Report

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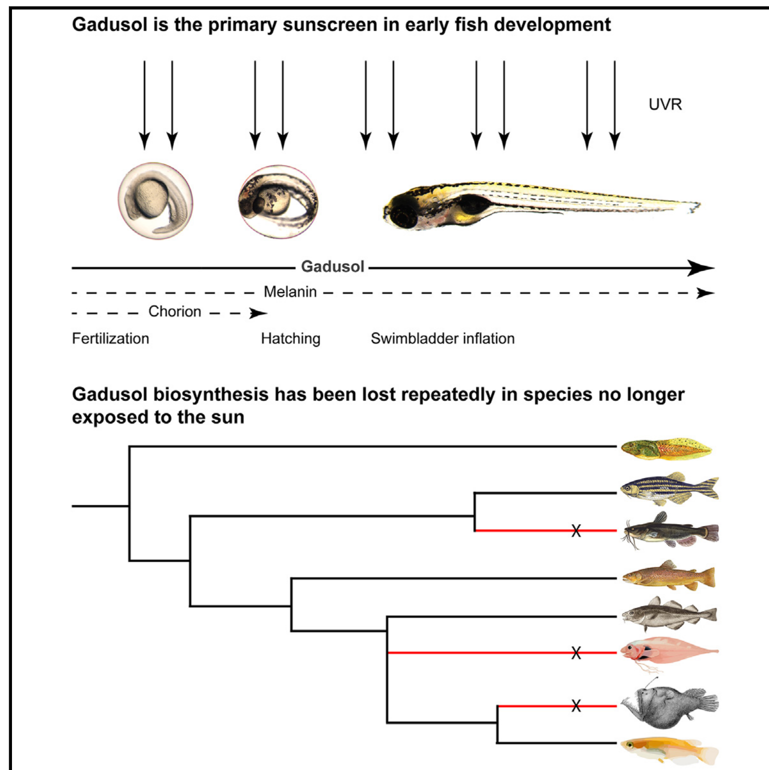
Gadusol is a maternally provided sunscreen that protects fish embryos from DNA damage

Graphical abstract



Highlights

- Gadusol is a powerful sunscreen that protects early stages of fish development
- Gadusol is produced by the mother and deposited into the egg
- Gadusol is a more efficient sunscreen during larval stages than melanin
- Gadusol synthesis has been lost in many species whose young are not exposed to UVR

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In brief

Sunscreens have evolved to mitigate ultraviolet radiation (UVR)-induced stress. Here Rice et al. show that a maternally provided transparent compound called gadusol is a powerful sunscreen that protects fish embryos. They find that gadusol synthesis genes have been repeatedly lost in fish species whose young are not exposed to UVR.

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Gadusol is a maternally provided sunscreen that protects fish embryos from DNA damage

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SUMMARY

Exposure to ultraviolet radiation (UVR) is harmful to living cells, leading organisms to evolve protective mechanisms against UVR-induced cellular damage and stress.[1,2] UVR, particularly UVB (280–320 nm), can damage proteins and DNA, leading to errors during DNA repair and replication. Excessive UVR can induce cellular death. Aquatic organisms face risk of UV exposure as biologically harmful levels of UVB can penetrate >10 m in clear water.[3] While melanin is the only known sunscreen in vertebrates, it often emerges late in embryonic development, rendering embryos of many species vulnerable during the earlier stages. Algae and microbes produce a class of sunscreening compounds known as mycosporine-like amino acids (MAAs).[4] Fish eggs contain a similar compound called gadusol, whose role as a sunscreen has yet to be tested despite its discovery over 40 years ago.[5] The recent finding that many vertebrate genomes contain a biosynthetic pathway for gadusol suggests that many fish may produce and use this molecule as a sunscreen.[6] We generated a gadusol-deficient mutant zebrafish to investigate the role of gadusol in protecting fish embryos and larvae from UVR. Our results demonstrate that maternally provided gadusol is the primary sunscreen in embryonic and larval development, while melanin provides modest secondary protection. The gadusol biosynthetic pathway is retained in the vast majority of teleost genomes but is repeatedly lost in species whose young are no longer exposed to UVR. Our data demonstrate that gadusol is a maternally provided sunscreen that is critical for early-life survival in the most species-rich branch of the vertebrate phylogeny.

RESULTS

Gadusol is maternally provided and protects embryos and larvae from UVR

To test if gadusol is a sunscreen in vertebrate embryos, we used CRISPR-Cas9 to delete most of exon 2 of zebrafish *eevs*, which encodes the enzyme essential for the first step in gadusol biosynthesis (Figures 1A, S1A, and S1C; Data S1). We chose zebrafish for these experiments because they live and spawn in shallow sunlit waters, they are known to produce gadusol,[6] and they are genetically tractable. Grown in our animal facility, where they are protected from UVR, homozygous *eevs* mutant females and males survived to fertile adulthood like their wild-type peers. Using reciprocal crosses between homozygous mutant adults ($eevs^{-/-}$) and wild-type adults ($eevs^{+/+}$), we generated heterozygous mutant embryos that lack maternal contribution of gadusol (hereafter referred to as M*eevs*) and heterozygous mutant embryos that retain this maternal contribution (referred to as $eevs^{+/-}$) (Figure 1B). Notably, M*eevs* and $eevs^{+/-}$ embryos have identical genotypes but either lack or possess maternally provided gadusol, as judged by mass spectrometry (Figure 1B) and UV spectrophotometry (Figure S1C). We generated maternal-zygotic homozygous mutant embryos (referred to as MZ*eevs*) from incrosses of homozygous mutant parents. Immediately after fertilization, gadusol was nearly absent in MZ*eevs* embryos and indistinguishable from M*eevs* (Figures 1C and S1C). We next asked how long maternally provided gadusol persisted in embryos and larvae. We compared gadusol abundances from whole embryos and larvae with the following genotypes: $eevs^{+/+}$ (wild type), M*eevs*, and MZ*eevs*. We found only a modest increase in gadusol abundance in M*eevs* relative to MZ*eevs* at 5 days post-fertilization (dpf) (Figure 1C). Together with transcriptomic data that show *eevs* mRNA is only present during early stages of oogenesis[7] (Figure S1B) and absent from embryos,[6,8] our data suggest that maternally synthesized and deposited gadusol is the source of nearly all gadusol in the developing zebrafish. This is an example of a maternal effect, where disruption of the *eevs* gene in mothers eliminates deposition of gadusol presence in their embryos, regardless of embryo genotype.

To determine if gadusol protects zebrafish embryos against UVB, we developed an assay to deliver precise doses of UVB to embryos and measure the effect on swim bladder inflation at 5 dpf (a hallmark of healthy development essential for survival; Figures S2A–S2D). We found that 450 J/m^2 of UVB

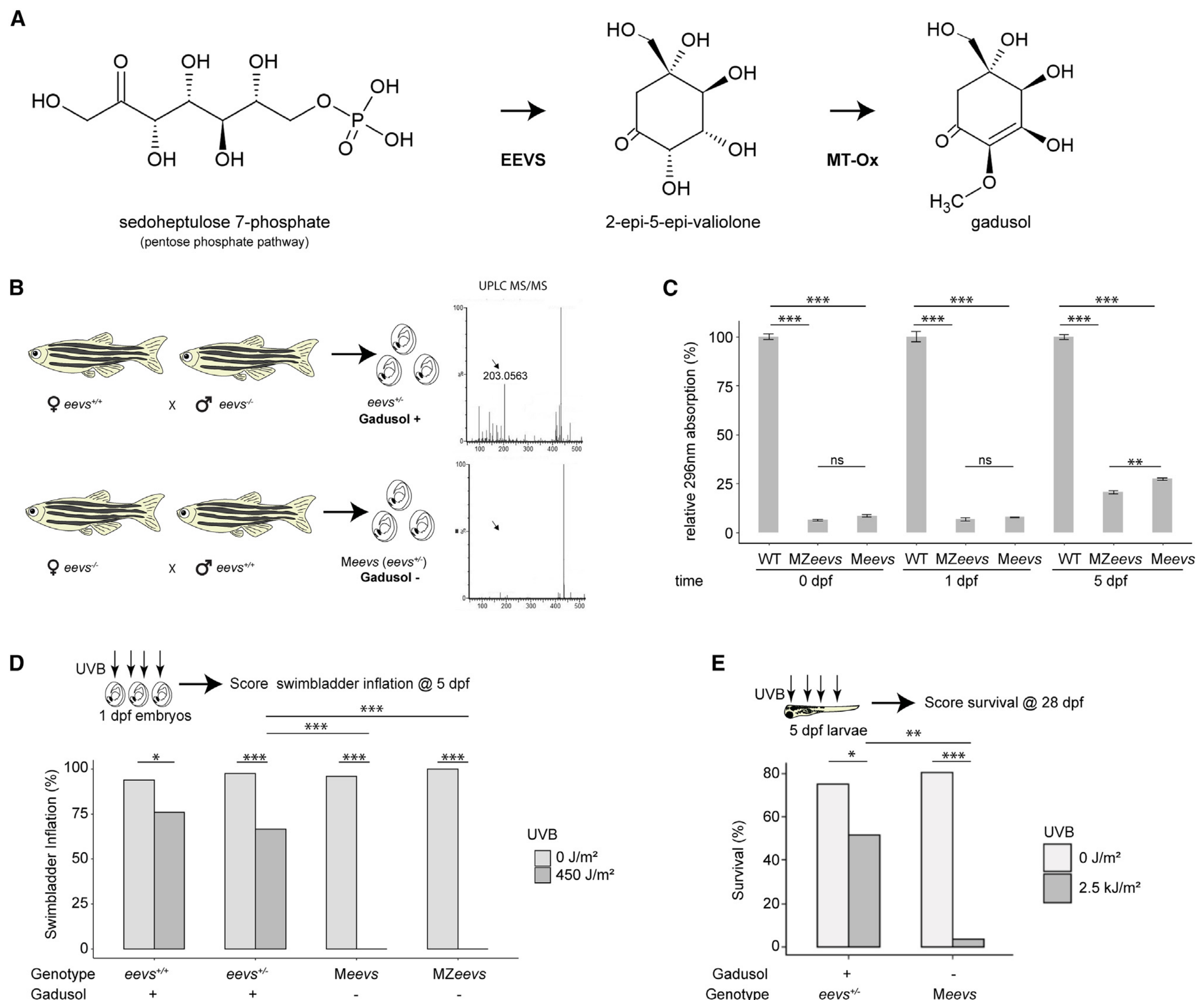

Figure 1. Gadusol is maternally provided and protects zebrafish embryos and larvae from UVB

(A) The biosynthetic pathway for gadusol production.

(B) Experimental diagram for generating heterozygous mutant *eevs*$^{+/-}$ embryos and larvae with identical genotypes but containing maternal contribution of gadusol (top) or depleted of maternally provided gadusol (bottom). On the right, UPLC-MS of 0 hpf egg extracts from each genetic cross; arrow indicates gadusol mass.

(C) Absorption values at 296 nm from the indicated genotypes at the indicated time points. All absorption values normalized to wild type. Error bars indicate SD from biological replicates.

(D) Distribution of swimbladder inflation scored in 5 dpf larvae, with genotypes and gadusol presence indicated, after mock exposure (gray) or UVB exposure (dark gray) at 24 hpf stage. All embryos resulted from crosses between TU and AB strain parents, except the TU in-cross that generated MZ*eevs* embryos. From left to right: n = 50, 50, 75, 75, 100, 97, 50, 50; C = 2, 2, 3, 3, 4, 4, 2, 2.

(E) Survival distribution scored at 28 dpf, with genotypes and gadusol presence indicated, after mock exposure (gray) or UVB exposure (dark gray) at 5 dpf. From left to right: n = 100, 95, 100, 97; C = 4 for all groups.

n = number of embryos/larvae. C = number of clutches. Student's t test (C), Fisher's exact t test (D and E); *p < 0.05; **p < 0.01; ***p < 0.0001.

See also Figures S1 and S2 and Data S1 and S3.

(fluence rate: 2.5 W/m^2; STAR Methods) delivered at 24 h post-fertilization (hpf) resulted in ~75% swim bladder inflation in wild-type and *eevs*$^{+/-}$ embryos, but did not result in gross developmental defects (Figures 1D and S2C). In stark contrast, MZ*eevs* and M*eevs* embryos were extremely vulnerable to the same dose of UVB; all embryos failed to inflate their swim bladders (Figure 1D).

Since zygotic production of gadusol was still minimal at 5 dpf (Figure 1C), we hypothesized that larvae lacking maternal gadusol should be highly sensitive to UVB at this later stage. We repeated UVB dosage curves on 5 dpf larvae and identified 2.5 kJ/m^2 for a significant impact on wild-type larvae survival (Figure S2E). We grew UV-exposed and control larvae in our fish facility nursery to 28 dpf, which requires developing animals

to forage for food to survive. We found that only 2% of exposed M*eevs* larvae survived, compared to ~50% of controls exposed to the same dose of UVB (Figure 1E). Together, these data demonstrate that maternally provided gadusol provides powerful UVB protection to early embryos and older larvae.

Gadusol prevents DNA damage and apoptosis

Next, we sought to understand the mechanism by which gadusol protects embryos from UVB. In other species, gadusol and related molecules were hypothesized to function as antioxidants as well as sunscreens.[5,6,9] To test if gadusol serves as an antioxidant in zebrafish embryos, we exposed 24 hpf embryos to hydrogen peroxide to induce oxidative stress. At 5 dpf, gadusol-depleted M*eevs* and control $eevs^{+/-}$ embryos had similar responses to oxidative stress, suggesting that gadusol does not function as an antioxidant *in vivo* (Figures S3A and S3B).

To test if gadusol serves as a sunscreen by absorbing UVB, we measured the production of cyclobutane pyrimidine dimers (CPDs), a signature of UVB-induced DNA damage.[10] If gadusol acts as a sunscreen, then it would absorb UVB photons and shield the underlying DNA from CPD formation. We exposed 24 hpf embryos to UVB and used immunohistochemistry to detect CPDs and quantify fluorescence intensity. Embryos that lacked gadusol had significantly higher levels of CPD formation after UVB exposure compared to controls containing gadusol (Figures 2A and 2B). CPDs are cytotoxic and induce apoptosis at high abundance. We used immunohistochemistry to detect a fast-acting apoptotic marker (activated caspase-3) in embryos exposed to UVB[11] (Figures 2C and S3C). We found that embryos lacking gadusol had increased levels of apoptotic nuclei, relative to controls (Figure 2C), supporting a role for gadusol in absorbing UVB and preventing DNA damage.

To characterize transcriptional responses to UVR in the absence of gadusol, we performed RNA sequencing (RNA-seq) comparing gadusol-depleted M*eevs* and wild-type embryos. Five hours after exposure to UVB, embryos lacking gadusol had significantly higher expression of many key stress response genes (*tp53*, *gadd45aa*, *ddb2*, and *cdkn1a*) relative to UVB-treated controls (Figure 2D). GO terms enriched in UV-exposed gadusol-depleted embryos included response to UV, response to DNA damage, response to light, and other stress response terms (Figure S3D; Table S1). Several of these genes were also modestly induced by visible light (Figure S3E), consistent with previous reports,[13,14] but their induction was similar between M*eevs* and wild-type embryos. Together, our imaging and gene expression data confirm that gadusol in zebrafish embryos acts as a true sunscreen to provide efficient protection against UV-induced DNA damage, cellular stress, and cell death.

Gadusol is the primary sunscreen in early fish development

In light of our finding that gadusol acts as a sunscreen, we compared the relative sunscreening potency of gadusol with that of other potential UV-blocking/absorbing mechanisms in larval zebrafish. Melanin is a well-known sunscreen in many organisms including humans. In zebrafish, melanophores become pigmented around 36 hpf, ultimately forming stripes that partially cover the larval brain and body, a pattern that is stable until ~14 dpf.[15,16] Melanophores protect the hematopoietic niche in larval zebrafish,[17] but their role as a whole-body sunscreen remains untested. The *nacre/mitfa* mutant disrupts a key melanophore master regulator and lacks melanophores. We generated two groups of larvae, each with pigmented and unpigmented siblings. One group contained no maternal gadusol, while the other group contained gadusol (Figure 3A). We treated all 5 dpf larvae with 2.5 kJ/m^2 of UVB and assessed survival in the nursery at 28 dpf. Larvae with gadusol were highly resistant to UVB stress, regardless of pigmentation status (Figure 3B). All larvae that lacked gadusol were highly sensitive to UVB, and larvae that lacked both gadusol and melanin were slightly more sensitive to UVB than their pigmented siblings. At a lower UVB dose (1.5 kJ/m^2), we also found a modest but significant effect of melanophores in protecting against UVB (Figure S3F). We conclude that while melanin plays a minor role in UVR protection, gadusol is the primary sunscreen in early fish development.

Another potential UV-protective mechanism is the chorion, the nearly transparent eggshell that contains perivitelline fluid and the embryo from fertilization until 2–3 dpf. We tested the sunscreening role of the chorion by mechanically removing it with forceps and exposing these embryos, and sibling controls that retained the chorion, to 450 J/m^2 of UVB at 24 hpf. We found that the chorion provides significant protection from UVB as ~60% of dechorionated embryos failed to inflate their swim bladders, significantly less than sibling controls (Figure 3C). We examined if gadusol was present in the chorion or in the perivitelline fluid within the chorion but found little to none (Figure S1D). These results suggest that the chorion structure itself can shield some incoming UVB. However, we conclude that the chorion provides less UV protection than gadusol, as gadusol-depleted embryos—even with intact chorions—all failed to inflate their swim bladders when challenged with the same dose of UVB (Figure 1D).

Together, our findings support a model in which embryonic and larval fish are protected by multiple layers of UVB protection that span early development (Figure 3D). The egg is maternally loaded with gadusol, which provides the primary and most important layer of UV protection from fertilization until at least 5 dpf. The chorion and melanophores are secondary, and less effective, means of UVR protection. The chorion protects the developing embryo between fertilization and hatching (2–3 dpf), when pigmented melanophores emerge and modestly protect the growing larval fish.

Gadusol has been repeatedly lost in fish species whose embryos are no longer exposed to sunlight

The two-enzyme biosynthetic pathway necessary for gadusol production (Eevs and MT-Ox) is encoded in numerous vertebrate genomes, including fish, birds, reptiles, and amphibians.[6] Osborn et al. identified the loss of the gadusol biosynthetic pathway in the coelacanth genome and suggested the loss might be attributable to lack of UV penetration in the deep-sea habitat of this species.[6] To test for broader patterns of conservation and loss among fish, we surveyed additional genomes, including many species that live in habitats not exposed to UVR. We hypothesized that gadusol synthesis genes would not be required in species that live in deep waters, caves, are live bearers, or use electroreception to navigate habitats with poor light penetrance.[18] To test this hypothesis, we searched 136 teleost

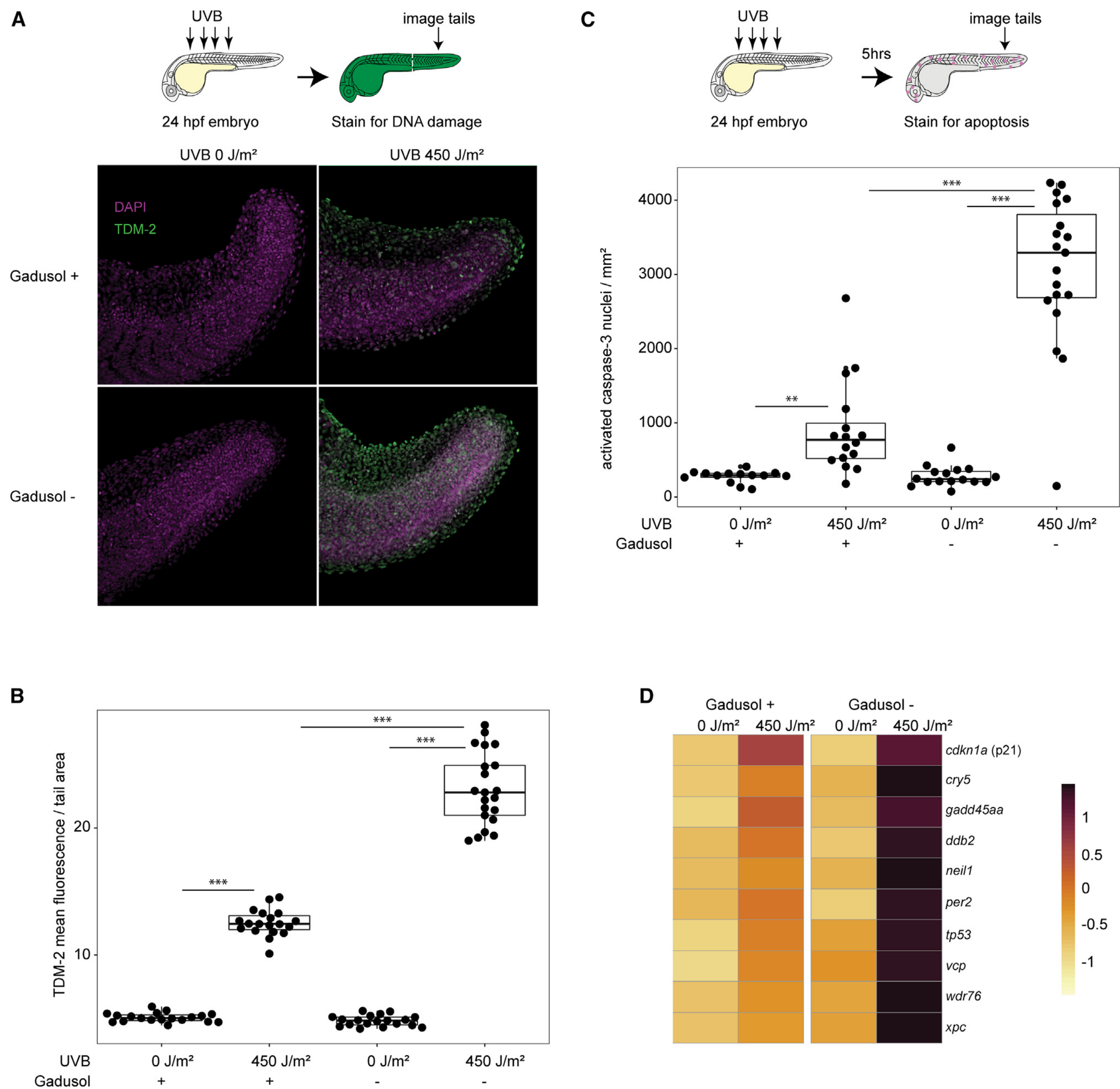

Figure 2. Gadusol functions as a sunscreen preventing DNA damage and apoptosis

(A) Immunohistochemistry, using an antibody that recognizes CPDs (TDM-2), on 24 hpf embryos immediately after mock or UVB exposure. Representative images shown.

(B) Quantification of CPD labeling normalized to tail area (mm^2). From left to right: n = 19, 19, 19, 21; C = 2 for all groups.

(C), Quantification of immunohistochemistry, using an antibody that recognizes activated caspase-3. n = 14, 16, 16, 20; C = 2 for all groups.

(D) Significant upregulation of select UVR response and DNA damage GO term-associated genes measured from the indicated conditions and genotypes using RNA-seq on 24 hpf embryos after mock exposure or UVB exposure. RNA was collected 5 h post mock or UVB exposure. Gene expression is scaled by rows. Significance determined via Fishenricher.[12]

Student's t test; *p < 0.05; **p < 0.01; ***p < 0.0001. n = number of embryos. C = number of clutches.

See also Figure S3, Data S3, and Table S1.

genomes for inactivation or loss of either *eevs* or MT-Ox. In all species, we identified a syntenic genomic region demarcated by highly conserved flanking genes and assessed the presence or absence of intact open reading frames (ORFs) encoding functional copies of *eevs* and MT-Ox. Our approach largely confirmed that the vast majority of teleosts have functional copies of *eevs* and MT-Ox.[6] However, our survey identified 16 independent losses of either the *eevs* or MT-Ox genes across the teleost phylogeny (Figure 4A, red species). Most of these genomes had lost orthologs of both *eevs* and MT-Ox, while others

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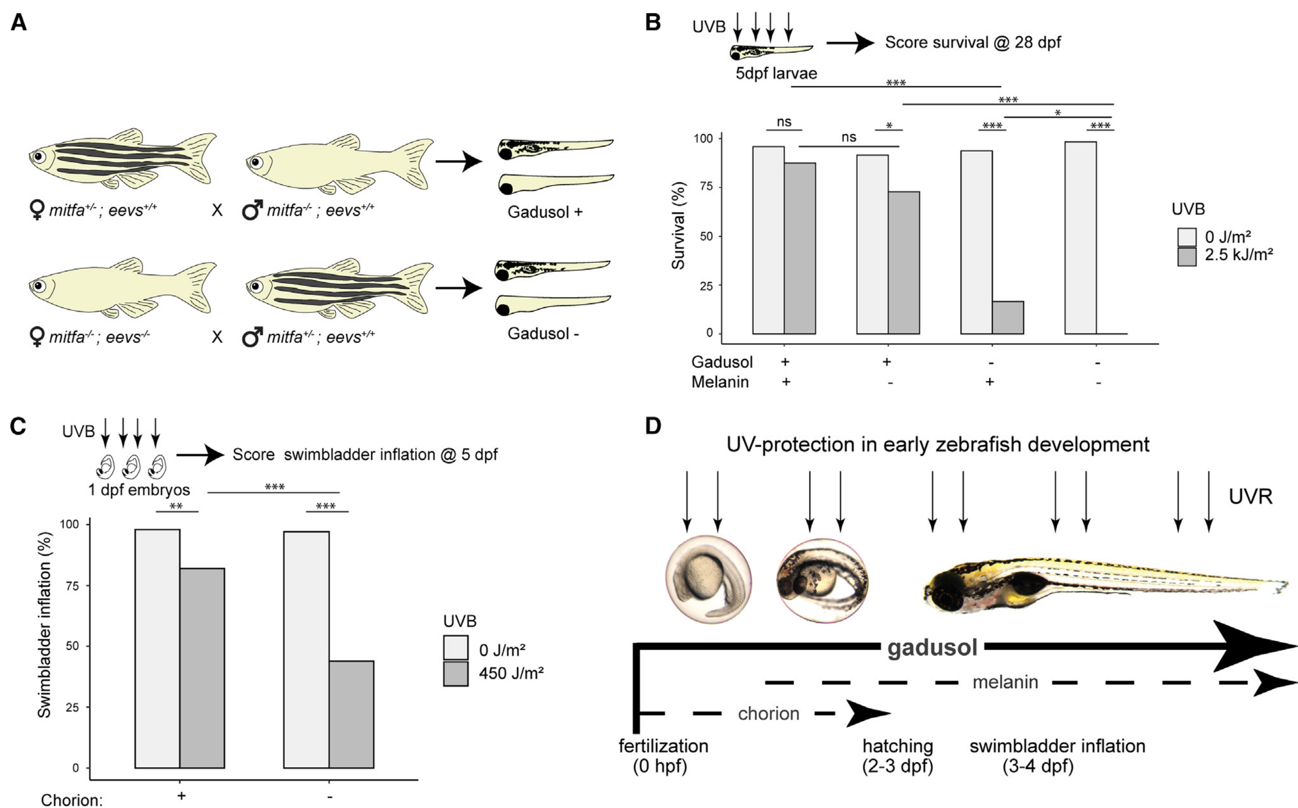

Figure 3. Melanin and the chorion serve as secondary UV-shielding mechanisms in embryonic and larval fish

(A) Experimental diagram for generating embryos that lack either melanin, maternally provided gadusol, or both.

(B) Survival distribution scored at 28 dpf, with presence of melanin and gadusol indicated, after mock exposure (gray) or UVB exposure (dark gray) at 5 dpf. From left to right: n = 48, 48, 48, 48, 48, 36, 60, 36; C = 2 for each group.

(C) Distribution of swimbladder inflation scored in 5 dpf larvae after mock exposure (gray) or UVB exposure (dark gray) at 24 hpf stage, with or without chorions. n = 100 for each group; C = 3 for each group.

(D) Model illustrating the relative importance and timing of multiple UV-shielding mechanisms used in early zebrafish development.

Fisher's exact t test; *p < 0.05; **p < 0.01; ***p < 0.0001. n = total number of individual embryos/larvae. C = total number of clutches.

See also Figure S1 and Data S3.

had lost only one gene or had pseudogene remnants (Figure 4B). The loss of genes involved in gadusol production was significantly correlated with lifestyle traits that identified species that live or spawn in habitats protected from the sun ($p = 0.012$) (Figures 4 and S4; Data S2). To corroborate the link between loss of *eevs* or MT-Ox and loss of gadusol, we measured gadusol levels in medaka embryos, which have intact *eevs* and MT-Ox genes, and ovaries of channel catfish, which have lost *eevs* and MT-Ox. We found a strict correlation between the presence of intact genes and maternally provided gadusol (Figure S1E). We conclude that gadusol production has been repeatedly lost during evolution in teleost species whose lifestyles protect them from UVR.

DISCUSSION

Plants and microorganisms use numerous UV-absorbing compounds as sunscreens.[2,4] However, other than melanin, the repertoire of vertebrate sunscreens—especially compounds that protect the most vulnerable early stages of development—remains essentially unknown. Here, we provide experimental and phylogenomic evidence that gadusol is an ancient sunscreen essential for protecting fish embryos from UVR. First, we use a CRISPR mutant that disrupts gadusol biosynthesis to show that gadusol is produced during oogenesis and persists in the embryo until at least 5 dpf. Second, we demonstrate that maternally deposited gadusol safeguards embryonic and larval development by preventing UV-induced developmental defects and improving survival. Third, we find that gadusol acts as a true sunscreen preventing the formation of CPDs, a signature of UVB-induced DNA damage, and consequently reducing levels of cell and organismal death. Gadusol does not have any obvious functions beyond protecting against UVR, as mutants survive to adulthood and are fertile. Together, these data demonstrate that gadusol is a maternally provided sunscreen employed during early fish development.

Our work explores two alternative mechanisms of UV protection during early development. We find that the chorion, a transparent eggshell that shields the developing embryo, also provides modest UV protection during embryogenesis. This protection is short lived (zebrafish hatch by 2–3 dpf) but

Figure 4. Gadusol production has been lost in several species no longer exposed to UVR

(A) For each of 136 teleost species (full tree in Figure S4), we assessed various life history traits that identify habitats that may not require embryonic protection from UVR, including electroreception, live-bearing, cave dwelling, and deep-sea dwelling, indicated with colors in the legend to the left of the phylogeny. For each

(legend continued on next page)

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may provide secondary protection during the most vulnerable stages of development. Melanin pigmentation emerges around embryo hatching and serves a relatively modest role as a whole-body sunscreen in 5 dpf larvae. Together, our results show that gadusol is the primary sunscreen across embryonic and larval development, while melanin and the chorion play secondary roles during distinct phases of development.

Finally, our phylogenetic analysis of gadusol biosynthetic genes, building on a previous study,[6] suggests that gadusol is an ancient sunscreen conserved broadly to protect teleost embryos. However, gadusol production has been repeatedly lost during teleost evolution. Intriguingly, these genes are absent in many fish species whose embryos are not exposed to UVR, including deep sea-dwelling and electroreceptive fish. We suggest that, similar to our protected fish facility environment, gadusol is also dispensable for embryonic development in natural environments that lack UVR. In microorganisms, the production of sunscreening compounds has been estimated to require >10% of all metabolic activity.[4] Perhaps the loss of gadusol production in nutrient-poor dark habitats provides some evolutionary advantage, analogous to the energy conservation hypothesis invoked to explain the repeated loss of eyes in Mexican cavefish.[19,20] Similar to loss of UV-responsive gene expression in a cavefish,[13,14] gadusol appears dispensable in species not exposed to sunlight. Once these genes have been lost, descendent species may enter an evolutionary fitness trap where they are confined to breeding environments lacking UVR.

It remains unclear what role gadusol might play in other tetrapods. Functional copies of *eevs* and MT-Ox have been found in numerous vertebrate genomes,[6] but to our knowledge the presence of gadusol has never been reported in vertebrates other than fish. Gadusol has been detected in the eggs or embryos of several aquatic invertebrates, including sponge,[21] starfish,[21] sea urchin,[22] and brine shrimp.[23] We hypothesize that gadusol may also protect early development in these diverse aquatic organisms.

Oxybenzone and octinoxate, two common ingredients in commercial sunscreens, have been recently banned in Hawaii due to concerns of toxicity to coral reefs.[24] Others have suggested gadusol might be a safe natural sunscreen replacement.[6,25] Our data support a role for gadusol as an effective natural sunscreen that warrants further investigation as a preventative agent.

Here, we show that aquatic vertebrates produce and employ an additional sunscreen to melanin. Melanin and gadusol both absorb well in the UVB spectrum. However, melanin also absorbs most wavelengths in the visible light spectrum, making it opaque and conspicuous while gadusol is transparent and invisible. Transparency as camouflage is a common trait in aquatic animals, especially in the open ocean where there is nothing to hide behind.[26] To date, gadusol has only been detected in aquatic organisms. We speculate that gadusol has been particularly advantageous to these animals as it offers protection from UVR, enabling an organism to stay in nutrient-rich sunlit areas while remaining optically inconspicuous. We propose that aquatic ecosystems exhibit unique ecological challenges that have selected for the use of a transparent sunscreen.

STAR★METHODS

Detailed methods are provided in the online version of this paper and include the following:

- KEY RESOURCES TABLE
- RESOURCE AVAILABILITY
  - Lead contact
  - Materials availability
  - Data and code availability
- EXPERIMENTAL MODEL AND SUBJECT DETAILS
  - *In vivo* animal studies
- METHOD DETAILS
  - Generation of *eevs* mutant lines
  - Gadusol extraction and UPLC MS/MS detection
  - Gadusol detection via nanodrop
  - UV exposure, swim bladder inflation, and survival assays
  - Determination of CPDs in 24 hpf embryos
  - Apoptosis assay
  - RNAseq sample prep, library prep, sequencing, and analysis
  - qRT-PCR
  - Generating embryos that lack melanin and gadusol
  - Chorion UV protection assay
  - Phylogenetic analysis of eevs and MT-Ox presence
- QUANTIFICATION AND STATISTICAL ANALYSIS

SUPPLEMENTAL INFORMATION

Supplemental information can be found online at https://doi.org/10.1016/j.cub.2023.06.012.

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species, we identified the presence of intact ORFs for *eevs* and/or MT-Ox. Species that have lost the genes required for gadusol production are indicated in red. We found 16 independent losses across this phylogeny. We found that fish with these traits are more likely than by chance to lose gadusol ($p = 0.012$).
(B) Examples of gadusol synthesis gene loss and pseudogenization in select species. Note *Danio rerio* has intact *eevs* and MT-Ox genes and is capable of gadusol production.
See also Figures S1 and S4 and Data S2.

AUTHOR CONTRIBUTIONS

M.C.R. and J.A.G. conceived of the study. M.C.R. created *eevs* knockouts. M.C.R. carried out and designed UV experiments. J.M. and M.C.R. performed RNA-seq experiments. M.C.R. analyzed RNA-seq data with the University of Utah bioinformatics core. D.L.F. analyzed and ran samples on UPLC-MS/MS. J.H.L. explored fish phylogeny and determined gene loss with input from N.L.C. and M.C.R. M.C.R. and J.A.G. wrote the manuscript with input from all authors.

DECLARATION OF INTERESTS

The authors declare no competing interests.

INCLUSION AND DIVERSITY

One or more of the authors of this paper self-identifies as an underrepresented ethnic minority in their field of research or within their geographical location. One or more of the authors of this paper self-identifies as a member of the LGBTQIA+ community. One or more of the authors of this paper received support from a program designed to increase minority representation in their field of research.

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STAR★METHODS

KEY RESOURCES TABLE

REAGENT or RESOURCE	SOURCE	IDENTIFIER
Antibodies		
Mouse monoclonal Anti-cyclobutane pyrimidine dimers	Cosmo Bio	Cat# NM-DND-001; RRID: AB_1962813
Rabbit anti-active Caspase-3	BD Biosciences	Cat# 559565; RRID: AB_397274
Alexa Fluor 546 goat anti-mouse IgG	ThermoFisher	Cat# A-11030; RRID: AB_2534089
Alexa Fluor 594 goat anti-rabbit IgG	ThermoFisher	Cat# A-11012; RRID: AB_2534079
Biological samples		
Danio rerio embryos	This paper	N/A
Danio rerio larvae	This paper	N/A
Danio rerio ovaries	This Paper	N/A
O. latipes embryos	This Paper	N/A
I. punctatus ovaries	This Paper	N/A
Chemicals, peptides, and recombinant proteins		
DAPI	Sigma-Aldrich	Cat# D9542
Methanol	Sigma-Aldrich	Cat# 34860
Critical commercial assays		
QuanTiTect Reverse Transcription Kit	Qiagen	Cat# 205311
PowerUp SYBR Green Master Mix	ThermoFisher	Cat# A25742
Deposited data		
Code for bioinformatic exploration of loss of gadusol	This paper	https://github.com/nclark-lab/gadusol
RNAseq data (raw and analyzed)	GEO	GEO: GSE229587
Experimental models: Organisms/strains		
Zebrafish *D. rerio* (Tübingen strain)	ZIRC	ZL57
Zebrafish *D. rerio* (AB strain)	ZIRC	ZL1
Zebrafish *D. rerio* (*mitfa*$^{w2/w2}$)	ZIRC	ZL2104
Zebrafish *D. rerio* (*eevs zj2/zj2*)	This paper	N/A
Zebrafish *D. rerio* (*eevs zj5/zj5*)	This paper	N/A
Oligonucleotides		
Primers for sequences cloning, see Table S1	This paper	N/A
Primers for qRT-PCR, see Table S1	This paper	N/A
Software and algorithms		
GraphPad	GraphPad Software	https://graphpad.com
ImageJ	NIH	https://imagej.nih.gov/ij/
Microsoft Excel Data Analysis Student t test	Microsoft	N/A
Other		
Benchtop UV Transilluminator	UVP	M-15V P/N 95-0456-01
UVB broad band bulb (306nm)	Ushio	G8T5E
Digital UV Radiometer UVB	Solarmeter	6.0 UVB

RESOURCE AVAILABILITY

Lead contact

Further information and requests for resources and reagents should be directed to and will be fulfilled by the Lead Contact, James A. Gagnon (james.gagnon@utah.edu).

Materials availability

The fish *eevs* knockout mutant lines generated in this paper (zj2 and zj5) are maintained in the laboratory of James A. Gagnon and are available upon request.

Data and code availability

- RNA-seq data have been deposited at GEO under accession number GEO: GSE229587 and are publicly available as of the date of publication. Accession numbers are listed in the key resources table. Microscopy data reported in this paper will be shared by the lead contact upon request.
- All original code has been deposited at GitHub and is publicly available as of the date of publication. DOIs are listed in the key resources table.
- Any additional information required to reanalyze the data reported in this paper is available from the lead contact upon request.

EXPERIMENTAL MODEL AND SUBJECT DETAILS

In vivo animal studies

Zebrafish D. rerio *(Tübingen and AB strains) embryos and larvae*

All zebrafish work was performed at University of Utah's CBRZ zebrafish facility. This study was conducted under the approval of the Office of Institutional Animal Care and Use Committee (IACUC no. 18-2008) of the University of Utah's animal care and use program. Zebrafish were maintained in a water circulation system at 28° C with a 14hr light and 10hr dark cycle. Fish were fed twice daily. Embryos were either exposed to UVB at 1 dpf or 5 dpf.

METHOD DETAILS

Generation of *eevs* mutant lines

To generate a stable gadusol-depleted mutant line, *eevs* was targeted using CRISPR-Cas9 mutagenesis. Four gRNAs (Data S1) were designed using ChopChop,[27] targeting exon 2 (Figure S1A) due to the lack of suitable target sites within the small exon 1. Guide RNAs were synthesized from DNA oligos using standard protocols.[28] Freshly laid wild-type TU-strain embryos were injected with SpCas9 protein (NEB) mixed with gRNAs (~300 ng/μl), KCl, and phenol red. 1-2 nL were injected into each embryo. Mosaic mutant embryos were raised to adulthood and outcrossed to wild-type Tübingen strain. Primers designed from ChopchopV2[27] were used to amplify the region targeted for CRISPR editing and to select for edited alleles with large deletions. A compound deletion allele was identified by Sanger sequencing that removes 393 bp from the *eevs* open reading frame (Figure S1A; sequences in Data S1) (Genewiz). This *eevs* mutant allele was given the designation zj2 and can be genotyped using PCR with allele specific primers (Data S1). Sibling fish with the zj2 allele were crossed to produce homozygous *eevs*$^{zj2/zj2}$ fish, labeled as *eevs*$^{-/-}$ in Figures 1B–1E and 2A–2D. Because *mitfa* and *eevs* are adjacent genes in the zebrafish genome, an additional *eevs* mutant line was generated in the *mitfa*$^{w2/w2}$*; mpv17*$^{-/-}$ mutant background using the CRISPR protocol described above. A compound deletion allele was identified by Sanger sequencing that removes 161 bp from the *eevs* open reading frame (Figure S1A; sequences in Data S1). This *mitfa*; *eevs* double mutant allele was given the designation zj5, and was used in Figures 3A and 3B. Embryos resulting from crosses of *eevs*$^{-/-}$ mothers had little to no gadusol compared to wild-type embryos, confirming the successful generation of gadusol-depleted lines.

Gadusol extraction and UPLC MS/MS detection

Gadusol was extracted twice from embryos (7.5mg of vacuum dried egg material, crushed with a microfuge pestle) using 150 μl of a (80:20, v/v) methanol:water solution. The extraction supernatant was analyzed using ultraperformance liquid chromatography (Waters Acquity I-Class, 2.1 × 100 mm BEH Amide column) and mass spectrometry (Waters Xevo G2 QToF) (UPLC-MS) in negative ionization mode (detector range of 50-2000 Da). We used a regular phase chromatography method starting with 95% acetonitrile (+0.1% formic acid) and 5 % water (+0.1% formic acid) following a linear gradient over 12 min ending with 30% acetonitrile (+0.1% formic acid). Analytical standards of pure gadusol were run during the same acquisition run to match the retention time and observed mass between embryo samples and the pure standard.

Gadusol detection via nanodrop

To monitor gadusol production the UV-vis spectrometry on a Nanodrop was employed to determine relative gadusol concentrations. Briefly, 25 embryos/larvae were placed in a microfuge tube. All excess water was removed with a Pasteur pipette. 100 μl of 80:20 (v:v) methanol:water was added to embryos. Embryos were mashed with a microfuge pestle for 15 s. Samples were left to extract for at least 15 min, and then centrifuged at 12,000 g. Clear supernatant, containing polar compounds such as gadusol, was separated and analyzed on the nanodrop.

UV exposure, swim bladder inflation, and survival assays

24 hpf embryos were exposed to 450 J of UVB as measured on a radiometer (Solarmeter UVB) at a fluence rate of 2.5 W/m^2 in 30mL of clear E3 media. This is a conservative estimate of a physiologically relevant UVB dose that fish embryos would routinely experience in the wild.[17] A raised and inverted UVP transilluminator with 306 nm broadband UVB bulbs was used (Ushio G8T5E) on the "low" setting (see Figures S2A and S2B). Embryos were returned to the incubator and kept in the dark after mock or UV exposure. Swim bladder inflation was scored at 5 dpf by adding ice to the petri dish to stun the larvae, followed by manual counting on a dissection scope. A standard dose curve was conducted to determine that 450 J/m^2 was an appropriate dose (Figure S2C). 5 dpf larvae were exposed to a dose curve to determine that 2.5 kJ/m^2 was an appropriate dose (Figure S2E). After mock or UV exposure, larvae were placed in an incubator for 1 day (dark) and then placed in the nursery at 6 dpf. Survival was scored at 28 days post-fertilization to ensure that all living juveniles could feed on their own and were not being sustained on maternal yolk. See also Data S3.

Determination of CPDs in 24 hpf embryos

24 hpf embryos were dechorionated to obtain more consistent UV exposure. Embryos were exposed to 450 J/m^2 of UVB and then immediately fixed after exposure in 4% PFA for 1 h at 25°C. After exposure to UVR, embryos were kept in the dark and covered in tin foil while being fixed in PFA to avoid photoreactivation. Fixed embryos were then washed in PBST. Embryos were exposed to 2 M HCl for 1 h to break apart dsDNA and expose CPD epitopes. Samples were blocked in 5% NGS + PBST. Mouse anti-CPD primary antibody (TDM-2, Cosmo Bio) was used to stain for CPDs. Goat anti-mouse AF546 secondary antibody (Invitrogen) was used to visualize CPDs. Embryos were also stained with DAPI to visualize nuclei. Prior to imaging on a confocal microscope, tails were removed from embryos and placed on a flat glass slide with a small drop of PBST. A cover slip was mounted over the tails and sealed with nail polish. Tails were then imaged on an inverted confocal microscope with a 20x objective (Zeiss 880). Images were analyzed using ImageJ[29] to determine mean fluorescence intensity / tail area using the DAPI channel to create a mask for the tail. See also Data S3.

Apoptosis assay

24 hpf embryos within chorions were exposed to 450 J of UVB and then placed in the incubator for 5 h. Chorions were removed and embryos were fixed for 1 h in 4% PFA. Embryos were stained with an activated caspase-3 antibody (BD Biosciences, anti:Rabbit) to mark apoptotic cells. Goat anti-rabbit AF594 secondary antibody (Invitrogen) was used to visualize apoptotic cells. Embryo tails were removed, processed, and imaged as above. ImageJ was used to process images and count the number of activated caspase-3 positive nuclei/mm^2. See also Data S3.

RNAseq sample prep, library prep, sequencing, and analysis

After 5 or 24hrs post UV exposure embryos were smashed with a microfuge pestle (MTC Bio) and RNA extracted using TRI Reagent (Zymo) and purified via Direct-zol RNA Miniprep Plus (Zymo). Library prepared using NEBNext Ultra II Directional RNA Library Prep with poly(A) mRNA Isolation. Samples then sequenced with Total RNA (eukaryote) NovaSeq SP Reagent Kit v1.5_50x50 bp. Each sample sequenced to a depth of 25 million reads. Reads aligned using STAR[30] and zebrafish reference genome (GRCz11). Optical duplicates removed and adapters trimmed. Differential expression analysis conducted with DESeq2[31] and specifically the Bioconductor package.[32] See also Table S1.

qRT-PCR

24 hpf embryos were exposed to 5hrs of cool white LED light while control embryos were kept in constant darkness. RNA was extracted as described above. cDNA was synthesized using QuanTiTect Reverse Transcription Kit (Qiagen). PowerUp SYBR Green Master Mix (Thermo Fisher Scientific) was used for qPCR reactions in a QuantStudio 3 (Thermo Fisher Scientific). Primers for *cry5, ddb2, per2,* and *xpc* were obtained from Weger et al.[33] *Neil1* was obtained from Zhao et al.[13] Housekeeping control gene *elfa* was obtained from McCurly and Callard.[34] Fold expression was calculated using the 2(-Delta Delta C(T)) method.[35] Primers listed in Data S1C.

Generating embryos that lack melanin and gadusol

To generate embryos that lacked melanin, *mitfa*$^{w2/w2}$ fish were crossed with *mitfa*$^{+/w2}$ fish to produce clutches of 1:1 pigmented:unpigmented siblings, all with maternally provided gadusol (Figure 3A). To generate embryos that lack both melanin and gadusol, *mitfa*$^{+/w2}$*; mpv17*$^{-/-}$; *eevs*$^{zj5/zj5}$ females were crossed to *mitfa*$^{w2/w2}$*; eevs*$^{+/+}$ males to produce 1:1 pigmented:unpigmented siblings that all lacked maternal gadusol. Lack of gadusol was confirmed using Nanodrop.

Chorion UV protection assay

24 hpf wild-type TU-strain embryos were manually dechorionated with forceps in a dish with a thin film of 0.5% agar on the base of the dish. Embryos were moved with a fire-smoothened Pasteur pipette. Embryos were exposed to 450 J of UVB as described above and then placed in incubator and swim bladder inflation was scored at 5 dpf. See also Data S3.

Phylogenetic analysis of eevs and MT-Ox presence

123 genomes were gathered from the UCSC genome ark (GenArk) and additional 11 genomes for deep sea and electro-receptive fish were gathered NCBI genomes for all except the Yap Hadal snailfish[36] and pseudoliparis swirei.[37] A BLAST database for each species

was created by using the zebrafish sequence spanning from FRMD4B to FOXP1 to find the same region in all curated genomes. If there was no BLAST hit for FOXP1 or MITF then the genome was dropped for low quality. We then performed a tBLASTn search on the created databases for the remaining genomes, using the zebrafish EEVS and MtOX translated nucleotide sequence as the query. If there was no hit for EEVS or Mt-OX in the tblastn search, we expanded the search from the FRMD4B-FOXP1 region to the entire genome. If there were still no hits at an e-value <10e-50 that species was labeled as not having gadusol. If a species did have a tblastn hit and an e-value < 10e-50 the hits were analyzed for pseudogenization by aligning back to the zebrafish mRNA sequence. The aligned regions were checked for missing exons, frameshifts causing premature stop codons, as well as checked for potential genome masking. If the alignment showed either a deletion of an exon or a premature stop codon and had no evidence of masked regions, the gene was called a pseudogene and marked as "absent" for that species.

To correlate the presence/absence of gadusol with life history traits we first collected life history data for all species (Data S2) from information available on fishbase.[38] The life history traits that we annotated were: deep-sea, nocturnality, live-bearing, electro-reception, and cave dwelling. We then built a species tree using fishtree[39] and added the Yap hadal snailfish[36] and Pseuolapris swirei[37] using the phylogenetic relationship determined in Mu et al.[36] Due to gene loss in sister species not being independent, we used Bayestraits[40] to perform the correlation test. We used discrete model testing and a likelihood ratios test comparing each of the five life-history traits to loss of gadusol (Data S2).

When running Bayestraits the loss of gadusol (parameter beta1 in the independent model and q31 and q42 in dependent model) was set as trait one and the various life history traits were set as trait two. The rate at which gadusol can be regained after loss was constrained to zero because we were scoring for loss of the gene, and assumed it is nearly impossible to regain the gene, especially in the short time span we are investigating. The parameters that estimate the rate of life history traits changing from absent to present (q12,q34,q21,q43) were constrained to equal to each other, under the assumption that it is unreasonable that a fish would change its lifestyle after loss of gadusol. When comparing the cave life history to gadusol loss, the parameter that estimates the rate of moving from cave to surface (q21 and q43) was constrained to zero under the assumption that species do not re-emerge from a cave after adapting to that life-style.

The significance of the correlation between life-history trait and loss of gadusol was determined using a likelihood ratio test which is calculated by 2*((dependent model likelihood)-(independent model likelihood)). The significance is then determined using a chi-sq distribution with 2 degrees of freedom.

All parameters and code to re-run these models can be found in https://github.com/nclark-lab/gadusol.

QUANTIFICATION AND STATISTICAL ANALYSIS

Data were analyzed using GraphPad (https://www.graphpad.com/quickcalcs/contingency1/) when Fisher's exact t-test was being performed. Student's t-test was performed using Microsoft Excel. Two-tailed p values were calculated assuming equal variance in samples. P values < 0.05 were considered statistically significant.