play an important role particularly at the 12 hour time point of exposure. While we observed no difference in whole-embryo lipid peroxidation between 12 hpf and 24 hpf, many studies demonstrate that the redox status of embryos also undergoes great changes throughout development. Two contradicting reports demonstrate the changes in redox status of medaka embryos throughout development. Wu et al. measured changes in oxidative stress in the whole embryo each day post fertilization using Dichloro-dihydro-fluorescein diacetate (DCHFDA; a dye that fluoresces following oxidation) and found overall reactive oxygen species (ROS) increased gradually until hatch. In contrast, another group studying medaka development and silver nanoparticle toxicity, found total ROS decreased throughout development. However, in addition to total ROS, both of these studies examined multiple biomarkers for oxidative stress and found no common patterns between them. Hence, the current studies on redox status throughout medaka development confirm that further studies are necessary in order to understand this complex process.

Another adverse outcome pathway that may contribute to SeMet toxicity in medaka at this early life stage is the UPR. Methylselenic acid (MSA) induced the UPR in PC-3 cells, a human prostate cancer cell line and we observed increases in Methylselenic acid (MSA) induced the UPR in PC-3 cells, a SeMet toxicity in medaka at this early life stage is the UPR. BiP is necessary in order to understand this complex process.

The UPR has a different response to DTT. The comparison of apoptosis taken at 12 hpf and 24 hpf clearly shows that these important processes fluctuate greatly during development. Indeed, apoptosis has been shown to fluctuate in tissue-specific patterns throughout Japanese medaka development, occurring mostly in the head, spinal column, and tailbud. Furthermore, apoptosis plays a key role in neural development.

The overwhelming evidence for changes in apoptosis, oxidative stress, and the UPR throughout development, indicates that developing Japanese medaka embryos have important windows of susceptibility to SeMet and hypersaline stress. Thus, while the roles of oxidative stress and apoptosis in the developmental toxicity of SeMet may be limited from 12 to 24 h, they may be increased as the oxidation state of the embryos increases, or as apoptosis is more active at later stages of development. It is important to map these processes throughout embryogenesis, so that we may better understand the developmental toxicity of SeMet and hypersaline conditions.

In summary, hypersaline conditions derived from the San Joaquin River Valley, CA, enhanced the toxicity of SeMet in the developing medaka embryo. While the UPR may have played a role, oxidative stress and apoptosis measured in the whole embryo were not associated with SeMet induced mortality and teratogenesis at this early stage. Additional studies will further consider the role of oxidative stress and the UPR throughout...
medaka development and investigate developmental periods most susceptible to SeMet and hypersaline toxicity.

■ AUTHOR INFORMATION

Corresponding Author
*Phone: 520-240-7893; fax: 951-827-3993; e-mail: allison.kupsco@email.ucr.edu.

Notes
The authors declare no competing financial interest.

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Figure 4. Effects of combined exposure of SeMet (50 μM) and hypersaline conditions on BiP, ATF6, and ATF4 transcripts in Japanese medaka embryos after 12 and 24 hpf. (A) Expression of BiP, ATF6, and ATF4 in whole embryos at 12hpf and 24hpf in freshwater. Expression of (B) BiP, (C) ATF6, and (D) ATF4 in 48hpf control embryos and SeMet treated embryos in freshwater and saltwater. Each value represents the mean ± SE of 5–10 replicates. EF1-α was run as a housekeeping gene. Statistical significance is indicated by differing letters (Two-way ANOVA, Tukey HSD Test p < 0.05).
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