Transcriptomic and genomic evolution under constant cold in Antarctic notothenioid fish

Zuozhou Chen1*, C.-H. Christina Cheng1#, Junfang Zhang1*,5, Luxue Cao4*, Lei Chen*, Longhai Zhou*, Yudong Jin*, Hua Ye9, Cheng Deng5, Zhonghua Dai5, Qianghua Xu*, Peng Hu8,9, Shouhong Sun*, Yu Shen*, and Liangbiao Chen*8

*Key Laboratory of Molecular and Developmental Biology, Institute of Genetics and Developmental Biology, §Graduate School of Chinese Academy of Sciences, Chinese Academy of Sciences, Beijing 100101, China; and ‡Department of Animal Biology, University of Illinois, 515 Morrill Hall, Urbana, IL 61801

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The antifreeze glycoprotein-fortified Antarctic notothenioid fishes comprise the predominant fish suborder in the isolated frigid Southern Ocean. Their ecological success undoubtedly entailed evolutionary acquisition of a full suite of cold-stable functions besides antifreeze protection. Prior studies of adaptive changes in these teleost fishes generally examined a single genotype or phenotype. We report here the genome-wide investigations of transcriptional and genomic changes associated with Antarctic notothenioid cold adaptation. We sequenced and characterized 33,560 ESTs from four tissues of the Antarctic notothenioid Dis-sostichus mawsoni and derived 3,114 nonredundant protein gene families and their expression profiles. Through comparative analyses of same-tissue transcriptome profiles of D. mawsoni and temperature/tropical teleost fishes, we identified 177 notothenioid protein families that were expressed many fold over the latter, indicating cold-related up-regulation. These up-regulated gene families operate in protein biosynthesis, protein folding and degradation, lipid metabolism, antioxidation, antipoptosis, innate immunity, choriogenesis, and others, all of recognizable functional importance in mitigating stresses in freezing temperatures during notothenioid life histories. We further examined the genomic and evolutionary bases for this expressional up-regulation by comparative genomic hybridization of DNA from four pairs of Antarctic and basal non-Antarctic notothenioids to 10,700 D. mawsoni cDNA probes and discovered significant to astounding (3- to >300-fold, \( P < 0.05 \)) Antarctic-specific duplications of 118 protein-coding genes, many of which correspond to the up-regulated gene families. Results of our integrative tripartite study strongly suggest that evolution under constant cold has resulted in dramatic genomic expansions of specific protein gene families, augmenting gene expression and gene functions contributing to physiological fitness of Antarctic notothenioids in freezing polar conditions.


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1Z.C., C.-H.C.C., and J.Z. contributed equally to this work.

*To whom correspondence should be addressed. E-mail: lichen@genetics.ac.cn.

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Transcriptome Comparisons Between D. mawsoni and Warm-Water Teleost Fishes. The highly biased transcription patterns and the notable stress-responding nature of the transcriptomes of the cold-adapted D. mawsoni may represent manifestations of cold-adaptive gene expression. To investigate this possibility and uncover potential cold-adaptive genes in D. mawsoni, we performed same-tissue comparative analyses of D. mawsoni transcriptomes and available transcriptomes of model warm-water teleost fishes (because transcriptome data from related non-Antarctic notothenioids were lacking). We downloaded the sequences of 11 (four brain, three liver, three ovary, and one head kidney) unnormalized high-volume EST datasets of five temperate/tropical teleosts, Danio rerio, Salmo salar, Gasterosteus aculeatus, Fundulus heteroclitus, and Orzias latipes, and processed them with the same bioinformatics pipeline used for D. mawsoni EST data. The ratio of the expression level (EST frequency) of each of the 3,114 D. mawsoni protein gene families and its homolog in the same tissue in the warm-water reference teleost was calculated (detailed methods in SI Appendix, SI Text, File s1, in SI Appendix). Through rigorous statistical evaluation and stringent consistency checks for all comparison pairs (false discovery Rate 10%; corrected P value of comparisons at <0.05 in Fisher’s exact test; SI Text, File s1, in SI Appendix), we identified 189 gene families as differentially expressed (Table S3 in SI Appendix) relative to the warm-water species. Of the 189, 177 (94%) were up-regulated in D. mawsoni (Fig. 2A), remarkably consistent with the observation in carp Cyprinus carpio that 97% (252 of 260) of differentially expressed genes during cold acclimation were up-regulated (20). Furthermore, 85 (48%) of the up-regulated genes in D. mawsoni were homologs of the cold-induced C. carpio transcripts (20), attesting to the feasibility of our comparative transcriptomics approach in detecting cold-related gene expression in D. mawsoni despite phylogenetic distances of the reference species.

Many of the differentially expressed gene families are tissue-specific, with brain and liver showing greater numbers (Fig. 2A). Approximately 20% of the differentially up-regulated genes are found in two or more tissues, suggesting the presence of a common underlying set of enhanced cellular responses. The lack of qualified head kidney transcriptomes in the database limited us to a single comparison for this tissue; nevertheless, a trend toward up-regulation was visible (Fig. 2A), most notably in the coevolution of 29 ribosomal protein genes (Table S3 in SI Appendix). The increase in ribosomal biogenesis suggests enhanced protein synthesis capacity in this tissue, which was reported for the gill and white muscle of another cold-adapted Antarctic teleost, the zoarcid Pachycentr us brachycephalum (21).

To assist in functional interpretations and evaluate potential cold-adaptive response, we performed Gene Ontology (GO) annotation and literature searches by which most of the differentially expressed genes were assigned to 15 functional groups (Table S3 in SI Appendix). The main subsets of genes with related biological functions in 11 of the 15 GO groups are presented in Fig. 2B i-ix. These are: (i) molecular chaperones including the conserved heat-shock proteins and protein-specific chaperones (22); (ii and iii) genes that operate in ubiquitin-dependent and -independent protein degradation machineries; (iv) lipid transport and membrane metabolism-related genes; (v) reactive oxygen species (ROS) scavengers; (vi) apoptosis regulators, especially those of antiapoptotic functions; (vii) genes involved in metal ion and solute homeostasis; (viii) plasma proteins of blood coagulation and innate immunity; (ix) diverse eggshell (ZP) proteins; (x) components of the Ras/MAPK signal transduction pathway; and (xi) factors involved in messenger RNA and ribosomal RNA transcription. The concomitant differential transcript frequencies, i.e., the total number of ESTs within a gene family, were tabulated for each of the four D. mawsoni transcriptomes (Table S2 in SI Appendix) and compared as TreeView plots (15). Blue and white indicate the presence or absence, respectively, of the specific transcripts. Transcript abundance is indicated by the intensity of the blue color. (B) The top tissue-specific and all-tissue-expressed genes and their transcript percentages in the transcriptome(s). Total number of ESTs in the transcriptomes is shown in parentheses.
up-regulation of multiple genes within these functional categories
in *D. mawsoni* compared with warm-water teleosts suggests
greater demands for these functions in the cold Antarctic
environment. The cold-adaptive relevance of their expression
is supported by the fact that the majority (75%) of the top
expressed genes in *D. mawsoni* tissues (Fig. 1B) are found in this
differentially up-regulated gene set (Table S3 in SI Appendix).
The co-up-regulated putative cold-adaptive genes identified lend support to reported evidence and provide grounds for
testable hypotheses regarding mechanisms of cold adaptation at
cellular and systems levels. One example is the co-up-regulation
of three groups of genes operating in protein homeostasis, i.e.,
maintenance of functioning protein pools (Figs. 1B, 2Bi, 2Bii, and
2Biii). The concerted up-regulation of HSPs and protein-specific
chaperones (Figs. 1B and 2Bi) in *D. mawsoni* strongly indicates
a constitutive system-wide need for mitigating cold denaturation
of proteins (23) in the face of chronic cold. It supports the
reported constitutive HSP70 expression and loss of HSP70
inducibility in some notothenioids (24). The augmented need for
cellular protein homeostasis is also indicated in the up-
regulation of genes in ubiquitin-dependent protein degradation (Fig. 2Bii)
respectively, of transcript abundance in *D. mawsoni* relative to the compared species (*P* < 0.05 in Fisher’s exact test). Black indicates no difference. The color scale
is derived from log2 transformation of differential expression ratios in Table S3 in SI Appendix. Rows dotted with a small black square indicate the homologs
of cold-responding genes identified in *C. carpio* (20). (Right) Names and EST counts of the highly up-regulated genes in each functional group. Sequentially, the
11 groups represent: molecular chaperones, ubiquitin-dependent protein degradation, proteolytic enzymes and inhibitors, lipid metabolism, antioxidant, apoptosis regulation, ion and solute homeostasis, immunity, zona pellucida proteins, Ras/MAPK pathway, and transcription factors.

Fig. 2. Transcriptome comparison between *D. mawsoni* and temperate/tropical teleosts revealed co-up-regulation of multiple gene groups in the Antarctic
nototheniid fish. (A) Tissue distribution and expression ratio of 189 differentially expressed genes between *D. mawsoni* and five temperate/tropical teleost
tissues identified from same-tissue transcriptome comparisons. Lanes B1-B4, L1-L3, O1-O3, and K1 showed comparison results of *D. mawsoni* brain vs. *D.
rerio*-1522, -14409 and *G. aculeatus*-17209, -18921 brain libraries; *D. mawsoni* liver versus *D. rerio*-14410, *F. heteroclitus*-15870 and *O. latipes*-17414 liver libraries; *D. mawsoni* ovary versus *S. salar*-15459 and *D. rerio*-9767, -15519 ovary libraries; and *D. mawsoni* head kidney versus *S. salar*-15454 head kidney library, respectively. The graph was derived from data in Table S3 in SI Appendix, clustered and rendered by the same programs described for Fig. 1A. (B) Eleven functional
groups of differentially up-regulated genes in *D. mawsoni*. (Left) TreeView plot. Red and green indicate statistically significant up- or down-regulation,
respectively, of transcript abundance in *D. mawsoni* relative to the compared species (*P* < 0.05 in Fisher’s exact test). Black indicates no difference. The color scale
is derived from log2 transformation of differential expression ratios in Table S3 in SI Appendix. Rows dotted with a small black square indicate the homologs
of cold-responding genes identified in *C. carpio* (20). (Right) Names and EST counts of the highly up-regulated genes in each functional group. Sequentially, the
11 groups represent: molecular chaperones, ubiquitin-dependent protein degradation, proteolytic enzymes and inhibitors, lipid metabolism, antioxidant, apoptosis regulation, ion and solute homeostasis, immunity, zona pellucida proteins, Ras/MAPK pathway, and transcription factors.
creased levels of ROS in their tissues and thus system-wide oxidative stress (26). This hypothesis finds support in three GO categories of up-regulated genes in D. mawsoni involved in ROS and oxidative stress mitigations. Group v (Fig. 2Bv) contains genes that act in ROS scavenging, including PHGPx, peroxiredoxin-5, and superoxide dismutase, and the potent antioxidant haptoglobin that alleviates hemoglobin-driven oxidative stress in blood cells (27). Group vi (Fig. 2Bvi) contains antiapoptotic genes, including B1-1 and TCTP with known roles in mitigating ROS-mediated apoptosis (28) and cellular oxidative stress (29). Group vii (Fig. 2Bvii) contains genes involved in iron transport and storage, including ferritin and serotransferrin that bind free iron and the small peptide hormone hepcidin that reduces iron absorption (30). The up-regulation of these genes would be consistent with the need for controlling free Fe²⁺-catalyzed ROS generation (31). Simultaneous up-regulation of these three groups of genes suggests augmented defenses against oxidative stress likely constituted an important aspect of evolutionary adaptation of the Antarctic notothenioids in their oxygen-rich environment. The identification of cold-specific adaptive evolution of novel hepcidin isoforms in the Antarctic notothenioids (32) lent support to this hypothesis.

Collectively, the results of transcriptome expression patterns in D. mawsoni and comparative transcriptome analyses provided us with an overview of the nature of transcriptomic changes potentially important in cold adaptation. The spectrum of up-regulated gene groups we identified will enable further in-depth investigations into mechanisms of nototheniid evolutionary cold adaptation in much wider physiological contexts than currently examined in the field.

**Gene Duplication in Antarctic Notothenioid Fish Genomes.** To assess genomic contribution to the transcriptional up-regulations in D. mawsoni, we performed array-based Comparative Genomic Hybridization (aCGH) of genomic DNA from selected pairs of related Antarctic and non-Antarctic notothenioids, using 10,700 D. mawsoni ESTs as probes. Three Antarctic species, D. mawsoni, Pagophenia borengrevinki, and Chaenocephalus aceratus, were referenced to the phylogenetically basal New Zealand species Bovichtus variatus or the South American species Eleginus maclovinus (phylogenetic relationship in Fig. S1 in SI Appendix) in four aCGHs (Fig. 3A). Differential hybridization signal intensity of a gene across all four aCGH pairs would indicate a gain or loss of gene copy number during the evolutionary radiation of Antarctic notothenioids.

Using a stringent selection criterion (Antarctic:non-Antarctic signal ratio of 2.4:1) (detailed methods in SI Text, File s2, and Fig. 3A in SI Appendix), we identified 101 protein-coding genes distinct from the well known AFGP gene and 17 long interspersed nuclear elements (LINEs) that were duplicated or potentially newly acquired in the three Antarctic species, and only 12 genes that were contracted (signal ratio <0.43, i.e., 1/2.4) (Fig. 3A; gene details and signal ratios in Table S4 in SI Appendix). Thus, duplicated genes greatly outnumbered contracted ones, by 10:1. Also, mean duplication ranged from ~3- to >300-fold (P < 0.05), far greater than reduction, which are 0.25 and 0.3 (P < 0.01), respectively, for the two most contracted genes, a LINE and Na⁺/K⁺ ATPase α subunit (Table S4 in SI Appendix). The majority of duplicated genes are homologs of known or hypothetical proteins, indicating duplication of pre-existing genes to augment gene function. The dominance of gene duplication over contraction held true when the aCGH selection ratio was lowered and non-protein coding genes were included (Table S5 in SI Appendix). Southern blot hybridizations using probes derived from four of the putative duplicated genes (FBP32II, ZPC5, a 10-aa repeat protein, LINE, with ß-actin as control) showed many intense hybridizing bands in the Antarctic species vs. few and weakly hybridizing bands in B. variatus and E. maclovinus (Fig. 3B), validating the occurrence of extensive gene duplications in the Antarctic lineages inferred from aCGHs.

Among the 118 duplicated genes (Table S4 in SI Appendix), 15 encode different enzymes potentially adaptive for cold survival. For example, matrix metalloproteinases MMP9 and nephrisin are involved in extracellular matrix remodeling, a known process of adaptive importance in cold acclimation (20). Seven genes are involved in protein folding and ubiquitin-dependent protein degradation, and nine genes function in stress-response, anti-ROS, and antiapoptotic processes. Duplication also occurred in nine genes participating in Ras/MAPK and TGF-β signal transduction pathways and in 10 genes encoding proteins of cellular structural components, such as the nuclear pore and cytoskeleton. Also duplicated are five genes involved in RNA binding and processing known to be important for mRNA function in the cold, and five zinc finger proteins potentially involved in gene transcription (Table S4 in SI Appendix). Many of these duplicated genes correspond to genes showing elevated transcription, either being the same genes or genes in the same biological processes (Figs. 1B and 2B), similarly for genes involved in innate immunity, such as hepcidins that act as antibacterial agents besides iron regulator, and FBP32II that encodes F-type lectins. FBP32II showed a 14-fold average increase in gene copies in the three Antarctic species compared with B. variatus and E. maclovinus (Table S4 in SI Appendix, Fig. 3Bj), and >53-fold greater transcription in D. mawsoni liver relative to warm-water teleosts (Fig. 2Bvii, Table S3 in SI Appendix). Lectins recognize specific carbohydrate domains on bacterial cell surfaces and elicit immune responses. Interestingly, fish type II antifreeze proteins evolved from C-type lectins (33), suggesting lectin gene
duplications may be a prevalent response in coldwater teleosts, preceding the cooption of one copy to become an antifreeze.

The highly expressed eggshell protein genes (Figs. 1B and 2B) are also substantially duplicated: ZPC1, -2, -3, -5; ZPX; ZPB and ZPD genes expanded by 2.7- to 7.7-fold (P < 0.05) in the Antarctic notothenioids (Table S4 in SI Appendix). Extensive duplication of ZPC5 was verified in Southern hybridization of genomic DNA (Fig. 3B2). The enlarged ZP gene family undoubtedly contributed to the abundant ZP transcripts seen in D. mawsoni. Antarctic notothenioids synthesize a dense and thick chorion (34). Embryos of Gymnodraco acuticeps and Pleuragramma antarcticum develop in ambient icy freezing seawater but have drastically inadequate amounts of AFGPs to avoid freezing; thus, the dense chorion was considered to act as a crucial physical barrier against ice penetration (35). The need for a durable protective eggshell could have been a strong selection for the extensive duplication of ZP genes.

We also identified >20 duplicated genes without known biological functions (Table S4 in SI Appendix). Two of these, which encode proteins of 10- (EKLNGTMSDE) and 11-residue (RHDGVNETNDV) repeats, respectively, have no homologs in other organisms and may be new acquisitions in Antarctic notothenioids. The extensive duplication of the repetitive 10-residue protein is shown in Fig. 3B3. Both gene types highly transcribed in D. mawsoni ovary (Fig. S3 in SI Appendix), suggesting a role in ovarian function that remains to be elucidated.

The most astounding Antarctic-specific gene duplication revealed by aCGH occurred in the LINEs, genetic elements prevalent in eukaryotic genomes capable of retrotransposition through self-coding enzymes. Seventeen LINE genes encoding reverse transcriptase and endonuclease showed Antarctic-specific duplication by 8- to >300-fold (P < 0.05), whereas only one underwent contraction (Table S4 in SI Appendix, Fig. 3B4). The LINE expansions correlate with increasing genome sizes from the phylogenetically basal notothenioids to the derived species. The C values of B. variegatus, D. mawsoni, P. borchgrevinki, G. acuticeps, and C. aceratus are 0.84, 0.97, 1.28, 1.83, and 1.86 pg, respectively (C.-H.C.C., unpublished data), suggesting the gain of LINEs contributed to genome size enlargement in notothenioid evolution. The LINE expansion might have in turn facilitated the observed protein gene duplications in Antarctic notothenioids through retrotransposition-mediated sequence transduction (36). Although these putative roles of LINEs in Antarctic notothenioids remain to be explored, the remarkable Antarctic-specific LINE expansion and diversification represent a direct link between LINE evolution and cold-selection pressure.

Conclusion
Through integrated genome-wide investigations, we conducted a comprehensive elucidation of the genotypic changes accompanying evolutionary adaptation of Antarctic notothenioid fishes in the freezing Southern Ocean. Evolution in constant cold had apparently resulted in remarkable transcriptomic shifts and gene expansion in Antarctic notothenioids with respect to related and unrelated temperate species. From 6,208 unique protein-coding genes expressed in four tissues of D. mawsoni, we identified 177 gene families that are transcriptionally elevated, and 118 protein genes that are substantially duplicated, bringing about a nonredundant set of >200 genes that are specifically augmented in the Antarctic notothenioids. These genes in concert would enhance an array of biological processes including protein synthesis, protein folding and degradation, lipid metabolism, antioxidation, antapoptosis, innate immunity, and choriogenesis, forging a collective stress-responding or mitigating phenotype to overcome various physiological challenges posed by the freezing and oxygen-rich Antarctic environment during notothenioid life histories. Because many of the duplicated protein genes correspond or are functionally similar to the dominant transcripts, gene family expansions likely contributed significantly to augmentation of cold-adaptive functions. The extensive duplications of the eggshell protein genes, LINEs, and two putative newly acquired proteins of repetitive sequences in the Antarctic species are suggestive of new cellular or genomic mechanisms of cold adaptation that will stimulate further research. The results of this study provide a comprehensive view into the making of notothenioid cold fitness and, in addition, a framework for exploring the genomic adaptability of the ecologically vital Antarctic notothenioids in face of global climate change.

Materials and Methods
Sample Collection. Antarctic notothenioids D. mawsoni and P. borchgrevinki were captured from McMurdo Sound, Antarctica, and C. aceratus from the Antarctic Peninsula. The two basal non-Antarctic notothenioids, B. variegatus and E. maclovinus, were obtained from Otago Harbor, New Zealand, and Ushuaia, Argentina, respectively. Tissues were dissected from anesthetized specimens and kept frozen at ~80°C until use.

Construction of cDNA Libraries and Sequencing. Complementary DNA libraries of D. mawsoni tissues were constructed by using the pCMV-Script XR cDNA Library Construction Kit (Stratagene). Seven micrograms of polyA+ RNA was reverse-transcribed with the addition of 0.6 M d-Trehalose to improve first-strand cDNA length, and cDNAs with sizes >0.7 kb were recovered and directionally cloned. Analysis of random recombinant clones showed an average insert size of ~1.8 kb (37). Plasmid DNA was isolated using AXyprep Easy-96 isolation kit (Axogen Biosciences) and sequenced from the 5’ end of the insert using BigDye v.3.1 (Applied Biosystems) and resolved on an ABI 3730 sequencer.

Sequence Annotation and Transcriptome Comparison. Raw sequence data were edited with PHRED (38) by using a stringent error rate of 0.1%. The ESTs were assembled with CAP3 (39) and annotated with BLAST against National Center for Biotechnology Information databases. A bioinformatics pipeline was designed to generate accurate and nonredundant D. mawsoni gene families. The assembled genes have ~50% homology to known proteins and are at least 70 aa in length (details in SI Text, File s1, in SI Appendix). Associated ESTs of each gene family were summed to obtain the expression level in each tissue. Eleven unnormalized EST libraries of warmtropical teleost fishes were downloaded from the UniGene database. EST frequencies of a homologous gene family in D. mawsoni and the comparison species were divided to obtain the expression ratio. Gene families showing at least 1.5-fold of expression difference and passing Fisher’s exact tests at FDR < 0.1 were retained and further screened for multiple-comparison consistency. Those consistently up- or down-regulated in at least three of the four brain comparisons, all three liver comparisons, all three ovary comparisons, or the head kidney comparison were regarded as differentially expressed genes in D. mawsoni relative to the reference teleost fish (details in SI Text, File s1, in SI Appendix).

Construction of cDNA Microarray and aCGH. Microarray slides containing 10,700 D. mawsoni ESTs and positive (AFGP coding sequence) and negative (DMSO) controls were constructed (details in SI Text, File s2, in SI Appendix). Genomic DNA from livers of D. mawsoni, P. borchgrevinki, C. aceratus, B. variegatus, and E. maclovinus were purified, sheared, and quantified. Three micrograms of sheared DNA from each species of a aCGH pair were labeled with Cy5- or Cy3-dCTP. Dye-reversed controls were performed for all aCGH pairs. Hybridizations followed the procedure developed by CapitalBio (SI Text, File s2, in SI Appendix).

Data Normalization, Selection of Duplicated Genes, and Verification. Hybridized microarrays were scored and the fluorescence intensities extracted using GenePix Pro 4.0 (Axon Instruments). The Cy5 and Cy3 signals were normalized by setting the log2 average Cy5/Cy3 ratio of all of the elements equal to 0. The consistency of the replicated spots was evaluated by Student’s t-test, and the average Cy5/Cy3 ratios of spots that passed the test were calculated from the replicas, whereas failed spots were eliminated. In interhominoid aCGHs, a log2 signal ratio of 0.5 between the test and reference species was used to infer gene duplication (13). To reflect the longer divergence times among the notothenioids, we adopted the signal ratio of a known duplicated gene, hepcidin between D. mawsoni and E. maclovinus as the selection criterion (Fig. S2 in SI Appendix), which was 2.4 (log2 2.4 = 1.26). EST spots with signal ratios >2.4 or <0.43 in all four CGHs were selected as duplicated or contracted genes and annotated (detail see SI Text, File s2, in SI Appendix). A geometric mean duplication/contraction fold of an individual EST was calculated.
from the four aCGH signal ratios (Pbl/Bv, Dm/Em, Pb/Em, Ca/Em) to refer the extent of duplication or contraction for this gene between Antarctic and non-Antarctic notothenioids. To validate aCGH detected gene duplication, genomic DNA of the five notothenioids used in aCGH was digested with EcoRI and probed with five selected ESTs by Southern blot hybridizations.


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