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<td><strong>Author(s)</strong></td>
<td>Alli Shaik, Asfa; Wee, Sheena; Li, Rachel Hai Xia; Li, Zhen; Carney, Tom J.; Mathavan, Sinnakaruppan; Gunaratne, Jayantha</td>
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Functional Mapping of the Zebrafish Early Embryo Proteome and Transcriptome

Asfa Alli Shaik,†⊥ Sheena Wee,†⊥ Rachel Hai Xia Li,† Zhen Li,‡ Tom J. Carney,†§ Sinnakaruppan Mathavan,‡ and Jayantha Gunaratne§⊥‡

ABSTRACT: Zebrafish is a popular system for studying vertebrate development and disease that shows high genetic conservation with humans. Molecular level studies at different stages of development are essential for understanding the processes deployed during ontogeny. Here, we performed comparative analysis of the whole proteome and transcriptome of the early stage (24 h post-fertilization) zebrafish embryo. We identified 8363 proteins with their approximate cellular abundances (the largest number of zebrafish embryo proteins quantified thus far), through a combination of thorough deyolking and extensive fractionation procedures, before resolving the peptides by mass spectrometry. We performed deep sequencing of the transcripts and found that the expressed proteome and transcriptome displayed a moderate correlation for the majority of cellular processes. Integrative functional mapping of the quantified genes demonstrated that embryonic developmental systems differentially exploit transcriptional and post-transcriptional regulatory mechanisms to modulate protein abundance. Using network mapping of the low-abundance proteins, we identified various signal transduction pathways important in embryonic development and also revealed genes that may be regulated at the post-transcriptional level. Our data set represents a deep coverage of the functional proteome and transcriptome of the developing zebrafish, and our findings unveil molecular regulatory mechanisms that underlie embryonic development.

KEYWORDS: Zebrafish, proteomics, transcriptomics, early embryo, development, functional mapping

INTRODUCTION

The zebrafish (Danio rerio) is an attractive experimental model organism for exploring the molecular mechanisms of vertebrate development.1 Mutant phenotypes commonly emulate the pathogenesis/phenotype of human diseases and disorders, hence making it a premier model for genetic and phenotypic analysis.2-4 Genetic screens have also broadened our understanding of the various factors that control cell differentiation and fate as well as organogenesis, allowing us to chart the sequential events involved during the transition from embryo to adult.5 There are more than 26,000 coding regions in the zebrafish genome, many of which are orthologous to those in humans.6 7 To understand the complex interplay among the expressed genes, large-scale analysis that extensively captures the expression variation at the mRNA and protein levels is important.8-11 A recent systematic analysis identified a total of over 56,000 transcripts, including alternative splice variants, during zebrafish embryogenesis.12 However, the maximum number of proteins that have been identified to date has been limited to only 5267 and 8475 in embryos and adults, respectively.13,14

Although transcript abundance can provide valuable information on the status of the cell at any point in time, proteins are the fundamental biological effectors that orchestrate key events within the cells. Gene expression patterns derived from large-scale transcriptomics, including those involving microarrays and RNA-seq, have been routinely used to estimate protein abundance. However, only a modest correlation has been observed between mRNA and protein levels across different species from yeast through higher eukaryotes.15-18 A recent comparative transcriptomic and proteomic study in the late-stage zebrafish embryo suggests that such differences between transcript and protein levels may underlie important translational and post-translational regulatory mechanisms.19 Hence, the need for a thorough representation of the proteome is increasingly recognized.20

Currently, mass spectrometry-based shotgun proteomics is the only available high-throughput method for identification and quantification of the whole proteome. In recent years,
global proteomic profiling of zebrafish adults and embryos has been extensively carried out, primarily to understand developmental processes as well as to recapitulate disease mechanisms. With zebrafish being recognized as a powerful model for chemical toxicity and drug safety assessment, proteomics-based methods along with transcriptomics are increasingly applied for large-scale system-wide studies. Although applications of quantitative proteomic approaches have been limited in zebrafish, recent studies used stable isotope-labeled zebrafish for studying cardiac morphogenesis and profiling various organs in the adult, indicating the possibility of performing large-scale quantitative proteomics studies in zebrafish in the future.

Despite the gaining popularity of proteomic studies in zebrafish, thorough protein identification is largely dependent on sample complexity and the dynamic range of the proteins within the sample. However, in the case of zebrafish, this endeavor is even more challenging owing to the high proportion of yolk proteins, particularly during the early stages of development. Hence, most of the proteomic studies in zebrafish have been performed in late-stage embryos or adults. Early embryonic stages are highly dynamic in nature and are marked by events that accompany cell differentiation and morphogenesis. A thorough representation of the proteome during these stages is essential to map the key biological events that occur during embryogenesis.

Here, we report a comprehensive map of the quantitative proteome profile of early stage zebrafish (24 h post-fertilization embryo) containing 8363 proteins, the highest number of proteins reported so far for early embryonic stages. We establish that the protein functions are linked to their abundances, wherein high-abundance proteins are predominantly associated with cellular core functions and low-abundance proteins perform regulatory functions that mediate development. The high coverage proteome was also compared to the corresponding transcriptome derived from the same early embryonic stages to provide a comprehensive functional map of the quantified proteome and transcriptome. The integrative approach identified biological processes that are modulated differently by transcript and protein levels in the early stage zebrafish embryo.

**MATERIALS AND METHODS**

**Sample Preparation**

Adult zebrafish were maintained on a 14 h light/10 h dark cycle at 28 °C in the AVA (Singapore) certified IMCB zebrafish facility. Zebrafish embryos were obtained through crosses of...
114 TL/AB hybrid parents, and zebrafish were raised at 28 °C in 115 zebrafish embryo medium. At 24 h post-fertilization (hpf), 116 approximately 1000 embryos were dechorionated with Pronase 117 and subsequently washed extensively in embryo medium. 118 Deyolking was performed largely as per Link et al.,35 using 119 three washes in calcium-free Ringer’s solution with mechanical 120 disruption through a flame-narrowed glass Pasteur pipet. The 121 resulting cell pellet was lysed by brief sonication in CSH buffer 122 (50 mM Tris-HCl, 250 mM NaCl, 1 mM EDTA, 1% NP40) 123 supplemented with protease inhibitors (Roche). Insoluble 124 material was removed by centrifugation. Protein concentration 125 was determined using the BCA protein assay, reducing agent 126 compatible (Thermo Fisher Scientific).

127 SDS-PAGE and In-Gel Digestion

128 Five-hundred micrograms of lysate, obtained from two 129 biological replicates, was separated on a SDS-PAGE gel from 130 3.5 to 260 kDa using a NuPAGE 4–12% Bis-Tris 1.0 mm, 10 131 well gel (Invitrogen). The gel was cut into 6 bands as shown in 132 Figure 1A, and each band was excised to 1.5 × 1.5 mm² using a 133 scalpel. In-gel digestion was carried out as previously described 134 with minor modifications.36 Briefly, the gel pieces were washed 135 with 3 mL of 50 mM ammonium bicarbonate. Reduction was 136 carried out by the addition of 10 mM DTT, covering the gel 137 pieces, and incubation for 30 min at 56 °C. Alkylation was 138 performed with the addition of 55 mM iodoacetamide and 139 incubation for 20 min in the dark at room temperature. The gel 140 pieces were destained with 6 mL of 50% acetonitrile/25 mM 141 ammonium bicarbonate. In-gel digestion was carried out as previously described 142 with minor modifications.36 Briefly, the gel pieces were washed 143 with 3 mL of 50 mM ammonium bicarbonate. Reduction was 144 carried out by the addition of 10 mM DTT, covering the gel 145 pieces, and incubation for 30 min at 56 °C. Alkylation was 146 performed with the addition of 55 mM iodoacetamide and 147 incubation for 20 min in the dark at room temperature. The gel 148 pieces were destained with 6 mL of 50% acetonitrile/25 mM 149 ammonium bicarbonate for 10 min. Six milliliters of 100% 150 acetonitrile was used to shrink the gel pieces, which was done 151 twice for 10 min each. One to two milliliters of 13 ng/μL 152 sequencing-grade trypsin (Promega) was added to each well, 153 and the gel pieces were allowed to swell for 60 min at 4 °C 154 before enough 25 mM ammonium bicarbonate was added to 155 cover the gel pieces. The samples were incubated for 3 h at 37 156 °C. All supernatants were collected by centrifugation. One and 157 a half milliliters of 5% formic acid was added to each well 158 followed by 1.5 mL of 100% acetonitrile for peptide extraction. 159 Both steps were repeated.

160 Off-Gel Isoelectric Focusing

161 Off-gel isoelectric focusing (IEF) was carried out using a 3100 162 OFFGEL fractionator (Agilent) as described in the manufac- 163 turer’s manual with slight modifications. In short, 13 cm IPG 164 strips for the pH range of 3–10 (GE Healthcare) were used, 165 resulting in 12 peptide fractions. The concentration of glycerol 166 and IPG buffer pH 3–10 (GE Healthcare) was halved in the 167 peptide OFFGEL stock solution as described previously.37 The 168 voltage gradient during the run was 250 V for the first hour 169 followed by a gradient of up to 1000 V over the next 2 h and 170 1000 V for an additional hour. Then, the voltage was increased 171 up to 3000 V over the next 7 h and held at 3000 V until a total 172 voltage of 20 000 V hours was reached. After the run, 30 μL of 173 1% TFA was added to each well for acidification.

174 NanoHPLC–ESI–MS/MS

175 Peptides resulting from the different fractionation methods 176 were desalted using self-packed C18 StageTips.38 The C18 177 StageTip was conditioned with 100 μL of methanol followed by 178 100 μL of 0.1% formic acid at 6000 g for 2 min. The extracted 179 peptides were loaded onto the C18 StageTips and washed with 180 100 μL of 0.1% formic acid. The peptides were eluted with 60 181 μL of 0.1% formic acid/80% acetonitrile. All eluents were dried 182 using a SpeedVac and reconstituted in 12 μL of 0.1% formic 183 acid. A total of 144 IEF fractions (72 fractions per biological 184 replicate) were analyzed in duplicate using an EASY-nLC 185 (Proxeon) coupled to a LTQ Velos (Thermo Fisher Scientific).

186 Samples were directly loaded at 400 nL/min onto a PicoFrit 187 column (HALO, C18, 90 Å, 2.7 μm, 75 um (i.d.) × 100 mm 188 length) (New Objectives). The HPLC gradient was created 189 using buffer A consisting of 2% acetonitrile/0.1% formic acid 190 and buffer B consisting of 80% acetonitrile/0.1% formic acid; 191 buffer B was increased from 0 to 8% over the first 4 min, 192 followed by an increase to 25% over the next 58 min, an 193 increase to 45% over the subsequent 32 min, an increase to 194 70% over the following 10 min, and an increase to 100% over 195 the next 3 min. This condition was maintained for 5 min. Buffer 196 B was then decreased to 5% over the subsequent 3 min and 197 retained at 5% for another 5 min. This results in a HPLC 198 gradient run of a total of 120 min. The flow rate was 250 nL/ 199 min for the first 104 min and 400 nL/min for the last 16 min. 190 MS analysis was online-coupled to the LC using a LTQ Velos 191 with the following settings: MS scans ranging from 300 to 1600 193 m/z, AGC target of 3 × 10⁶, and maximum injection time of 10 194 ms. The 10 most intense ions with an ion intensity above 1000 195 and a charge state excluding one were sequentially isolated to a 196 maximum AGC target value of 4 × 10⁵ for a maximum of 100 197 ms and fragmented by collision induced dissociation (CID) 198 using a normalized collision energy of 35%. A dynamic 199 exclusion list was applied using an exclusion list size of 500, 200 one repeat count, a repeat duration of 45 s, an exclusion 201 duration of 90 s, and mass widths of 1.0 (low) and 1.5 (high). 202 Expiration count was set to 3, and its S/N threshold, to 3.0. 203 Data processing and emPAI Calculation

204 All Velos raw data were first converted to peak lists in the 205 centroid mzXML file format and then to the mgf file format. 206 The conversion was performed with ReAdW.exe (version 207 4.0.2), which is part of the Trans-Proteomic Pipeline (TPP) 208 (version 4.4.0).39

209 A target-decoy database was compiled using Sequence 210 Reverser (part of MaxQuant v1.0.13.13) with the ipli.dan- 211 re.v3.85.fasta downloaded from ftp://ftp.ebi.ac.uk/pub/ 212 databases/IPI/last_release/current/ and 262 contaminant 213 sequences in Sequence Reverser. The final database containing 214 81 476 sequence entries was searched on the mgf peak list files 215 using Mascot (version 2.3).

216 Mascot search parameters were set as follows: full trypsin 217 specificity was required (cleavage after lysine or arginine 218 residues at two peptide termini), two missed cleavages were 219 allowed, carbamidomethylation (C) was set as 219 fixed modifi- 220 cation, and acetyl (protein N-term) and oxidation (M) were 221 set as variable modifications. Peptide charge was set to 2+, 3+, 222 and 4+. Mass tolerance of the precursor ion and the fragment 223 ions was set at 2 and 0.5 Da, respectively. 224

225 All of the mascot search outputs were combined in TPP 225 (version 4.4.0). First, mascot outputs (dat file) were converted 226 to pepxml file format. Then, PeptideProphet with a minimum 227 length of 7aa, a probability of 0.9, and an accurate mass model 228 was applied. iProphet was used to integrate all of the 229 PeptideProphet results.46 Finally, proteins were assembled 230 with PeptideProphet on the iProphet results with a minimum 231 probability of 0.9.

232 The PeptideProphet output with peptide count was used to 233 calculate the relative protein abundances. Relative protein 234 abundances were calculated using the emPAI algorithm as 235 described by Ishihama et al.41
Analysis of Detected Protein Bias

The IPI identifiers of all of the detected proteins were mapped to the corresponding UniProt IDs and subjected to protein parameter analysis in ExPaSy (http://www.expasy.ch). The ProtParam tool was used to calculate the protein length and pI values. For mapping the chromosomal bias of the detected proteins, the Entrez database (ftp://ftp.ncbi.nlm.nih.gov/gene/DATA) and the annotations from the Zv9, as implemented in the UCSC Genome Browser (http://genome.ucsc.edu/cgi-bin/hgTables), were used.

RNA-Seq Analysis

Zebrafish embryos (wild-type AB line) were collected and incubated at 27 °C. Synchronously developing embryos were collected at 24 hpf and frozen in liquid nitrogen. The frozen embryos were used for RNA extraction. Total RNA was extracted using TRIzol reagent (Invitrogen, USA). RNA concentration was determined using a NanoDrop 2000 (Thermo Scientific), and 60 μg of total RNA was used as starting material. The integrity of the RNA samples was determined using an Agilent RNA 6000 Nano chip on an Agilent 2100 Bioanalyzer. The RNA sample with RIN > 9.0 was selected for mRNA purification using the MicroPoly(A) purist kit (Ambion, USA). Five-hundred nanograms of mRNA was applied for RNA library construction with the solid total RNA-seq kit (ABI, USA) according to the manufacturer’s instructions. RNA was sequenced in a SOLiD3 (ABI) platform, generating 50 bp single-ended reads. We generated about 40 million tags for this library and mapped them to the genome (ZV9). The RNA-seq reads were mapped in a strand-specific manner to the reference seq genes (RefSeq), and the expression is presented as reads per kilobase of exon per million reads mapped (RPKM).

Comparison with RNA-Seq Data

The complete RNA-seq data comprised 10 101 transcripts with RPKM abundance greater than 2. The IPI identifiers of the quantified proteome were mapped to Entrez nucleotide identifiers and subsequently to the corresponding genes. Five-thousand two-hundred and fifty four IPI identifiers from the total 8363 quantified proteins could be successfully mapped to a corresponding transcript from RNA-seq. Some of the IPI identifiers could be mapped to more than one gene identifier. Excluding the events of alternative splice variants, a total of 5084 different protein-coding genes could be mapped. The anatomical enrichment of the quantified proteome and transcriptome was carried out using DAVID.42

GO Pathway Analysis-Based Clustering for Protein and Transcript Groups

The high and low protein abundance groups were identified on the basis of the quantile density distribution of the emPAI and RPKM values across the quantified proteome and transcriptome, respectively. Genes in the top 20% quantile (upper) with respect to the abundance values were categorized into high-abundance groups (very high + high cluster), and those in the bottom 20% (lower) comprised the low-abundance groups (very low + low cluster). The remaining genes were considered to be expressed at moderate levels. For analyzing the enrichment across the high- and low-abundance proteins in accordance with the GO terms, biological process, molecular function, and cellular component (BINGO), as implemented in Cytoscape, was used.35 The enrichment was done using hypergeometric testing followed by Benjamini–Hochberg false discovery rate correction. The frequency of over-representation in the high- or low-abundance groups was calculated by comparing against the enrichment across the whole quantified proteome. For integrative proteomic and transcriptomic analysis based on GO categorization (biological process), the genes were first categorized according to their abundance values into different groups. The GO enrichment along with their p values was obtained for each of the groups and then filtered to retain only those groups that were significantly enriched (p value <0.05) in at least one of the analyzed groups. The filtered p values were log-transformed and z-score normalized before being subjected to hierarchical clustering based on Euclidean distance and average linkage.

MicroRNA Prediction

Genes that displayed low protein levels and high mRNA levels were analyzed for potential microRNA (miRNA) regulation using TargetScanFish version 6.2,44,45 Only those predicted miRNA families with a target score (total context score) ≤ −0.3 were considered to be reliable. For genes with multiple 3’UTRs (untranslated region), the predictions were specifically carried out on those that are curated to be expressed at 24 hpf developmental stage.

Network Analysis

The proteins in the high- or low-abundance groups that could be distinctly mapped to corresponding transcripts were used for network reconstructions. Protein–protein interactions, as described in reactome functional interaction (reactome FI) in Cytoscape visualization software and in GeneGO MetaCore, were used to unravel the connectivity between the protein groups.66,67 Zebrafish shares many orthologous genes and pathways with other vertebrate species and hence the human orthologous proteins corresponding to the zebrafish genes were identified from ZFIN (http://zfin.org/) and InParanoid (http://inparanoid.sbc.su.se/) databases and used in network analysis.48 Only interactions between the quantified proteins were retained, and other linker candidates (not in our data set) were excluded. The direct interactions from MetaCore were downloaded and parsed into Cytoscape as a SIF (simple interaction file) network. This was combined with the functional interaction network derived from reactome FI, and subsequent pathway enrichment was performed for the combined network. The densely connected regions in the network were identified using molecular complex detection (MCODE) algorithm.49 The highest-ranking modules were extracted and visualized. MCODE could not be successfully applied to the low-abundance protein groups owing to the less-dense nature of the network. Hence, the clusters were visualized purely on the basis of their significant pathway enrichment.

RESULTS

Extensive Analysis of Zebrafish Embryo Proteome

To generate an extensive map of the zebrafish proteome, lysates were obtained from deyolked embryos representing the 24 h post-fertilization (hpf) developmental stage and resolved on a 1D SDS-PAGE gel. Deyooling ensures that high-abundance yolk proteins that would otherwise interfere with the deep mining of zebrafish embryos are depleted.33 After deyooling, the sample still consists of a complex mixture of proteins. In order to reduce this complexity, we carried out extensive fractionation of the deyoolked protein mixture using 1D SDS-
PAGE proteome level fractionation followed by off-gel isoelectric focusing (IEF) of the tryptic peptides from each gel band, as these methods were observed to be the best among the other evaluated approaches in our previous study. A total of 72 IEF fractions obtained from two biological replicates each were finally subjected to LC-MS/MS analysis, summing to a total of 288 runs including those of the technical replicates. MS/MS data analysis was performed using the Trans Proteomics Pipeline, and a minimum probability of 0.9 was set for confident peptide assignment. The mass error in parts per million (ppm) for precursor ions of all identified peptides is shown in Figure S1 in the Supporting Information. After assembling the proteins using ProteinProphet, we identified a total of 8363 different proteins including splice variants (at a false discovery rate < 1.2%), which, to our knowledge, is by far the most comprehensive proteome map of the zebrafish embryo (Tables S1 and S2 in the Supporting Information). This translates to an improvement of more than 2-fold in proteome coverage in comparison to that in our previous report.13 Of these proteins, 6475 proteins (78%) were detected by at least two peptides. About 10% of the proteins (857) were detected via the same single peptide multiple times, whereas about 12% (1031) were detected once by a single peptide (Figure 1B). The median number of peptides identified per protein was 4, and the corresponding tandem spectra detected per protein was 17. The median sequence coverage per protein was found to be 13.504.

The coverage of each protein attained by the corresponding peptide matches can be used to estimate the abundance of the identified proteins. To calculate the approximate abundance of the proteins in the zebrafish embryo, we used the exponentially modified protein abundance index (emPAI) algorithm, which normalizes the number of sequenced peptides per protein by the number of theoretically observable peptides of the protein.41 In order to conduct a comprehensive study of the zebrafish proteome, we explored the possibility of studying all detected proteins, including those that were detected in only one biological replicate. The emPAI-based semiquantitative protein abundance values showed a high correlation between the two biological replicates (Spearman’s correlation, 0.862) (Figure 1C). We, therefore, merged the data from both biological replicates for subsequent analysis (Table S3 in the Supporting Information).

The ranked distribution of all identified proteins allowed for evaluation of individual protein contribution to the total mass. It is revealed that 97.2% of the total protein abundance is contributed by the most abundant 25% of identified proteins (Figure 2A). Ninety percent of the total quantified proteome is within a range of log2 emPAI between 2.82 and −3.64 around the median abundance value. For further analysis, the quantified proteins were categorized into five quantiles based on their protein abundance values. Accordingly, the upper quantile constituted the very high and high categories, representing greater than the 90th percentile and 80–90th percentile, respectively, and the lower quantile comprised the very low and low categories, representing less than the 10th percentile and 10–20th quantile, respectively, of the estimated protein abundance values (Figure S2 in the Supporting Information). The quantile corresponding to 20–80% was considered to be moderate.

Assessing the physiochemical features across all of the identified proteins revealed that proteins of shorter length (<100 amino acids (aa)) are more abundant than proteins that are over 1500 aa in length (Figure S3A and Table S4 in the Supporting Information). On dissecting the length distribution within the individual clusters, we noticed that a majority of the highly abundant proteins (very high + high) are shorter than 450 aa (Figure S4A,B in the Supporting Information). The bias toward shorter length is also reported for highly abundant transcripts.60 This serves as an efficient means of minimizing energy cost, and the short proteins, generally in high abundance, play key roles in various cellular processes including signaling, cell-cell communication, and other basic metabolic processes. The pI distribution, on the other hand, showed a

Figure 2. Quantitative analysis of expressed proteins. (A) Ranked protein abundances from highest to lowest across the global quantified proteome. The contribution of each of the ranked quantiles to the total quantified embryonic protein abundance is indicated. (B) Enrichment based on GO categorization in each of the high- and low-abundance protein groups. Frequency corresponds to the preferential enrichment against the enrichment of total detected proteins. The GO categories are represented as MF (molecular function), BP (biological process), and CC (cellular component).
432 drop in abundance for proteins with midrange pI values (Figure S3B and Table S4 in the Supporting Information). However, comparison of the pI value distribution between the upper and lower quantile displayed similar trends across the pI range, with an under-representation of proteins with basic pI values (Figure S4C,D in the Supporting Information).

To map the chromosomal distribution of the quantified proteome, the identified proteins were traced back to their genomic loci by mapping against the annotated protein-coding genes (Table S3 in the Supporting Information). However, only about 78.3% of the detected proteins could be successfully mapped to each of the 25 chromosomes (Table S5 in the Supporting Information). Most of the identified proteins mapped to genomic regions in chromosome 5, and the least number of identified proteins were represented by chromosomes 4 and 24 (Figure S5 in the Supporting Information).

This is in agreement with the genome data that demonstrated relatively fewer protein-coding genes in chromosome 4.6 On comparing the coverage of the identified proteins to the total protein-coding genes annotated for each of the chromosomes, we observed a maximum coverage of ~31.5% for chromosome 19, closely followed by ~30.4% for chromosome 5 (Figure S6A in the Supporting Information). The least coverage (~14.7%) was observed for chromosome 4, suggesting that the genes present here are not protein coding or do not express at 24 hpf. Intriguingly, chromosome 4 is unique in the large number of noncoding RNAs and repeats that it harbors as well as the presence of a large family of genes that are specific to D. rerio.6

Of note, chromosomes 3, 6, 11, and 19 are particularly enriched in abundantly expressed (very high + high cluster) protein-coding genes (Figure S6B in the Supporting Information).

To obtain functional insights into the biological processes and cellular organization that are active at this developmental stage, we performed GO-slim analysis for the high- and low-abundance protein groups. Enrichment was performed by hypergeometric testing, and significant GO categories were identified in each group. The cluster frequencies of each of the significant GO categories were used to calculate the frequency of over-representation with respect to the overall quantified proteome, as shown in Figure 2B. We observed that proteins related to basic metabolic functions, primarily translation-related processes, are the most significantly enriched in the highly abundant protein cluster. Also, processes related to protein transport and organelles involved in trafficking, including the endoplasmic reticulum and nuclear envelope, are abundantly enriched. As observed in other systems, regulatory proteins associated with kinase activity, enzyme regulation, and protein binding have lower expression levels in the zebrafish embryo. At 24 hpf, the embryo is still in a very early stage of development, and important morphogenetic features including pigmentation, the cardiac tube, and fin fold begin to appear. Accordingly, we observed that the lower quantile proteins are enriched in functions relating to cell differentiation, structure morphogenesis, and embryonic development. While metabolic processes related to protein and carbohydrates are functionally enriched, lipid metabolic processes are low in abundance.
Concordance with Transcript Abundance

It is widely appreciated that the regulation of a protein’s level occurs at multiple levels beyond RNA transcription. To determine the extent of such regulatory systems, we sought to correlate our proteome data with the transcriptome of embryos at the same stage; hence, we performed RNA-seq on the 24 hpf embryos to determine transcript abundance (Table S6 in the Supporting Information). The RPKM measure obtained from the RNA-seq is a representation of transcript abundance. By including only those transcripts with an abundance greater than 2 and subsequently mapping the reads to the zebrafish reference genome (ZV9), a total of 101 unique transcripts were obtained. Some of the transcripts included alternative splice variants of the same genes. Thus, a total of 9601 different protein-coding genes were successfully identified. The different protein-coding genes showed a similar distribution among the 25 chromosomes as that from the quantified proteome. The maximum number of identified transcripts mapped to chromosome 5, similar to that observed for the proteome, and the least number of genes mapped to chromosome 24 (Figure S5 and Table S5 in the Supporting Information).

The quantified proteome and transcriptome data showed similar percentages across the different anatomical enrichment categories (Figure 3A). A majority of the transcripts and proteins (∼65% for transcripts and ∼75% for proteins), however, were not annotated to any specific anatomical feature and hence the distinct roles of these genes in the development of zebrafish remain to be explored. In fact, many of the known morphological developments that occur at 24 hpf, including the development of the retina, fin, and myotome, are represented with higher percentages of proteins, suggesting that the deyolking and extensive fractionation have enabled a thorough representation of the proteome.

On comparing the transcriptome and proteome data on the basis of gene annotations, we observed that some of the IPI (International Protein Index) identifiers mapped to more than one transcript (Table S7 in the Supporting Information). In all, we identified a corresponding transcript for a total of 5254 proteins in our quantified proteome. On comparing the distribution of abundances for the overall quantified transcriptome, we observed that no proteins were identified for a considerable number of transcripts in the lower abundance range (Figure S7AB in the Supporting Information). We also noticed that the distribution of protein abundance is broader than that of the corresponding transcript abundance values, although both of the abundance distributions share the same general shape (Figure 3B). Altogether, there was a 64% overlap between our proteome and transcriptome data on the basis of common protein-coding genes, excluding references of alternative slice variants (Figure 3C).

The RPKM and emPAI values are a proxy for the cellular abundance of transcripts and proteins, respectively, at a given point in time; hence, we analyzed the correlation between these two measurements. We observed a moderate correlation between the RPKM-based transcript abundance and emPAI-based protein abundance (Spearman correlation, 0.498) (Figure 3D). The level of correlation obtained is comparable to that observed previously in other organisms including human (Spearman correlation, 0.6), Drosophila (Spearman correlation, 0.66), Caenorhabditis elegans (the Spearman correlation, 0.59), and yeast (Spearman correlation, 0.58). Although proteins modulate key events within the cells, up- or downregulation of mRNA from large-scale transcriptomic studies is directly associated with protein expression levels based on the assumption that there is high correlation between transcript and protein abundances. Comparative studies performed in various organisms, however, suggest that the correlation coefficients generally range between 0.3 and 0.6, highlighting that protein levels are regulated beyond transcription.

We next analyzed the concordance between the transcript and protein abundance across functional categories that were arbitrarily grouped to represent core cellular and regulatory and developmental functions (Figure 4A,B). Translational process-related genes were found at the extreme end of the distribution with highest the correlation (Spearman correlation, 0.683), suggesting that these genes have elevated expression at both the transcript and protein levels (Figure S8 in the Supporting Information). The transcriptional machinery proteins, on the other hand, showed moderate abundance at the protein level but were more elevated on the transcript scale. Those belonging to carbohydrate metabolism, although generally considered to be abundant, spanned over almost the entire distribution with a moderate correlation. The lipid metabolic process-associated genes were frequently of low abundance, and this category was the least correlated. Focusing on the proteins important in development and signal transduction, we found that with the exception of a few proteins at the top end of the distribution that are associated with embryonic development, the rest had moderate-to-low protein expression levels (Figure 4B and Figure S9 in the Supporting Information). This suggests that processes relating to cell organization and development are modulated more at the mRNA level than at the protein level at this early stage of development.

Further investigating the cellular compartmentalization across the distribution, we noticed that the ribosomal proteins form one tight cluster at the top end of the distribution and a second additional cluster at moderate expression levels (Figure S10A,B in the Supporting Information). The nucleus, represented with the maximum number of proteins, extended over a large range traversing the entire range of abundance distribution. Although we noticed a slight bias for organelles like mitochondria and endoplasmic reticulum when based on

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Figure 4. Functional correlation of protein and mRNA levels. (A) Scatter plot of the mRNA and protein abundances across cellular core functions and (B) regulatory and developmental functions based on GO terms. Significantly enriched groups based on GO categorization are shown.
the quantified proteome, no such distributional bias was observed for the quantified transcriptome. 

**Distinct Functional Regulation of Proteins and Transcripts**

Embryonic development is associated with highly regulated processes that need to be precisely controlled at both the mRNA and protein levels. The analysis above provided a functional overview of the transcriptome and proteome across the entire distribution of cellular abundance. To delineate biological processes that are modulated at different levels of mRNA and protein abundances, we performed a combined hierarchical clustering of the observed transcriptome and proteome. We categorized genes based on their mRNA or protein level into various abundance groups (detailed below) and performed GO enrichment (biological process) by hypergeometric testing on all of the individual groups. Those GO categories that were significantly enriched in at least one of the seven groups are depicted on the heat map. The red arrow corresponds to high abundance of quantified proteins or transcripts, the green arrow, low, and the black symbol, moderate. The shaded red triangle corresponds to moderate-to-high protein or transcript abundances, and the shaded green triangle represents moderate-to-low protein or transcript abundances. High (yellow) and low (blue) in the heat map represent statistical over- or under-representation, respectively.

**Figure 5.** Functional modulation of protein and transcripts in the developing embryo. Proteins and mRNAs were grouped into seven groups based on abundance values as follows: high protein and mRNA, low protein and mRNA, moderate protein and mRNA, high protein and moderate-to-low mRNA, low protein and moderate-to-high mRNA, low mRNA and moderate-to-high, and high mRNA and moderate-to-low protein. The clustered GO biological process terms enriched in at least one of the seven groups are depicted on the heat map. The red arrow corresponds to high abundance of quantified proteins or transcripts, the green arrow, low, and the black symbol, moderate. The shaded red triangle corresponds to moderate-to-high protein or transcript abundances, and the shaded green triangle represents moderate-to-low protein or transcript abundances. High (yellow) and low (blue) in the heat map represent statistical over- or under-representation, respectively.
We first categorized the quantified 5254 genes (including splice variants) into four groups that represented only the upper and lower quantile as follows: high protein and high mRNA, high protein and low mRNA, low protein and high mRNA, and low protein and low mRNA. After performing hierarchical clustering on the four groups, we observed that genes associated with core cellular functions such as energy metabolism, protein transport, and cellular biosynthetic processes are fine-tuned by high levels of both mRNAs and their corresponding proteins (Figure S11 in the Supporting Information). The regulatory (post-translational modification) and developmental process (embryo morphogenesis, nervous system development) are modulated by genes with high mRNA expression and low protein levels. For the remaining two groups, very few GO categories passed the p value threshold, and these often coincided with enriched categories in the high protein and high mRNA and low protein and high mRNA groups.

To gain further functional insights on the entire range of the quantified proteome and transcriptome, we additionally categorized the quantified 5254 genes to include genes from the high-, low-, and moderate-abundance groups. For this purpose, we identified seven groups based on their abundances as follows: high protein and mRNA, low protein and mRNA, moderate protein and mRNA, high protein and moderate-to-low mRNA, low protein and moderate-to-high mRNA, low mRNA and moderate-to-high protein, and high mRNA and moderate-to-low protein (Figure 5). We observed that processes associated with core cellular functions such as metabolism, protein transport, and cellular biosynthetic processes are generally modulated by high protein levels, whereas the mRNA levels show considerable variation. The genes associated with nucleoside metabolic and biosynthetic processes display both high and low levels of mRNA and protein. Interestingly, genes associated with eye development have high mRNA and protein levels, with the optic system being in the mid-to-late phase of its development. Other cellular processes including folding and DNA replication are also modulated by high expression at both levels. Functions associated with DNA damage and repair, cell death, protein localization, and sensory organ development exhibit moderate abundance of expressed genes.

Genes that displayed a low protein level but moderate-to-high transcript abundance encoded for those functionally important proteins that are involved in regulatory (post-translational modifications), signal transduction, migratory, and developmental processes. We observed that genes associated with the development of the fin, cartilage, and embryonic skeletal system are highly represented within this group. We note that all of these tissues have not yet initiated their developmental programs and thus these may represent poised conditions. Other processes, such as those associated with the development of the central nervous system, and important morphogenetic events, such as pattern specification process, cell projection organization, and appendage development, are also characterized by differential levels of mRNA and protein.

Genes that displayed a high mRNA level but moderate-to-low protein level were associated with tissue and organ development and regulation of cellular processes. Specifically, processes associated with the regulation of primary metabolic and biosynthetic processes, gene expression, mRNA processing, and cell cycle display significantly high mRNA expression levels and low protein levels. Functions pertaining to microtubule-based movement, lipid metabolic processes, and stress response show enrichment in the low transcript and moderate-to-high protein level group, possibly indicative of long-lived proteins with sentinel roles.

Although proteins are the ultimate biological effectors, we observed that most of the embryonic morphogenetic events and signal transduction processes are modulated by low protein and high mRNA levels, in contrast to the core cellular functions that are modulated by high protein and transcript levels. To assess if any of these low-abundance proteins could be regulated by putative miRNAs, we carried out miRNA target prediction using TargetScanFish 6.2, which predicts gene targets based on conserved sites (7-mer or 8-mer) that match the seed region within a miRNA. It is predicted that many such low-abundance proteins are specifically regulated by miRNA families in the zebrafish embryo (Table S8 in the Supporting Information). This underscores that finer control of protein levels through miRNA regulation or high turnover is constantly in action to ensure proper and coordinated development.

Potential interactions among proteins expressed at high and low abundance may modulate important functional processes within the cell. To unveil the functional connectivity among the high- and low-abundance proteins, we constructed protein–protein interaction networks based on curated information from the reactome pathways and GeneGO MetaCore. The reactome functional interaction (reactome FI) combines protein–protein interactions from various organisms alongside curated pathway maps to provide with a high-quality pathway- informed interaction resource, and MetaCore is an expert-curated reliable data source for protein interactions, primarily focusing on human, rat, and mouse pathways. The zebrafish shares many orthologous proteins with those from the mammalian groups and hence these resources may be useful in deriving possible interactions among the quantified zebrafish proteins.

High-Abundance Protein Interactome. A large number of the proteins present in the high-abundance protein groups shared direct interactions with each other and resulted in a closely connected network (653 nodes and 6881 edges) (Figure S12A in the Supporting Information). While a majority of the corresponding transcripts also displayed high to very high abundance, we noticed that a few genes were modulated differently at the protein and mRNA levels and that the rest remained at moderate-abundance levels. Using MCODE, the tightly connected clusters were identified (Figure S12B in the Supporting Information). Of the 10 significant clusters identified with a minimum number of five nodes, the highest-ranking cluster (80 proteins) belonged to Translation. The other clusters were primarily associated with core cellular functions including transcription and RNA transport, oxidative phosphorylation, protein folding, DNA repair, and carbohydrate metabolism. The very low transcript abundance of one of the splicing factor proteins that forms the U2 small nuclear ribonucleoprotein complex (U2snRNP), SF3B3, leads us to speculate that this protein may be highly stable or efficiently translated. RPS6KA1 is the other important protein whose abundance is different at the protein and mRNA levels. The gene encodes for a serine/threonine kinase that is involved in...
Various signal transduction processes, including MAPK and the nutrient sensing mammalian target of rapamycin (mTOR)\(^5\). Poor correlations between the mRNA and protein abundances may point to more control at the translational or post-translational level. Assessing the degree of variation of the mRNA expression at different time points during development may provide us with clues on the influence of transcriptional or translational control for the poorly correlated genes.

**Low-Abundance Protein Interactome.** In contrast to the high number of proteins that could interact directly within the higher abundance groups, only 163 low-abundance proteins could be potentially mapped onto a direct interaction network (Figure 6). The clustering coefficient was low (0.156), and the average number of neighbors for each node was only around 3. Hence, dense clusters, as observed in the high-abundance protein network, could not be visualized. The number of noncorrelated genes in terms of protein and mRNA abundance...
is high in the mapped network. While only about 20% of the
genomes were expressed at low levels that were on par with the
protein abundance, 13% of the genes had high transcript
abundance, and the rest had moderate mRNA levels. Protein
interactions significantly enriched within the low-abundance
protein groups primarily correspond to cell signaling pathways.
WNT and Notch signaling, which are important modulators of
growth and cell fate decisions in the developing embryo, are
highly represented. The Slit-Robo pathway, which is involved in
axon guidance and heart tube formation, and the Rap1 signaling
pathway, which is associated with cell adhesion and junction
formation and is also crucial for heart development in zebrafish,
are also significantly enriched.\textsuperscript{55,56} The mapped signaling
pathways also include genes that are differentially regulated at
the transcriptional and translational levels. The observed
discordance in expression of several of these genes may
possibly point to important regulatory processes at the level of
signal transduction and embryonic development

\textbf{DISCUSSION}

In this study, we present a deep proteome map of a selected
stage of early embryonic development in zebrafish. We
identified >8000 proteins, which is by far the most extensively
measured zebrafish embryo proteome. In addition, we also
performed transcriptome mining by RNA-seq to provide an
informative comparative map of the quantified proteome and
transcriptome during embryonic development. Such an
integrative approach links the transcript abundance to protein
levels and ultimately highlights specific developmental
processes that are modulated by changes in the mRNA or
protein species. Although similar studies have been performed
previously in other organisms,\textsuperscript{16,17,57,58} we unveil, for the first
time, large-scale combinatorial functional mapping in zebrafish.
Previous transcriptomics and proteomic studies are limited
by overall coverage.\textsuperscript{52,56} It may be possible that some of the
proteins are masked by high-abundance proteins, restricting
detection limits, or that the peptides are not amenable to
detection by mass spectrometry. Also, regardless of the
transcript levels, some of the mRNAs may not be efficiently
translated, resulting in low or undetectable protein levels. We
noticed that for a majority of the genes expressed at low levels
the corresponding proteins could not be identified. Never-
theless, we could successfully map \textasciitilde64\% of our quantified
proteome to a corresponding mRNA level. Our proteome data
identified a paucity of coding sequences on chromosome 4,
confirming the analysis of the genome.\textsuperscript{6} Moreover, our analysis
revealed that there was only a moderate correlation between
the protein and transcript levels for most of the cellular
processes, consistent with observations in other biological
systems.\textsuperscript{5,17,58} Although such low correlations may be
attributed to technical discrepancies, it is also suggestive of an
intricate functional regulatory mechanism that operates to
maintain proper levels of transcripts and proteins. There is
continuing debate on the concordance of transcript and protein
abundances, and the precise mechanisms that act at the post-
transcriptional level remain to be elucidated.\textsuperscript{57,59,60} In fact,
cross-species comparisons suggest that orthologous protein
levels correlate better than the corresponding transcript
abundances, hinting that the mechanisms to achieve a particular
protein abundance evolve rapidly\textsuperscript{16} and may include utilization
of altered protein stability, translational efficiency, and
ribosomal occupancy to achieve the final protein abun-
dance.\textsuperscript{61--63} Interestingly, a genome-scale study established
that genes that showed minimum variation at the mRNA level
through the cell cycle had poor correlation with final protein
levels, whereas those that displayed large variation had a high
degree of concordance.\textsuperscript{64} Analyzing the concerted variation in
transcript and protein abundances through the different
developmental stages may expose more insights on the
orchestration of the transcriptional and translational machi-
neries in zebrafish.

As observed previously in other biological systems, we find
that the most abundant proteins are of considerably shorter
length and are often associated with central pathways and core
processes, including energy and carbohydrate metabolism,\textsuperscript{51,65}
translation, and transport.\textsuperscript{51,65} For such core metabolic
processes, the correlation between the protein and mRNA
levels was considerably higher. The genes associated with these
processes indeed exhibit a highly conserved coexpression
pattern and are also highly correlated at the protein level
across species.\textsuperscript{16,66} The low-abundance protein groups are
primarily regulatory in function and are involved in signal
transduction, phosphorylation, and other protein modifications.

In spite of the low abundance, higher eukaryotes have a large
fraction of the protein mass dedicated to regulatory functions.\textsuperscript{67}
Furthermore, low-abundance species generally show high
sequence variability across species.\textsuperscript{16} These proteins, although
present in low abundance, are associated with various important
processes of development, including cell differentiation and
anatomical structure morphogenesis, in our quantified
proteome. Of note, it has been shown that regulatory proteins
display varied expression levels between different human cell
lines.\textsuperscript{58} This supports the notion that these regulatory proteins
are potent regulators of cell identity and behavior.

While we observed that most of the core metabolic and
biosynthetic processes are modulated by proteins that are
highly abundant, specific developmental processes are marked
by genes showcasing high transcript but low protein levels. It is
known that some transcripts are not efficiently translated, are
differentially degraded, and/or are stalled during the process of
translation. Those cohorts of transcripts, which showed
abundant transcripts and low protein levels, might be subjected
to the process of stalling during translation. Ribosome profiling
experiments identified the existence of differential translational
efficiencies of these transcripts.\textsuperscript{69,70}

Anatomical developments associated with morphogenesis of
embryonic skeletal systems, pattern formation, and neuronal
differentiation, for example, are fine-tuned by low protein and
high transcript levels. Such systems are yet to initiate full
developmental programs at 24 hpf, and we speculate that they
may be held in a poised state. As the embryo develops, some of
these processes may be primarily modulated by the levels of
proteins, transcripts, or both. Network mapping highlighted
instances of several genes involved in signaling pathways that
had low protein and high mRNA expressions. The low protein
abundance may be attributed to reduced stability of the
regulatory proteins or may point to post-transcriptional
regulation of protein abundance. miRNAs have emerged as
important modulators of post-transcriptional regulation, and it
is estimated that approximately 30% of the mammalian coding
genes are regulated by them.\textsuperscript{71} Indeed, most miRNAs in
zebrafish are primarily expressed from segmentation stage
onward, and some miRNAs have been shown to regulate
different processes during development.\textsuperscript{72--75} This leads us to
speculate that many of the low-abundance (or even
undetected) proteins may have been subjected to miRNA
regulation.
regulation to allow for the tightly regulated initiation of
differentiation. Of note, we observed that many of the critical
proteins involved in development, including LFNG,76
FGFRs,77 and CTNNB2,78 are predicted to be regulated by
miRNAs (Table S8 in the Supporting Information). Such
restriction of mRNA translation underscores the potency of the
derived protein products.

From our network mapping, we identified a particular
modulator of the Notch pathway, Lunatic Fringe (LFNG),
which has low protein and high mRNA levels. The Notch
signaling cascade plays a major role in the establishment of the
neural crest and binary fate decisions in the neural tube and
elsewhere, and it also regulates somitogenesis in developing
embryos.79–81 Formation of the somites (embryonic segments
of the vertebrate body) is regulated by oscillatory expression of
genes in the segmentation clock that define the spatial pattern,
and Notch functions to synchronize the segmentation clock.82
Such synchronization depends on tight control at the level of
mRNA half-life and translational efficiency. Interestingly,
LFNG is one such oscillatory gene whose expression is post-
transcriptionally regulated by miR-125a-5p (miR-125 was also
predicted by TargetScanFish 6.2) for proper somite formation
in chick embryo.83,84 In zebrafish, LFNG is expressed within
the presomitic mesoderm and is important for the formation of
gene regulatory processes may have resulted in
the observed differences in the gene and protein expression
levels of LFNG. Similar dissection of the network for other
protein modules, like FGFR, CLASP2, and DUSP6, may
provide functional insights into the differential regulation of the
transcriptional and translational machineries during zebrafish
embryogenesis.

CONCLUSIONS

The in-depth comparative and functional mapping presented
here highlights the usefulness of integrative proteomics and
transcriptomics to unveil molecular mechanisms regulating
early embryogenesis in zebrafish. We particularly highlight
differential modulation of various morphogenetic events during
embryogenesis. We believe that such an exhaustive approach
over the entire time course of development is likely to uncover
many novel mechanisms and various levels of gene expression
control during embryogenesis and provide a valuable resource
for systems biology-based modeling in the future.

ASSOCIATED CONTENT

Supporting Information

Table S1: List of all identified peptides and proteins from the
zebrafish embryo proteome. The data was retrieved using
Trans-Proteomic Pipeline analysis. Table S2: List of identified
proteins from the zebrafish embryo proteome. This table enlists
the spectral matches, sequence coverage, and peptide counts for
each protein. Table S3: Quantification of detected proteins.
The table enlists the calculated abundances as emPAI values
and information on chromosomal mapping for each protein.
Table S4: Assessment of length and pI for all quantified
proteins. Table S5: Chromosomal distribution of detected
proteins and transcripts. This table enlists the total number of
genes annotated for each chromosome and those that could be
distinctly mapped from the quantified proteome and tran-
scriptome to individual chromosomes. Table S6: Deep
sequencing of transcripts from the zebrafish early embryo.
Table S7: Mapping of quantified proteins with corresponding
transcripts. This table enlists the 5254 proteins, including
instances of splice variants, for which a corresponding transcript
was identified using deep sequencing of the zebrafish embryo.
Table S8: miRNA prediction for low-abundance proteins. This
table enlists the sites and the total context score for each
predicted miRNA. Figure S1: Mass error in parts per million
(ppm) of precursor ions of all identified peptides. Figure S2:
Density distribution of protein abundances as represented by
emPAI. Figure S3: Bias analysis of protein length and pI. Figure
S4: Distribution of length and pI in the high- and low-
abundance protein clusters. Figure S5: Distribution of mapped
genes from the proteome and transcriptome across different
chromosomes. Figure S6: Coverage and enrichment of mapped
genes from the proteome across different chromosomes. Figure S7:
Density distribution of transcript abundances as repre-
sented by RPKM values. Figure S8: Correlation of tran-
scriptome and proteome for core cellular processes. Figure S9:
Correlation of transcriptome and proteome for regulatory and
developmental processes. Figure S10: Correlation of protein
and mRNA levels for different subcellular compartments. Figure
S11: Functional modulation of proteins and transcripts in
the early stage embryo. Figure S12: Protein–protein
interaction network among the high-abundance proteins. This
material is available free of charge via the Internet at http://
www.ncbi.nlm.nih.gov. The protein data set information was uploaded to
the PeptideAtlas database and is available at ftp://
ftp.peptideatlas.org/.

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Notes

The authors declare no competing financial interest.

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ABBREVIATIONS

AA, amino acids; BP, biological process; CC, cellular
component; DAVID, Database for Annotation, Visualization
and Integrated Discovery; emPAI, exponentially modified
protein abundance index; ESI, electrospray ionization; FDR,
false discovery rate; GO, gene ontology; hpf, hours post
fertilization; HPLC, high-performance liquid chromatography;
IEF, isoelectric focusing; IPL, international protein index; LC--
MS/MS, liquid chromatography–tandem mass spectrometry;
LFNG, lunatic fringe; LTD, linear trap quadrupole; MF,
molecular function; miRNA, microRNA; mRNA, messenger
RNA; mTOR, mammalian target of rapamycin; PPM, parts per
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