Abstract  Insufficient blood supply during acute infarction and chronic ischemia leads to tissue hypoxia which can significantly alter gene expression patterns in the heart. In contrast to most mammals, some teleost fishes are able to adapt to extremely low oxygen levels. We describe here that chronic constant hypoxia (CCH) leads to a smaller ventricular outflow tract, reduced lacunae within the central ventricular cavity and around the trabeculae and an increase in the number of cardiac myocyte nuclei per area in the hearts of two teleost species, zebrafish (Danio rerio) and cichlids (Haplochromis piceatus). In order to identify the molecular basis for the adaptations to CCH, we profiled the gene expression changes in the hearts of adult zebrafish. We have analyzed over 15,000 different transcripts and found 376 differentially regulated genes, of which 260 genes showed increased and 116 genes decreased expression levels. Two notch receptors (notch-2 and notch-3) as well as regulatory genes linked to cell proliferation were transcriptionally upregulated in hypoxic hearts. We observed a simultaneous increase in expression of IGF-2 and IGFbp1 and upregulation of several genes important for the protection against reactive oxygen species (ROS). We have identified here many novel genes involved in the response to CCH in the heart, which may have potential clinical implications in the future.

Keywords  Hypoxia · Zebrafish · Heart · Hyperplasia · Gene expression

Introduction
Low oxygen levels (hypoxia) play important roles in clinical conditions such as stroke and heart failure. Insufficient blood supply leads to tissue hypoxia in the heart during acute infarction and chronic ischemia (Semenza 2001).

Effective protection of the heart against ischemia/reperfusion injury is one of the most important goals of experimental and clinical research in cardiology. Besides ischemic preconditioning as a powerful temporal protective phenomenon, adaptation to chronic hypoxia also increases cardiac tolerance to all major deleterious consequences of acute oxygen deprivation such as myocardial infarction, contractile dysfunction and ventricular arrhythmias (Kolar and Ostadal 2003).
Although many factors have been proposed to play potential roles, the detailed mechanism of this long-term protection remains poorly understood. Some of the molecular mechanisms of cardiac protection by adaptation to chronic hypoxia and chronic high-altitude hypoxia have recently been reviewed (Kolar and Ostadal 2003; Ostadal and Kolar 2007). $K_{\text{ATP}}$ channels, PKCδ as well as the different MAPK pathways were shown to be involved in the mechanism of increased tolerance of chronically hypoxic hearts and further the controversial role of ROS in hypoxia tolerance is discussed (Kolar and Ostadal 2003). Furthermore, a recent study has profiled the gene expression changes induced by chronic constant hypoxia (CCH) and chronic intermittent hypoxia (CIH) in newborn mice (Fan et al. 2005).

In contrast to most mammals (with the exception of some marine mammals), some teleosts, have developed the ability to withstand extreme chronic hypoxia (Stecyk et al. 2004). It is well assumed that these vertebrate species possess unique adaptations in order to survive short and long term oxygen deprivation. However, the molecular basis of these adaptations in fish has so far not been extensively investigated.

Several studies have profiled gene expression changes in teleosts exposed to hypoxia. Gracey et al. showed in adults of the euryoxic gobid fish *Gillichthys mirabilis* (the long-jawed mudsucker), that 5 days of hypoxia induced a complex transcriptional response, including a shut down of energy requiring pathways like protein synthesis and locomotion, and an induction of genes needed for anaerobic ATP production in different tissues (Gracey et al. 2001). Recently, we described phenotypic and behavioral adaptations to long-term hypoxia and described the gene expression changes induced by chronic constant hypoxia in the gills of adult zebrafish (van der Meer et al. 2005).

Ton et al. identified global gene expression changes in zebrafish embryos. Zebrafish embryos at 48 h post fertilization were exposed to water with 5% oxygen content for 24 h. The authors identified 138 genes responsive to short-term hypoxia and could also show that transcriptional changes indicated metabolic depression, a switch from aerobic to anaerobic metabolism and energy conservation (Ton et al. 2003).

In this study, we have identified CCH-induced gene expression changes in the zebrafish heart by looking at over half of the zebrafish genome. We have compared several of these novel changes described in other species and tissues. We have here identified the heart-specific molecular adaptations to CCH. Future functional experiments are warranted to determine whether some of the findings can be used to better adapt mammalian hearts to CCH.

### Material and methods

#### Animal handling

Adult wild-type zebrafish (*Danio rerio*) around 3 month of age, were obtained from a local pet store. Cichlids (*Haplochromis piceatus*) have been collected in the Mwanza Gulf of Lake Victoria in 1984 and were bred in our laboratory for about 20 generations. All animals were handled in compliance with animal care regulations. Our animal protocols were approved by the review board of Leiden University in accordance with the requirements of the Dutch government. Zebrafish were kept at 25°C in aquaria with day/night light cycles (12 h dark vs. 12 h light). Cichlids were kept at 25°C with the same day/night light cycles.

#### Hypoxia treatment

For gradual hypoxia treatment, oxygen levels were gradually decreased in 4 days from 100% air saturated water to 40% (day 1), 30% (day 2), 20% (day 3) and the final 10% air saturation (day 4). After day 4, the fish were kept for an additional 21 days at 10% air saturation (at 100% air saturation and 28°C the $O_2$ concentration is 8 mg/l and pO$_2$ is 15 Torr). In parallel, a control group was kept at 100% air saturated water. Both groups were kept in identical aquaria of 100 l. The oxygen level on the hypoxia group was kept constant by a controller (Applikon Biotechnology, The Netherlands) connected to an $O_2$-electrode and solenoid valve in line with an air diffuser. The oxygen level in the tank was kept constant by adding oxygen via the diffuser and thereby compensating the oxygen consumption of the fish. In case of immediate hypoxia exposure, tanks were pre-equilibrated to the respective pO$_2$ concentration and fish were then directly set in the equilibrated aquaria.

#### Perfusion of cichlid hearts

In order to minimize blood clotting, a perfusion protocol was developed in which the whole blood volume of clinically dead animals was initially replaced with isotonic buffer and with a fixative solution secondarily.

#### Heart dissection

The fish were killed with an overdose of anesthetic (MS-222; Tricaine Methanesulfonate from Argent Chemical Laboratories, USA). Hearts were dissected from the fish immediately after the anesthetic worked. For RNA preparation the hearts were immediately shock-freezed in liquid nitrogen. For histology and microscopy, the hearts were left intact and fixed immediately in Karnovsky
fixative (4% paraformaldehyde (PFA) and 2.5% glutaraldehyde in 0.1 M phosphate buffer, pH 7.2) for 4 h at 4°C. After three washes in 0.1 M phosphate buffer, pH 7.2, they were transferred to 70% ethanol.

Histology of adult fish hearts, statistical analysis and scanning electron microscopy

Hearts from zebrafish and cichlids raised either under normoxic or hypoxic conditions, were dissected from the fish and fixed for 24 h in 4% PFA in PBS. After fixation, hearts were washed with 1 × PBS, cut in halves (sagittal through the midline) and then dehydrated through ethanol series starting at 70% ethanol, followed by 80, 90, 96 and 100% ethanol, each step was done once for 1 h except the last one which was done twice. After dehydration the hearts were embedded in increasing gradients of Historesin (Technovit 7100, Heraus Kulzer, Germany) (25, 50, 75 and 100% Historesin in ethanol, for 2 h at room temperature; 100% Historesin, 24 h at 4°C). Afterwards the plastic with the hearts was polymerized at 40°C (overnight). A 5 μm sagittal sections were made using the Ultramicrotome (Reichert-Jung) and a glass knife. Approximately 500 sections on each side of the midline were visually analyzed per heart (1,000 sections in total per heart). Sections were left to dry and later stained with hematoxylin–eosin staining. Pictures were taken with Axioplan 2 imaging, (Carl Zeiss, Jena). Azan staining of histological sections were done as follows. Paraffin sections of the hearts were prepared as described before (van der Meer et al. 2006). Sections were incubated with Azokarmine solution, for 30 min at 60°C. Afterwards they were washed in water and differentiated in 0.2% Anilin alcohol. They were then rinsed in 1% acetic acid in 95% alcohol, followed by 45 min incubation period, in 5% phosphotungstic acid. After that, sections were rinsed in distilled water and incubated for 45 min with anilene blue. Finally, they were rinsed with distilled water, differentiated and dehydrated in 95% alcohol followed by absolute, cleared in xylene and mounted in Entallan. The protocols used here for SEM had been described before (van der Meer et al. 2005). The statistical analysis of cardiac myocyte nuclei per section was done using Statistica by performing an independent t test. A P value of less than 0.05 was considered significant.

Immunohistochemistry and statistical analysis

Zebrafish, raised either under normoxic or hypoxic conditions, were killed with an overdose of anesthetic MS-222 and frozen in liquid nitrogen. Subsequently, transversal cross-sections (10 μm thick) of the body, were cut using a cryostat at −20°C and mounted on glass slides coated with Vectabond (Vector Laboratories, Burlingame, USA). Sections were fixed in 4% formaldehyde in Tris-buffered saline (TBS; 50 mM Tris and 150 mM NaCl, pH 7.5) for 10 min and subsequently washed in TBS with 0.05% Tween-20 (TBST) (Sigma-Aldrich, Zwijndrecht, The Netherlands). Subsequently, sections were incubated for 10 min with 10% normal swine serum (Vector laboratories) in TBST after which sections were incubated for 24 h at 4°C with anti-phospho Akt polyclonal antibody (Santa-Cruz Biotechnology, USA) diluted 1:50 in TBST. After incubation with primary antibody, the slides were washed in TBST and subsequently placed in 0.25% (v/v) acetic anhydride in 0.1 M triethanolamine buffer (pH 8) for 10 min followed by rinsing in TBST. After this, sections were incubated for 60 min at 20°C with secondary anti-rabbit immunoglobulin G (IgG) antibody covalently coupled to alkaline phosphatase (Vector Labaratories) diluted 1:100 and washed in TBST. After this, sections were incubated for 5 min with alkaline phosphatase (AP) buffer (0.1 M NaCl, 0.1 M Tris, 50 mM MgCl2 and 0.1% Tween-20, pH 9.5) followed by incubation with BM Purple AP substrate (Roche Applied Sciences, Almere, The Netherlands) for 30, 45 or 60 min which was followed by rinsing in TBST. All sections were mounted in glycerine-gelatin and stored at 4°C in the dark until staining intensity was measured. The absorbance values of the BM Purple in the sections were determined using a Leica DMRB microscope (Wetzlar, Germany) fitted with calibrated gray filters using different interference filters. Absorbances for BM Purple were determined at 550 nm. Images were recorded with a ×20 objective and a Sony XC-77CE camera (Towada, Japan) connected to a LG-3 frame grabber (Scion; Frederick, MD) in an Apple Power Macintosh computer. Recorded images were analysed with the public domain program NIH-Image V1.61 (US National Institutes of Health, available at http://rsb.info.nih.gov/nih-image/). Gray values were converted to absorbance values per pixel using the gray filters and a third-degree polynomial fit in the calibrate option of NIH-image programme. Morphometry was calibrated using a slide micrometer and the set scale option in NIH-image, taking the pixel-aspect ration into account. An independent t test was used to test for differences in phospho-Akt levels in cardiac myocytes of normoxic and hypoxic fish. A P value of less than 0.05 was considered significant. Values are means ± S.E.M.

RNA preparation and biological sampling

After dissection hearts were homogenized in a Dounce homogenizer using 1 ml Trizol solution (GibcoBrl, Life technologies). The whole heart was used and for each biological sample hearts were pooled from five different animals. After Trizol extraction, total RNA was further purified using RNAeasy columns (Qiagen). RNA samples were analyzed for quality control by Lab-on-a-chip analysis.
(Agilent) and on agarose gels. For the array experiment five arrays were done for normoxic and 5 arrays for the hypoxic condition. Biological samples (BS) came from two independent experiments and one technical replicate (TR) was included (2BS + 2BS + TR for normoxia and 2BS + 2BS + TR for hypoxia). As mentioned above for each BS, hearts from five different animals were pooled.

Microarray analysis

The Affymetrix GeneChip® Zebrafish Genome Arrays containing 15,509 Danio rerio gene transcripts were used. Probe sets on the arrays were designed with 16 oligonucleotide pairs to detect each transcript and procedures were in full support of MIAME standards. Labeling and microarray hybridization were performed by ServiceXS (Leiden, The Netherlands), including prior a standard round of RNA amplification according to standard Affymetrix protocols. The criteria used for differential expression were greater or equal than 2-fold induced or reduced and P ≤ 0.02. Data analysis was done using Rosetta Resolver. All expression data was submitted to the NCBI Gene Expression Omnibus (http://www.ncbi.nlm.nih.gov/geo). The series entry number is GSE4989 and the following 10 accession numbers were assigned: GSM112796 and GSM112798-GSM112806. A complete list of differential regulated genes is shown in Supplemental Table 1.

Table 1 Statistical analysis of histological sections (5 μm) from zebrafish (Danio rerio) and cichlid (Haplochromis piceatus) hearts

|                     | Zebrafish (nuclei per 900 μm²) | Cichlid (nuclei per 10,000 μm²) |
|---------------------|--------------------------------|--------------------------------|
|                     | Normoxia                       | Hypoxia                        | Normoxia                       | Hypoxia                        |
| Mean                | 9.81                           | 13.67                          | 14.7                           | 24                             |
| SD                  | 0.30                           | 0.39                           | 0.50                           | 0.68                           |
| P value             | 9.9 × 10⁻¹⁷                    | 7.6 × 10⁻¹²                    |

Midline sections of zebrafish and cichlids raised under normoxic and hypoxic conditions were chosen and subdivided in smaller areas. In the case of D. rerio, each subarea of the section was 900 μm² whereas in the case of the H. piceatus it was 10,000 μm². Then, subareas were randomly picked and the amount of cardiac myocyte nuclei were counted. The vast majority of cardiac myocytes were mononucleated cells. Hundred subareas per section were counted per specimen. Three sections per heart were visually analyzed (with a magnification of 20×). In total 6 different zebrafish hearts (from three independent experiments) were investigated for hypoxic conditions and 6 for normoxic condition. In total 1,800 subareas were quantified per heart and condition used. The same was done for the cichlids (for which bigger subareas were counted). A two-tailed t test was applied and a significant difference (P < 0.001) in the amount of nuclei present in the hearts of normoxia versus hypoxia groups was observed for both species.

Gene ontology analysis

The Gene ontology analysis was performed using eGOn and the database of the Norwegian Microarray Consortium (http://www.genetools.microarray.ntnu.no/egon/) and the most recent updated Unigene numbers for zebrafish (May 2007). 13414 Unigene numbers were annotated to the Affymetrix gene chip used. eGOn Enrichment analysis was used for the entire Affymetrix set versus the identified 376 gene set based on the new Unigene numbers. A master target test was performed (using Fishers exact test) and the cut off for positives was set to P < 0.05. The available three categories, molecular function, biological process and cellular component were all tested and are shown in this dataset (Supplemental Data 1). The fold enrichment was based on the percentage found within the differentially expressed genes divided by the percentage on the total chip. A list of all Unigene numbers found in the different categories is given in Supplemental Data 2.

Real-time quantitative RT-PCR

For verification of gene expression data, we used quantitative real-time RT-PCR. Biological RNA samples were obtained from a third independent experiment and as mentioned above for the microarrays, for each BS hearts from five different animals were pooled. The Roche Master SYBR Green kit was used for the RT-PCR reactions. The annealing and synthesis temperature was 55°C alternating with 96°C for 45 cycles. Dissociation protocols were used to measure melting curves and control for unspecific signals from the primers. A measure of 100 ng of total RNA was used per reaction. A standard curve for β-actin using 1, 5, 10, 100 and 500 ng of total RNA was used for normalization. Samples were measured in the Roche LightCycler. The primer3 software (http://frodo.wi.mit.edu/cgi-bin/primer3/primer3_www.cgi) was used to design primers for short amplicons between 50 and 100 bases. The primers used are shown in Supplemental Table 2.

Results

We describe here the survival rates of adult zebrafish upon immediate and gradual exposure to different pO₂ concentrations. Immediate exposure to pO₂ of 15 Torr (O₂ concentration of 0.8 mg/l or 10% air saturated water) was lethal for the adult zebrafish and none survived for longer than 72 h (Fig. 1). If lowered gradually (see experimental procedures for regimen in Sect. “Hypoxia treatment”) zebrafish were able to grow and gain weight at O₂ levels of 10% air saturation. The zebrafish were able to survive for longer than 6 months and no mortality was observed, demonstrating
that they can well adapt to these conditions (data not shown). For the experiments with zebrafish and cichlids, the pO₂ concentration was gradually lowered to 10% air saturation (0.8 mg/l) (see “Material and methods”) and fishes were kept under 10% air saturation for 3 weeks. Control groups were always kept in parallel under normoxic conditions.

Phenotypic changes in the heart of adult teleosts under chronic constant hypoxia (CCH)

In comparison to the normoxic control groups, we observed a significantly smaller ventricular outflow tract and reduced lacunae within the central ventricular cavity and reduced lacunae around the trabeculae in midline sections of hearts of both zebrafish (Danio rerio) (Fig. 2a) and cichlids (Haplochromis piceatus) (Fig. 2b) exposed to CCH. In addition to the midline sections, none of the sections investigated from hypoxia treated fishes showed a ventricular outflow tract in comparable size to the normoxic controls (data not shown). In addition to the midline sections, none of the lateral sections (to both sides of the midline) investigated from hypoxia treated fishes showed a ventricular outflow tract as well as lacunae in a comparable size to the normoxic controls (data not shown). The larger cichlid hearts were also perfused and midline sectioned and showed similar results with a smaller ventricular outflow tract and reduced lacunae (Fig. 2b-H). This might represent ventricular hypertrophy or hyperplasia in both walls and trabeculae, which could lead to the observed cavity obliteration in these sections. We quantified the number of cardiac myocyte nuclei per area in the midline sections of both zebrafish and cichlids under normoxic, as well as hypoxic conditions. A significant difference in both species was observed and showed that hypoxia led to a 1.4- and 1.6-fold increase in the number of cardiac myocyte nuclei per area in zebrafish and cichlid hearts, respectively (Table 1). Cardiac myocyte nuclei in sections were clearly distinguishable from nuclei of other cells like erythrocytes and fibroblasts and only centralized nuclei in cardiac myocytes (which are more elongated than nuclei from erythrocytes) were counted. Furthermore, scanning electron microscopy (SEM) was used to confirm these findings in the smaller zebrafish hearts (Fig. 3). Future research is warranted in order to assess further how the cardiac myocytes adapt to CCH in the fish heart.

Gene expression changes in the heart of adult zebrafish under chronic constant hypoxia (CCH)

In this study, we used microarrays for the transcriptional profiling of up and downregulated genes in response to hypoxia in the zebrafish heart. We identified 376 genes that were differentially expressed under hypoxic conditions, out of which 116 genes showed a decrease in gene expression (30.9%) in comparison to 260 genes which showed increased expression levels (69.1%).

All 376 differentially expressed genes, including the ones with oligo sequences which could not be annotated so far (referred therein either as transcribed locus or zebrafish clone) are shown in the complete file (Supplemental Table 1). Functional groups are color coded and if possible, gene functions are briefly summarized and OMIM links given.

Functional groups of differentially expressed genes in the heart

We have clustered the differential expressed genes according to known functions (Table 2). Genes can have more than one particular function assigned, so some genes can appear in more than one group. In addition, a gene ontology analysis was performed using eGOn to determine gene enrichment and overrepresentation in the three categories of molecular function, biological processes and cellular components (Supplemental Data 1 and 2).

Proteinbiosynthesis (Translation): In the group linked to proteinbiosynthesis, only three genes showed decreased expression under hypoxic conditions. All three were found to be part of the mitochondrial translational machinery. In contrast, 9 non mitochondrial genes linked to proteinbiosynthesis showed all increased expression (Table 2).

Metabolism: The group with metabolic genes contains 11 repressed and 28 genes with enhanced expression (Table 2). The repression majorly involves metabolic genes linked to β-oxidation and lipid metabolism. Among the
Fig. 2  Histological changes of zebrafish and cichlid hearts after exposure to chronic constant hypoxia. a shows zebrafish hearts that were dissected, sectioned and stained with a hematoxylin–eosin staining with A, B and E representing normoxic and C, D and F hypoxic conditions. Cell nuclei are seen in dark (dark blue in online version) and cell cytoplasm in light (pink in online version). Pictures A, C, E and F have the same magnification (10×). Images B and D represent a 20× magnification of cardiac muscle (D). Abbreviations used are: a atrium; v ventricle; vo ventricular outflow tract and ca conus arteriosus. b (A–F) corresponds to sections of cichlid hearts, which were treated the same way as the ones above from zebrafish and G and H show cichlid hearts which have been perfused prior to dissection and were stained with either hematoxylin–eosin (A–F) or Azan blue (G, H). In A, B, C and G pictures of normoxic conditions are shown and D, E, F and H represent the corresponding hypoxic conditions. Similar results for both the zebrafish and the cichlid hearts were observed in three independent experiments.
metabolic genes with enhanced expression, are pyruvate kinase and aldolase which both are key enzymes for glycolysis indicating a shift from aerobic to anaerobic metabolism induced by hypoxia.

Protection against reactive oxygen species (ROS): The group of genes important for protection against ROS contains 6 genes which are all enhanced under hypoxia.

Apoptosis: We found four genes linked to programmed cell death to be enhanced by hypoxia. Two of these, the death receptor 5 (DR5) and the BNIP3 homologue, are considered to be pro-apoptotic, whereas apoptosis inhibitor 5 and Bax inhibitor have been shown to have anti-apoptotic properties (Tewari et al. 1997; Xu and Reed 1998).

Growth regulation: In the group of genes linked to growth regulation, we found 9 genes to be repressed and 8 genes with enhanced expression. Within the group of 8 repressed genes, we found 5 antiproliferative genes: spry4 and dual specificity phosphatase 5 both inhibit mitogen-activated kinases (MAPK), SOCS3 binds and inhibits Janus kinases (JAK) and thereby prevents STAT3 activation. BTG2 which is important in the G1/S transition and TIEG2 is a transcriptional repressor with antiproliferative functions. Although some genes involved in cell proliferation, like the transcription factor c-fos were repressed by hypoxia (see also “Discussion”), the regulation of the vast majority (13 of 17) of identified genes in this group suggests stimulation of proliferation (Table 2).

Inflammation: 14 genes involved in the inflammatory response were identified to be differentially regulated and all showed increased expression upon hypoxia treatment.

Heart-related function: Several genes linked to cardiac hypertrophy, cardiomyopathy (disease of the heart muscle) and cardiac infarction were identified in this study (see Table 2 and “Discussion”).

Muscle-related function: Several sarcomeric genes linked to hypertrophy showed decreased expression under hypoxic conditions (Table 2 and “Discussion”).

Development: The genes in this group were all found to be upregulated by hypoxia among them the gene for notch-2 and notch-3. Notch receptors are transmembrane receptors with essential roles in development including heart development.

Transport (cellular and vascular): A heterologous group containing the gene for embryonic hemoglobin beta e2, important for oxygen transport and hemopexin which is important for heme and iron transport and was found to be upregulated.

Angiogenesis: It is well known that hypoxia via the hypoxia inducible factor 1 (HIF1α) pathway leads to angiogenesis. In the zebrafish heart under CCH, we observed increased expression of HIF1α as well as fibrinogen-α and -γ. Fibrinogen-α has been shown to stimulate HIF1α and VEGF expression and thereby induces angiogenesis (Shiose et al. 2004).

Expression changes of known hypoxia responsive genes Examples for regulation of known hypoxia responsive genes are HIF1α, insulin growth factor-2 (IGF-2), insulin growth factor binding protein 1 (IGFbp1) and caveolin 3. The transcription factor Hif1α is pivotal in the cellular response to hypoxic stress (Semenza 1999). We further observed increased expression of the egl nine homolog, a gene which was shown to be induced by hypoxia through...
| Table 2  Functional groups of differentially expressed genes |
|---------------------------------|
| UniGene | GeneBank | Fold | Gene name |
|**Upregulated genes** |
| Angiogenesis |
| Dr.11575 | NM_173244 | 2.3 | T-cell acute lymphocytic leukemia 1; TAL1 |
| Dr.845 | BG729013 | 2.8 | Fibrinogen alpha-alpha-E chain |
| Dr.4907 | BC045868 | 4.2 | Fibrinogen, gamma polypeptide |
| Apoptosis |
| Dr.15862 | AF493987 | 2.1 | BCL2adenovirus E1b 19 interacting protein3 |
| DrAfx.1.39 | AF302789 | 2.3 | Death receptor |
| Dr.20106 | AI722277 | 2.8 | Apoptosis inhibitor 5 |
| Dr.4039 | BQ480688 | 21.8 | BAX inhibitor 1 |
| Cell adhesion |
| Dr.6007 | NM_131820 | 2.9 | Cadherin 1 |
| Dr.25140 | BQ262802 | 3.3 | Tumor-associated calcium signal transducer |
| Dr.4409 | BC049036 | 4.4 | CD9 antigen |
| Dr.25140 | BQ262802 | 7.7 | Tumor-associated calcium signal transducer glycoprotein |
| Development |
| Dr.11575 | NM_1732 | 2.3 | T-cell acute lymphocyte leukemia 1 (tal 1) |
| Dr.23348 | BE201653 | 2.6 | Bone morphogenetic protein 3b; (bmp 3) |
| Dr.25405 | BC013923 | 2.8 | SOX2 SRY-box 2 |
| Dr.6382 | AW165053 | 2.9 | Hedgehog-interacting protein |
| Dr.10879 | U97669 | 3.0 | NOTCH3 Notch homolog 3 (Drosophila) |
| Dr.15055 | BC050172 | 3.6 | Chemokine receptor 4a |
| Dr.6787 | BI533426 | 5.5 | Noelin |
| Dr.16720 | BI980847 | 6.3 | notch 2 |
| Disease related |
| Dr.6349 | AW116668 | 2.4 | Eparin cofactor II |
| Dr.21064 | BC046075 | 4.5 | 4hydroxyphenylpyruvate dioxygenase HPD |
| Dr.12584 | NM_131211 | 5.4 | Gata binding protein 3 (GATA3) |
| Dr.3530 | AI497545 | 79.3 | Prion protein (prp) gene |
| Growth regulation |
| Dr.8145 | NM_13143 | 2.2 | Insulin like growth factor 2 (IGF-2) |
| Dr.7609 | BI475857 | 2.4 | Prolactin receptor |
| Dr.8285 | NM_13136 | 2.4 | Mad homolog 2 |
| Dr.8947 | CD594735 | 2.5 | Spint 2 |
| Dr.822 | BM184127 | 2.5 | Spint 2 |
| Dr.3563 | CD014488 | 2.8 | Tetraspan membrane protein IL-TMP |
| Dr.8587 | NM_17328 | 2.9 | Insulin-like growth factor binding protein 1 |
| Dr.2596 | BM342901 | 3.2 | Cyclin I |
| Dr.8587 | AL910822 | 3.4 | Insulin-like growth factor binding protein 1 |
| Dr.26458 | BC053206 | 5.6 | m-ras |
| Heart related |
| Dr.15088 | BM181749 | 4.3 | Lectin galactoside-binding soluble 1; (galectin10-like 3) |
| Dr.4867 | AI496840 | 5.5 | Haptoglobin |
| Dr.3585 | AY049731 | 6.6 | Angiotensinogen |
| Dr.2452 | BQ284848 | 4.3 | Complement component C9 |
| Dr.18453 | BC044525 | 4.8 | Uridine phosphorylase |
| Dr.3025 | BG738204 | 2.7 | Alpha-2 macro-globulin; A2MG |
| Inflammation |
| Dr.12491 | BI672168 | 2.1 | Complement C4–2 |
### Table 2 continued

| UniGene | GeneBank | Fold | Gene name |
|---------|----------|------|-----------|
| Dr.4047 | NM_131627 | 2.3  | Small inducible cytokine A (scyba) |
| Dr.5053 | NM_131723 | 2.3  | Kruppel-like factor 4 |
| Dr25207 | X06465 | 2.5  | Complement component 8, gamma polypeptide |
| Dr.6845 | K02765 | 2.9  | C3 complement component 3 |
| Dr.5741 | BU710482 | 3.2  | Complement component b fb |
| Dr.7722 | BI878414 | 3.5  | Complement C3-H1 |
| Dr.22244 | AW019781 | 3.6  | Complement C1s |
| Dr.22133 | AW076768 | 3.7  | c1rs-A and c1rs-B |
| Dr.5528 | AI497212 | 4.2  | Complement component C9 |
| Dr.2452 | BQ284848 | 4.3  | Complement component C9 |
| Dr.1730 | AI721528 | 4.8  | cfr-B complement control protein factor I-B |
| Dr.2452 | BM778002 | 5.8  | Complement component C9 |
| Dr.20291 | BM036389 | 6.5  | Complement C3-S |
| Dr.190 | NM_131338 | 7.9  | Complement component factor B |
| Dr.1192 | AB071601 | 2.0  | Lipocalin-type prostaglandin D synthase-like protein |

**Metabolism**

| UniGene | GeneBank | Fold | Gene name |
|---------|----------|------|-----------|
| Dr.9492 | B1882244 | 2.0  | Sulfide dehydrogenase like |
| Dr.15574 | BM571467 | 2.1  | Hypoxanthine phosphor-ribosyltransferase 1 |
| Dr.3332 | AI943053 | 2.2  | Angiopoietin 5 |
| Dr.16130 | CD014898 | 2.3  | Alcohol dehydrogenase 8 b |
| Dr.3959 | BI43001 | 2.5  | 5'-nucleotidase |
| Dr.22205 | AW019477 | 2.6  | Oxidoreductase |
| Dr.1699 | A1667249 | 2.7  | Pyruvate kinase |
| Dr.5504 | BI879550 | 3.2  | Cystathionine-beta-synthase |
| Dr.1202 | AJ245491 | 3.9  | Apolipoprotein A-I |
| Dr.4111 | BC053267 | 4.2  | Fructose-1,6-bisphosphatase 1 |
| Dr.18834 | AW019321 | 4.2  | Urate oxidase |
| Dr.19224 | BC050167 | 4.3  | Aldolase b |
| Dr.4938 | NM_131645 | 4.4  | Fatty acid desaturase 2 |
| Dr.12654 | BC046901 | 14.8 | ELOVL family member 6, |
| Dr.5488 | AI545593 | 17.3 | Apolipoprotein A-IV |

**Proteolysis**

| UniGene | GeneBank | Fold | Gene name |
|---------|----------|------|-----------|
| Dr.20934 | AF541952 | 2.6  | Trypsin precursor |
| Dr.3025 | BG738204 | 2.7  | Alpha-2-macroglobulin |
| Dr.22139 | AW018965 | 3.0  | Alpha-1-antitrypsin |
| Dr.25331 | AI658072 | 4.1  | Alpha-2-macroglobulin-2 |
| Dr.12602 | NM_139180 | 4.3  | Lysozyme |
| Dr.1605 | BM185388 | 4.4  | Protease inhibitor 1 |
| Dr.17459 | CD586837 | 4.8  | Inter-alpha-trypsin inhibitor heavy chain H3 |
| Dr.3073 | AI585030 | 5.0  | Serine protease inhibitor alpha 1 |
| Dr.26371 | AI667676 | 5.4  | Prostasin |
| Dr.3025 | BM503427 | 5.6  | Alpha-2-macroglobulin-1 |
| Dr.3025 | BM316867 | 6.5  | Alpha-2-macroglobulin-2 |
| Dr.2960 | X67055 | 3.5  | ITIH3 pre-alpha (globulin) inhibitor, H3 polypeptide |
| Dr.25379 | BI326783 | 6.7  | Alpha-2-macroglobulin |
| Dr.4797 | AI959534 | 7.4  | 26–29 kD-Proteinase protein |
### Table 2 continued

| UniGene | GeneBank | Fold | Gene name |
|---------|----------|------|-----------|
| **ROS protection** | | | |
| Dr.20068 | NM_131075 | 2.1 | Metallothionein (mt) |
| Dr.5399 | AY957765 | 2.3 | Biliverdin I Beta Reductase |
| Dr.14058 | CD015351 | 3.5 | Glutathione S-transferase theta 1 |
| Dr.25160 | BC049475 | 5.9 | Metallothionein 2 |
| Dr.3613 | BC048037 | 6.0 | Ceruloplasmin |
| Dr.4905.1 | BC045464 | 6.5 | Uncoupling protein 4 |
| **Signal transduction** | | | |
| Dr.9852 | AW826425 | 2.1 | CAM kinase 1 |
| Dr.8591 | BM186508 | 2.9 | Rho guanine nucleotide exchange factor 10 |
| Dr.6236 | AW115973 | 3.1 | Rho guanine nucleotide exchange factor 5 |
| Dr.1267 | BC051157 | 3.4 | Phospholipase C delta |
| Dr.22129 | BC016668 | 3.9 | RRAGC Rag C (Ras-related GTP binding C) |
| Dr.7255 | AW116479 | 4.4 | Protein phosphatase 1, |
| Dr.4453 | BC044421 | 5.8 | Phosphoprotein phosphatase |
| **Translation** | | | |
| Dr.13234 | BM036471 | 2.0 | Ribonuclease P |
| Dr.382 | CB363830 | 2.1 | Nucleolin |
| Dr.6949 | AW078116 | 2.1 | RNA 3'-terminal phosphate cyclase-like protein (HSPC338) |
| Dr.13563 | BB90792 | 2.3 | Methionyl aminopeptidase 2 |
| Dr.26328 | AL723696 | 2.3 | Eukaryotic translation initiation factor 4A, |
| Dr.17693 | BQ078285 | 3.7 | 40 S ribosomal protein S6 |
| Dr.20270 | B1674050 | 5.9 | Ribosomal protein L12 |
| Dr.25224 | CD015330 | 20.4 | Ribosomal protein L12 |
| Dr.12439 | BM533848 | 17.5 | Heterogeneous nuclear ribonucleoprotein K |
| Dr.12439 | BM533848 | 24.2 | Heterogeneous nuclear ribonucleoprotein K |
| Dr.14821 | BM071714 | 33.8 | Heterogeneous nuclear ribonucleoprotein K |
| Dr.12502 | BQ284666 | 40.7 | Heterogeneous nuclear ribonucleoprotein K |
| Dr.12439 | BM534432 | 40.9 | Heterogeneous nuclear ribonucleoprotein K |
| Dr.12439 | BQ616930 | 45.4 | Heterogeneous nuclear ribonucleoprotein K |
| **Transport** | | | |
| Dr.1084 | BQ109772 | 3.0 | Clathrin coat assembly protein AP19 |
| Dr.5562 | X04506 | 3.0 | APOB apolipoprotein B (including Ag(x) antigen) |
| Dr.13231 | BM778646 | 4.2 | Solute carrier family 22 |
| Dr.30444 | AY329629 | 4.3 | Embryonic globin beta e2 |
| Dr.24250 | AF489105 | 2.0 | Uroporphyrinogen III synthase |
| Dr.10343 | NM_131687 | 4.7 | Na+K+ transporting, alpha 1a.2 polypeptide |
| Dr.7634 | AW115757 | 11.3 | Hemopexin |
| **Downregulated genes** | | | |
| Angiogenesis | | | |
| Dr.26411 | BQ783571 | −8.9 | Fast muscle troponin I |
| Dr.15501 | BM316040 | −2.1 | Similar to CYR6 HUMAN CYR61 protein precursor, Insulin-like growth factor-binding protein 10 |
| **Cell adhesion** | | | |
| Dr.251 | BQ285646 | −2.3 | Cadherin 11 |
| **Disease related** | | | |
| Dr.22774 | AW280206 | −5.7 | ras-like GTP-binding protein RAB27A |
| Dr.1816 | AL720262 | −4.4 | Ataxin 2-binding protein |
| Dr.9893 | BM036473 | −2.3 | Fibrillarin |
Table 2 continued

| UniGene | GeneBank | Fold | Gene name                        |
|---------|----------|------|----------------------------------|
| Dr.16726 | BI429372 | −2.0 | netrin G1                        |
| Growth regulation |
| Dr.12986 | CA787334 | −5.3 | v-fos                            |
| Dr.12986 | BI881979 | −5.0 | v-fos                            |
| Dr.12986 | BM957279 | −4.5 | v-fos                            |
| Dr.1221  | AW510198 | −4.3 | Pmx-1b (PHOX-1)                  |
| Dr.12986 | BI881979 | −4.2 | v-fos                            |
| Dr.12410 | NM_131826 | −2.4 | Sprout homolog 4                 |
| Dr.6431  | BC049326 | −2.3 | Suppressors of cytokine signaling 3 |
| Dr.6511  | NM_130922 | −2.2 | B-cell translocation gene 2      |
| Dr.3565  | Al601685 | −2.2 | Dual specificity phosphatase 5    |
| Dr.12062 | BC047814 | −2.1 | Epidermal growth factor receptor kinase substrate EPS8 |
| Dr.17286 | BM777144 | −2.0 | Hormone-regulated proliferation-associated 20 kDa protein |
| Dr.9448  | BM156058 | −2.0 | TGF-beta-inducible early growth response protein 2 |
| Heart related |
| Dr.20010 | BQ826502 | −7.0 | ATPase, Ca++ transporting, cardiac muscle (ATP2A1) |
| Dr.1448  | AL717344 | −3.5 | Fast skeletal myosin light chain 1a |
| Dr.20990 | AY033829 AY081167 | −2.4–2.1 | Titin |
| Metabolism |
| Dr.24950 | BC053305 | −4.1 | Creatine kinase CKM3             |
| Dr.9528  | BC045993 | −3.5 | Pyruvate dehydrogenase kinase    |
| Dr.146   | Al477401 | −2.9 | Carnitine O-palmitoyltransferase II |
| Dr.21501 | Al667180 | −2.4 | Short-chain acyl-CoA dehydrogenase |
| Dr.19643 | AL918850 | −2.4 | FabG beta-ketoacyl -reductase     |
| Dr.15059 | BM530407 | −2.2 | Elongation of very long chain fatty acids (Cig30) |
| Dr.21040 | BC045479 | −2.1 | Glucose-6-phosphatase, transport protein 1 |
| Dr.988   | AW154697 | −2.1 | Dodecenoyl-coenzyme A delta isomerase |
| Dr.11971 | BG727588 | −2.0 | Carnitine O-acetyl-transferase    |
| Dr.4777  | AW420997 | −2.0 | Succinate-CoA ligase              |
| Dr.11252 | BC047826 | −2.0 | Creatine kinase, mitochondrial 1  |
| Muscle related |
| Dr.21800 | AI883923 | −5.0 | Myosin binding protein C          |
| Dr.5066  | AF524840 | −3.4 | Alpha-actinin 3                   |
| Dr.24260 | NM_131619 | −3.0 | Myosin, light polypeptide 3       |
| Dr.2914  | BC045520 | −2.5 | Myosin light polypeptide 2; mylz2 |
| Dr.20990 | AY033829 AY081167 | −2.4–2.1 | Titin |
| Dr.1435  | AI353817 | −2.0 | Caveolin 3                        |
| Dr.18657 | BQ479700 | −2.1 | Carbonic anhydrase II             |
| Dr.26411 | BQ783571 | −8.9 | Troponin I                        |
| Proteolysis |
| Dr.3581  | BM101561 | −8.3 | Chymotrypsinogen B1               |
| Dr.3581  | BM101561 | −7.5 | Chymotrypsinogen B1               |
| Signal transduction |
| Dr.22841 | Al641080 | −2.4 | Serum deprivation response protein (SDPR) |
| Translation |
| Dr.7939  | AW281840 | −2.7 | Mitochondrial elongation factor G1 |
| Dr.1286  | BM036808 | −2.2 | Mitochondrial ribosomal protein L48 |
| Dr.18218 | AL909921 | −2.1 | Mitochondrial 28 S ribosomal protein S12 |
the Hif1α pathway (Pescador et al. 2005). IGF-2 gene- and protein expression had been shown to be upregulated by hypoxia (Beilharz et al. 1995). It was shown that IGFBP1 also is a hypoxia-inducible gene in zebrafish embryos and it mediates hypoxia-induced embryonic growth-inhibition and developmental-retardation (Kajimura et al. 2005a). Both IGF-2 and IGFBP1 were found upregulated in our study. We observed decreased expression of caveolin 3 in the hearts of zebrafish exposed to CCH, earlier findings showed that chronic myocardial hypoxia led to decreased caveolin-3 protein expression in rabbit hearts (Shi et al. 2000). These findings indicate that the hypoxic conditions used lead to hypoxic stress in the fish heart.

Evaluation of microarray results by quantitative real-time RT-PCR

To further verify our results, we used quantitative real-time PCR for 10 of the transcripts. We confirmed the gene expression changes found in the microarray studies for these 5 up- and 5 downregulated transcripts by this independent method (Fig. 4). The downregulated zebrafish genes tested were: c-fos, phox1, creatine kinase (ckm3), nebulin, titin, and the upregulated genes tested were: metallothionein, pyruvate kinase, apoptosis inhibitor 5, igfbp1 and notch-2. The fold induction values were not always directly comparable to the array data but in all cases induction or reduction was confirmed. Quantitative differences between array data and qPCR results have been reported before (Meijer et al. 2005; Ton et al. 2003; van der Meer et al. 2005).

Assessment of microarray results for the IGF/PI3K/Akt pathway by comparing phospho-Akt levels in cardiac myocytes of normoxic versus hypoxic zebrafish hearts

The IGF/PI3K/Akt pathway is activated by IGFs, which are antagonized by the IGFBP1. To test the effects of the upregulation of both IGF-2 and IGFBP1, we assayed phospho-Akt levels in cardiac myocytes and showed that phospho-Akt levels were not different between normoxic and hypoxic cells. Figure 5 shows cytoplasmic immunohistochemical staining of phospho-Akt. The antibody recognizes phosphorylated and detects the phospho-Akt1/2/3 forms. The incubation times for the primary and secondary antibody as well as the BM Purple were optimized to obtain a good signal to noise ratio. Absorances were linearly related to the time of incubation with the primary and secondary antibodies as well as that of BM Purple. Figure 5d shows for normoxic fish the absorbance values of the phospho-Akt staining in cardiac myocytes as well as skeletal muscle fibers from the tail as a function of the incubation time with BM Purple AP substrate. The absorbance values for the heart are considerably higher than for the skeletal muscle fibers. However, for both cardiac and skeletal muscle the absorbances are linearly related with the incubation time with BM Purple AP substrate and increase at the same relative rate. This implicates that the absorbance of BM Purple after 45 min incubation with BM Purple AP substrate did not reach saturation and therefore provides a semi-quantitative estimate.
of the phospho-Akt content in the cardiac myocytes. Absor- 
bances of staining for phospho-Akt in normoxic and hyp-
oxic cardiac myocytes were not shown to be signi-
cantly different (Fig. 5E, P < 0.48), which indicates that hypoxia 
did not change the activation of the Akt pathway in the 
cardiac myocytes.

Discussion

In the aquatic environment, oxygen concentrations can 
often vary, and being able to adapt to changes in oxygen 
levels can be advantageous for the survival of aquatic ani-
mals. This might be in part the reason why some teleosts 
have developed the ability to withstand extreme hypoxic 
conditions.

In this study, we have focused on the long-term response 
to hypoxia in the fish heart. The hypothesis is that the 
zebrafish heart, in contrast to most mammalian hearts, 
which are characterized by relative intolerance to injury or 
the lack of oxygen, are able to adapt to extreme hypoxic 
conditions.

We showed that chronic hypoxia of zebrafish caused a 
smaller ventricular outflow tract, reduced lacunae and 
increased cardiac myocyte densities in the heart. These 
findings suggest that hypoxia induced an increase of the 
cardiac myocyte volume or at least did not result in a loss of 
cardiac myocytes. This is in contrast to mammals where 
tissue hypoxia in chronic heart failure leads to apoptosis 
and considerable losses of cardiac myocytes (see for 
review, Sabbah et al. 2000a). In mammals, compensation 
for this loss of cardiac myocytes occurs mainly by hypertro-
phy of the remaining cardiac myocytes (Ostadal and Kolar 
2007), although regeneration of myocardium by prolifera-
tion of cardiac myocytes may occur also but to a limited 
extend (Beltrami et al. 2001). Our assay for phospho-Akt did 
not show any enhancement of Akt activity in response to 
the CCH, suggesting a lack of hypertrophic signaling via 
the phosphotadilinositol 3 kinase pathway. However, the 
density of cardiac myocyte nuclei increased by 50% during 
CCH, which indicates substantial proliferation of cardiac 
myocytes and/or nuclear hyperplasia. Recently, it has been 
shown that the zebrafish heart has the ability to regenerate 
from mechanical cardiac injury by proliferation of cardiac 
myocytes (Poss et al. 2002). If the zebrafish heart responds 
to chronic hypoxia in a similar way as to mechanical dissec-
tion, this will be beneficial in preventing apoptosis of car-
diac myocytes as the diffusion distance for oxygen are not 
increased as during hypertrophy, which will help to prevent 
the development of anoxic cores in the cardiac myocytes 
(Des Tombe et al. 2002; van der Laarse et al. 2005). In the 

![Fig. 5](https://example.com/fig5.png)

**Fig. 5** Effects of chronic constant hypoxia of the activation of Akt in zebrafish cardiac myocytes. Immunohistochemical staining of phospho-Akt in cardiac myocytes of zebrafish raised under hypoxic a or normoxic b conditions. Specificity is shown by the control sections obtained from normoxic fish which were not incubated with primary antibody against phospho-Akt c. For both cardiac myocytes and skeletal muscle fibers, the absorbance of the BM Purple is linearly increasing at the same relative rate with the incubation time with BM Purple AP substrate at the same relative rate d. Mean absorbances of phospho-Akt staining (+S.E.M.) from normoxic and hypoxic cardiac myocytes was not different e)
investigation of the mechanisms underlying the general morphological adaptations of teleost fish heart in response to CCH requires a nuclear stain in combination with a clear membrane stain, which allows determination of cell sizes in addition to counts of cardiac myocyte nuclei.

We were interested in the underlying gene expression changes of these adaptations. We found gene regulations in a transcriptional network of the serum response element (SRE), which are opposed to the ones described in mammals. In the zebrafish heart hypoxia repressed c-fos and phox1 expression. Both genes are important in the same transcriptional network, phox1 can transduce serum-responsive transcriptional activity to the c-fos (SRE) by interacting with serum response factor (SRF) (Simon et al. 1997). In mammals many studies showed increased expression of c-fos by hypoxia, e.g. in rats hypoxia induces c-fos expression in the LV and RV (Deindl et al. 2003) and the same held true in tissue culture cells (Webster et al. 1993). Interestingly, repression of c-fos by hypoxia has also been shown for the short-term response to anoxia in anoxia tolerant turtles (Greenway and Storey 2000). Is it possible that repression of c-fos and phox-1 are important adaptations in hypoxia tolerant animals?

Several novel gene expression changes induced by CCH have been identified in this study (Table 2 and Supplemental Table 1). An example is the two Notch receptors, notch-2 and notch-3, whose expression were induced by CCH. Notch receptors have been shown to be important for heart patterning and differentiation (Armstrong and Bischof 2004), cell fate determination and self renewal of stem cells (Androutsellis-Theotokis et al. 2006; Silvia Bianchi 2006) but so far have not been linked to the hypoxic response.

Several genes with links to human heart pathology were found to be upregulated in this study. Among these were two markers for myocardial infarction, complement component C9 and haptoglobin. C9 is used as a marker for myocardial infarction (Doran et al. 1996) and a polymorphism for haptoglobin predicts 30-day mortality and heart failure in patients with diabetes and acute myocardial infarction (Levy et al. 2002). We also observed increased expression of fetuin-z, a circulating calcium- regulatory glycoprotein that inhibits vascular calcification. Low fetuin-z levels have been associated with heart failure in mice (Merx et al. 2005). Upregulation of fetuin-z in the zebrafish heart could help to better tolerate CCH. Selenoprotein P (SEPP1) is a heparin-binding protein that appears to be associated with endothelial cells and has been implicated as an oxidant defense in the extracellular space. Human populations that are selenium deficient are susceptible to the development of Keshan disease, a potentially fatal form of cardiomyopathy (Nezelof et al. 2002). SEPP1 expression was found to be increased in our study indicating a potential protective mechanism against oxidative stress in the heart. This was further supported by the increased expression observed for several genes important for the protection against reactive oxygen species (ROS) (Table 2). Among them were metallothionein and glutathione S-transferase, which are both known ROS scavengers. Metallothionein has further been described to be protective against hypoxia-induced apoptosis when overexpressed (Wang et al. 2001). Our findings suggest ROS protection as an important adaptation to CCH in the fish heart.

Furthermore, our data show for the first time that CCH simultaneously induced upregulation of IGF-2 and the insulin-like growth factor binding protein 1 (IGFbp1). In isolated cardiomyoblasts, angiotensin II stimulates IGF-2 expression which is involved in the induction of apoptotic signaling in rat hearts (Lee et al. 2006). We have observed here both angiotensin and IGF-2 upregulation (Table 2). The upregulation of IGFbp1 seems to be a general response to hypoxia in zebrafish embryos, where it mediates hypoxia-induced embryonic growth retardation and developmental delay (Kajimura et al. 2005b). IGFbp1 is a secreted protein, which binds to IGFs in the extracellular environment and prevents receptor activation (Florini et al. 1996; Stewart and Rotwein 1996). Here, IGFbp1 by binding IGF-2, may have prevented both cardioprotective as well as apoptotic effects of enhanced IGF-2 expression. To test the effects of the upregulation of both IGF-2 and IGFbp1, we assayed phospho-Akt levels in cardiac myocytes and showed that phospho-Akt levels were not different between normoxic and hypoxic cells. This suggests that either hypoxia-induced changes in mRNA expression did not occur at the protein level or the effects of increased IGF-2 expression on the IGF-1 receptor (which can be activated by IGF-2) were blunted by the upregulation of the IGFbp1. The fact that we did not observe increased phospho-Akt levels suggests a lack of hypertrophic signaling as enhanced phospho-Akt is required for cardiac hypertrophy (DeBosch et al. 2006). This may be beneficial for the heart as hypertrophy of cardiac myocytes implicates an increase in the diffusion distance for oxygen which may cause the development of anoxic cores (Des Tombe et al. 2002), release of cytochrome c (van Beek-Harmsen and van der Laarse 2005) and production of ROS (Lee et al. 2006; Powers et al. 2007) and eventually causing apoptosis of cardiac myocytes. The importance of the IGF-2/IGFbp1 signaling in the protection of the zebrafish heart and its underlying mechanisms remain to be determined.

The gene expression changes we observed for the specific response to CCH in the fish heart were very different from the responses we observed in an earlier study in the gills (van der Meer et al. 2005). Under the same criteria as used in this study, we found that the majority of differentially regulated genes in the gills showed a decrease in gene expression (68.1% or 250 genes in total) in comparison to
genes, which showed increased expression levels (31.9% or 117 genes in total). This is opposed to the heart where 69.1% of differentially regulated genes (260 genes in total) showed increased and 30.9% (116 in total) decreased expression levels. Many genes linked to protein synthesis showed a similar trend, and were downregulated in the gills (van der Meer et al. 2005) and upregulated in the heart (Table 2). The major differences observed in gene regulation between heart and gills point to very tissue specific responses to CCH. A list of genes identified in both studies, as identified by a direct comparison based on Accession numbers and old Unigenes numbers are shown in Supplemental Data 3.

Teleosts have developed specific phenotypic adaptations to low oxygen, due to the natural occurrence of hypoxia in the water environment. Here, we identified the changes in gene expression as well as associated morphological changes of the fish heart to hypoxia. We observed repression of c-fos, which differs compared to the described increase in expression in mammals (Deindl et al. 2003). Other changes found like upregulation of the two Notch receptors have not been described before to the best of our knowledge. Similarly, the simultaneous increase in expression of IGF-2 and IGFbp1 has not been shown. The changes identified here can contribute to the ability of teleosts to adapt to severe hypoxia (for example, the upregulation of fetuin-α and sepp1 levels). Future functional studies are warranted to validate the role of the identified genes in cardiac protection to hypoxia.

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