Characterisation of the Fibroblast Growth Factor Dependent Transcriptome in Early Development

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Abstract

Background: FGF signaling has multiple roles in regulating processes in animal development, including the specification and patterning of the mesoderm. In addition, FGF signaling supports self renewal of human embryonic stem cells and is required for differentiation of murine embryonic stem cells into a number of lineages.

Methodology/Principal Findings: Given the importance of FGF signaling in regulating development and stem cell behaviour, we aimed to identify the transcriptional targets of FGF signalling during early development in the vertebrate model Xenopus laevis. We analysed the effects on gene expression in embryos in which FGF signaling was inhibited by dominant negative FGF receptors. 67 genes positively regulated by FGF signaling and 16 genes negatively regulated by FGF signaling were identified. FGF target genes are expressed in distinct waves during the late blastula to early gastrula phase. Many of these genes are expressed in the early mesoderm and dorsal ectoderm. A widespread requirement for FGF in regulating genes expressed in the Spemann organizer is revealed. The FGF targets MKP1 and DUSP5 are shown to be negative regulators of FGF signaling in early Xenopus tissues. FoxD3 and Lin28, which are involved in regulating pluripotency in ES cells are shown to be down regulated when FGF signaling is blocked.

Conclusions: We have undertaken a detailed analysis of FGF target genes which has generated a robust, well validated data set. We have found a widespread role for FGF signaling in regulating the expression of genes mediating the function of the Spemann organizer. In addition, we have found that the FGF targets MKP1 and DUSP5 are likely to contribute to the complex feedback loops involved in modulating responses to FGF signaling. We also find a link between FGF signaling and the expression of known regulators of pluripotency.

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Introduction

Fibroblast growth factors (FGFs) are small polypeptides that have multiple functions in early development and homeostasis of the adult organism. FGFs are present in all animal groups and are one of relatively few families of extracellular signaling molecules that are involved in regulating animal development. 22 FGFs have been identified in higher vertebrates [1].

FGF signaling has a key role in specifying the primary germ layers that give rise to all the tissues of the adult organism. Experiments initially carried out in amphibians, and later supported by studies in mammals, birds and fish, demonstrated that FGF signaling is required to regulate gene expression within the early vertebrate mesoderm, which is the germ layer giving rise to muscle, skeleton, connective tissue, blood and organs such as the kidney [2–6]. As well as regulating mesodermal gene expression, FGF signaling is involved in regulating the complex morphogenetic activity exhibited by mesoderm cells during vertebrate gastrulation [3,7]. FGF signals produced by the mesoderm, acting on the adjacent ectoderm, are also required for induction and patterning of the vertebrate nervous system [8–11].

More recently it has been shown that FGF signaling plays a critical role in the commitment of mouse embryonic stem (ES) cells to mesodermal, as well as both neural and non-neural ectodermal lineages [12,13]. FGF signaling is also important for maintaining self renewal in human ES and induced pluripotent stem (iPS) cells in culture [14–16].

Given the importance of FGF signaling in adult and embryonic life, the downstream transcriptional targets involved in mediating the activities of the FGFs are of great interest. In order to identify genes that respond to FGF signaling in early development we have compared gene expression in normal embryos with embryos in which FGF signaling has been inhibited. Our analysis identifies 67 genes which are significantly down regulated and 16 genes which are up regulated in response to FGF inhibition. A high proportion of the putative FGF target genes have predicted functions associated with cell signaling and transcriptional regulation.

We show that many of the targets are expressed in known regions of FGF activity during development. Our analysis reveals some interesting features of the FGF dependent transcriptome. We find that FGF signaling is required for the normal expression of multiple genes in the Spemann organizer, a structure orthologous...
to the node of higher vertebrates and which is required for the establishment of the main body axis.

Interestingly, we find that inhibition of FGF signaling down regulates expression of *Lin28* and *FoxD3*, two genes which have been implicated in regulating the pluripotential state of embryonic stem (ES) cells [14,17–19].

Cluster analysis, based upon temporal expression, identified a number of distinct waves of expression from FGF targets following the initial activation of FGF signaling in the amphibian embryo. Following the activation of endogenous FGF signaling in blastula stages, we show that two of the earliest expressed target genes are the MAP kinase phosphatase 1 (*MKP1*) gene and a novel *Xenopus* gene related to human Dual Specificity Phosphatase 5 (*DUSP5*). We show that both *DUSP5* and *MKP1* inhibit FGF dependent ERK/MAP kinase phosphorylation and mesoderm formation induced by FGF in *Xenopus* tissues. Our analysis indicates that *DUSP5* and *MKP1* are members of the FGF synexpression group and are components of the negative feedback network required to limit the extent of FGF signaling in the early embryo.

Results

Timing of FGF signaling in the *Xenopus* embryo

The aim of this study was to identify FGF targets that are induced shortly after the initial activation of zygotic FGF signaling in the embryo. It was therefore necessary to accurately determine when FGF signaling is activated in the embryo. We examined the temporal profile and spatial distribution of activated diphospho-ERK (*dp-ERK*), which is a key signal transduction effector of FGF signaling in the *Xenopus* embryo [20–22]. Figure 1A is a Western blot showing levels of *dp-ERK* in embryos from early cleavage to late blastula stages (NF stage 3 to 9.5). We detect constant low levels of activated ERK up to stage 8.5. The initial rise in the level of *dp-ERK* is detected at mid-blastula stage 8.5 and there is robust increase by stage 9 (7 hours post-fertilization (pf) at 23°C), corresponding to the onset of major zygotic transcription at the mid-blastula transition (MBT) [23].

Our data show that the initial activation of ERK is in a dorsal to ventral gradient within a broad belt of tissue around the equator of the embryo at late blastula stage 9 (Figure 1B). Initial *dp-ERK* activation is not limited to the presumptive mesoderm of the marginal zone but extends a considerable distance into the animal hemisphere on the dorsal side of the embryo. Figures 1C and D show that this early zygotic ERK activity is blocked by over expression of a dominant negative FGF receptor (dnFGFR). The dnFGFRs used in this study are carboxy-terminal truncations of the receptors lacking tyrosine kinase activity and block FGF signaling by associating with endogenous receptors to form non-functional dimers [24,25]. We conclude that zygotic activation of the FGF signaling pathway commences at mid-blastula stage 8.5 (6 hours pf at 23°C).

Timing of transcriptional responses to FGF signaling in the *Xenopus* embryo

The genes coding for the brachyury, *MycD* and *CdX4* transcription factors have previously been identified as targets of the FGF signaling pathway. These genes are activated by FGF even in the presence of the translation inhibitor cycloheximide and are defined as immediate early responses to FGF signaling [26–28]. We have used expression of these genes to indicate when the initial transcriptional responses to FGF signaling occur in the *Xenopus* embryo. Figure 1E is an RNAase protection analysis (RPA) showing that *Brachury* expression is detected by early gastrula stage 10 and *CdX4* by stage 10.25. However, robust expression of all three immediate early FGF response genes is not detected until stage 10.5 (11 hours pf at 23°C). Furthermore, we show that the initial expression of all three genes is blocked by over expression of a dnFGFR. Based on the timing of ERK activation and transcriptional activation of known FGF target genes, early gastrula stage 10.5 (11 hours of culture at 23°C) was chosen as the stage for the analysis of FGF targets.

Identifying transcriptional responses to FGF signaling

In order to identify transcriptional targets of FGF signaling gene expression in control embryos was compared to sibling embryos in which FGF signaling was inhibited by over expression of dominant negative versions of FGFR1 (dnFGFR1) or FGFR4 (dnFGFR4) [24,25].

In order to undertake statistical analysis of the microarray data three biological replicates were carried out. Each replicate set consisted of control embryos and embryos injected with dnFGFR1 or dnFGFR4 collected at stage 10.5 (11 hours pf at 23°C). Before proceeding to microarray analysis each replicate set was checked for effective FGF inhibition. Sibling embryos from each replicate set were analysed for *dp-ERK* levels and expression of *CdX4*, *Brachury* and *MycD* by both RPA and *in situ* hybridization. Figures 1F–1H are analyses of a representative biological replicate set showing that, with the experimental conditions used, overexpression of dnFGFR1 or dnFGFR4 results in potent inhibition of ERK activation (Figure 1F) and down regulation of transcription from known FGF target genes relative to sibling controls (Figures 1G and 1H).

Changes in gene expression resulting from dnFGFR1 and dnFGFR4 overexpression

After these quality control checks, the RNA samples from each of the three biological replicates were analysed using Affymetrix GeneChip *Xenopus laevis* Genome Arrays, which allow the analysis of more than 14,400 transcripts expressed in early development. Figures 2A and 2B are scatterplots of log2 gene expression values in controls versus dnFGFR1 and dnFGFR4 injected embryos. Probe sets showing greater than 2-fold changes in expression in control versus experimental groups are indicated by red and green points. These data show that the expression of a considerable number of genes is altered in response to FGF inhibition by both dnFGFR1 and dnFGFR4.

In contrast to Figure 2A and 2B, in dnFGFR1 versus dnFGFR4 injected embryos we see that expression levels of relatively few probe sets are greater than two-fold different between the two groups (Figure 2C). Our analysis shows that only four genes, using the criteria of ≥2-fold change in expression and a significance level of *p*≤0.01, exhibit different expression levels in dnFGFR1 versus dnFGFR4 injected embryos (Table S1). This indicates that at this stage of development the genes affected by inhibition with the different dominant negative receptors are largely the same. The differences in gene expression profiles of dnFGFR1 and dnFGFR4 injected embryos are quantitative differences in the levels of expression from the same set of target genes. This conclusion is supported by Figure 2D, which shows that both dnFGFR1 and dnFGFR4 down regulate the expression of several known FGF targets. However, in all cases dnFGFR4 has a more potent effect on gene expression. The data in Figure 2A and 2B follow a similar trend, in which there are greater fold changes in gene expression following dnFGF4 injection than with dnFGFR1. We conclude that inhibition with dnFGFR1 and dnFGFR4 affects the same sets of genes but that on a per mass of injected mRNA basis, dnFGFR4 is more potent.
Classification of putative FGF target genes

Lists of genes affected by FGF inhibition were compiled by comparing gene expression changes in dnFGFR4 injected embryos versus control embryos. The criteria of at least a 2-fold change in expression and a significance level of p < 0.01 were used to compile the gene lists. After the elimination of multiple probe sets representing the same gene, using the stated criteria, we find that 67 genes are significantly down-regulated by FGF inhibition, indicating that in normal development these genes are positively regulated by FGF signaling. Table 1 shows these genes in order of mean fold inhibition in dnFGFR injected embryos relative to controls. The T-box gene brachyury shows the highest fold inhibition (>19-fold). We find that only 16 genes are significantly up-regulated by FGF inhibition, indicating that in normal development these genes are negatively regulated by FGF signaling (Table 2).

Where possible we have classified FGF target genes based on cellular function. We find that a large proportion of genes positively regulated by FGF signaling are either involved in transcriptional regulation (24%) or cell signaling (18%), with a
A. Control vs dnFGFR1
B. Control vs dnFGFR4
C. dnFGFR1 vs dnFGFR4

D. Inhibition of known FGF target genes

E. Genes positively regulated by FGF signalling

F. Genes positively regulated by FGF signalling
   - Esr6
   - purine phosphorylase
   - glycogen phosphorylase
   - ephrin receptor A4
   - apobec2
   - unknown xls.5479

G. Genes negatively regulated by FGF signalling
   - Wig-related
   - glucocorticoid inducible leucine zipper
Derived expression values are based on the average of three biological replicates. Relative expression is calculated as a percentage of the expression in control embryos. Standard deviation bars are indicated. (D) shows pie charts of genes positively and negatively regulated by FGF signaling. Percentages of each group classified according to their putative cellular function are indicated. Details of the up regulated and down regulated genes are contained Tables 1, 2 and Tables S2 to S11. (E) shows the expression patterns of genes positively regulated by FGF signaling at determined by in situ hybridization at early gastrula stage 10.5, early neurula stage 14, post-neurula stage 22 and early tailbud stage 30. Gastrula embryos are vegetal views with dorsal to the top. Neurula embryos are dorsal views with anterior to the left. Post-neurula and tailbud embryos are lateral view with dorsal to the top and anterior to the left. (G) shows the expression patterns of genes negatively regulated by FGF signaling. doi:10.1371/journal.pone.0004951.g002

A relatively small number of genes (9%) involved in various aspects of metabolism. The corresponding figures for genes negatively regulated by FGF signaling are 29% involved in transcriptional regulation, 6% in cell signaling and 29% in metabolism. These data are represented as pie charts in Figure 2E and the detailed breakdown of FGF target classification, together with relevant references are presented in Tables S2, S3, S4, S5, S6, S7, S8, S9, S10 and S11. Tables S12 and S13 show the Gene Ontology (GO) term classifications for FGF targets derived from the available Affymetrix annotation files.

Expression of FGF target genes

Consistent with the pattern of FGF activity in the late blastula and early gastrula stage embryo (Figure 1) many of the genes identified as being down regulated following FGF inhibition are expressed in the mesoderm or dorsal ectoderm at the start of gastrulation. Expression data for previously charcterised genes, along with new data from this study are summarized in Table S14.

We have undertaken a more detailed expression analysis of a number of these positively regulated FGF targets at early gastrula, early neurula, post-neurula and early tailbud stages. (Figure 2F) We find that all of these genes are expressed in the mesoderm around the mesoderm. In post-gastrula stages the expression patterns of these genes diversify; however, some common patterns are detected. For example, the posterior mesoderm, the paraxial mesoderm and the tailbud are common sites of expression. Other sites of expression include the anterior CNS and branchial arches. In the case of a putative methyltransferase the post-gastrula expression is remarkably restricted, being limited to a very tight domain around the closed blastopore and later in the developing otic vesicle.

The expression patterns were also determined for two of the genes that are up regulated in response to FGF inhibition, which we predict will be negatively regulated by FGF signaling in normal development (Figure 2G).

Wig-related codes for a protein similar to Xwig1, which is a putative endoplasmic reticulum resident protein [29]. GILZ (glucocorticoid inducible leucine zipper), is a member of the Tsc-22 family of transcription factors related to Drosophila bunched [30]. In contrast with genes positively regulated by FGF, the early expression of these genes is excluded from the circum-blastoporal region. The Wig-related gastrula and post-gastrula expression pattern is highly dynamic, before resolving to a stable pattern of expression in the CNS, branchial arches and lateral mesoderm. After gastrulation GILZ is expressed in the neurogenic region of the open neural plate and subsequently in the neural tube.

Validation of FGF targets

It is generally accepted that gene lists identified by microarray analysis should be validated by independent methodology. For a number of the putative target genes independent validation of their response to FGF signaling was undertaken by in situ hybridization. Figure 3A shows the effects of FGF inhibition on the spatial expression of genes identified as being down regulated in the microarray-based screen. Injection of dnFGFR leads to dramatic inhibition of the circum-blastoporal expression of these genes in gastrula stage embryos.

Conversely, the size of the expression domains of two putative targets negatively regulated by FGF signaling are dramatically increased in embryos injected with dnFGFR mRNA (Figure 3B). In the case of CP2-like and GILZ, inhibition of FGF signaling leads to elevated expression in the circum-blastal region indicating that in normal development FGF signaling is required to exclude their expression from this region of the embryo.

Further validation of the FGF target genes was undertaken by showing that FGF signaling is not only necessary but is also sufficient for their expression. Figure 3C shows an RNase protection assay (RPA) on control animal hemisphere tissue explants (animal caps) and animal caps treated with recombinant FGF4 protein. With all genes tested, FGF treatment leads to marked increase in transcription as compared to control explants. Taken together, our analyses indicate that the microarray based screen has provided a well supported list of candidate FGF target genes.

FGF signaling and dorsal gene expression

The identified FGF target genes included several genes, including chordin and noggin, which are required for the function of the Spemann organizer in the dorsal mesoderm [31,32]. This region of the embryo plays a key role in regulating the formation of the main body axes. We investigated if there is a general role for FGF in regulation of dorsal gene expression. The data in Data in Figure 4A and Table 3 show that many dorsally biased genes, including Egr1 and FoxD5 are highly down regulated when FGF signaling is inhibited (>12-fold and >10-fold respectively). Many of the dorsally expressed FGF target genes have been shown to be directly involved in mediating organizer function, including Frzb1, chordin, noggin, FoxD3, FoxD5 and goosecoid. Other dorsally expressed genes, such as Xnr3 and cerberus, are not significantly down regulated, and others such as Otx2 show small increases in expression (not significant at the p = 0.01 level).

The in situ hybridizations in Figure 4B show that FGF inhibition dramatically reduces the spatial extend of chordin, FoxD5 and Frzb expression through early gastrula stages. We also show by RPA that chordin and noggin are strongly down regulated in early gastrula stage embryos following FGF inhibition (Figure 4C), indicating a role for FGF in regulating the expression of secreted BMP inhibitors.

Stimulation of BMP signaling leads to the phosphorylation and activation of SMAD1. Figure 4D is a Western blot for phospho-
Table 1. Genes positively regulated by FGF signaling.

| Gene                                | Fold inhibition | GenBank accession | Affymetrix probe set |
|--------------------------------------|-----------------|-------------------|----------------------|
| Brachyury                           | 19.2            | M77243            | XI.514.1.S1_at       |
| Egr1                                 | 12.4            | AF250345          | XI.637.1.A1_at       |
| FoxD5A                               | 10.8            | AF162782          | XI.642.1.S1_at       |
| SIP1                                 | 9.8             | AB038353          | XI.958.1.S2_at       |
| Cdx4                                 | 8.6             | UO2034            | XI.10269.1.S1_at     |
| Esr5                                 | 8.5             | BJ057112          | XI.14524.1.S1_at     |
| Purine phosphorylase                 | 7.9             | BM172525          | XI.16206.1.A1_at     |
| Marginal coil                        | 7.7             | BJ044312          | XI.544.1.S1_at       |
| Paraxial protocadherin               | 7.3             | AW782445          | XI.617.1.A1_at       |
| Glycogen phosphorylase              | 7.0             | BJ056085          | XI.781.1.A1_at       |
| NADH dehydrogenase sub-unit          | 6.5             | BJ051675          | XI.12993.1.A1_at     |
| FoxO3A                               | 6.0             | AB014611          | XI.525.1.S1_at       |
| G-coupled receptor P2YS              | 5.7             | BQ401062          | XI.19933.1.S1_at     |
| Related to DC-STAMP domain receptor  | 5.6             | BI447679          | XI.15270.1.A1_at     |
| Meso05                               | 5.1             | BF615090          | XI.7720.1.A1_at      |
| Uncharacterised protein C2orf32      | 5.1             | CB756627          | XI.25136.1.A1_at     |
| Frzb1                                | 5.1             | U78598            | XI.212.2.S1_at       |
| XPO                                 | 5.0             | BJ051206          | XI.5908.1.S1_at      |
| Ephrin receptor A4                   | 4.8             | BJ080037          | XI.132.1.A1_at       |
| XSp2                                 | 4.5             | BJ049843          | XI.2755.1.S1_at      |
| Zic3a                                | 4.5             | AB005292          | XI.7969.1.S1_at      |
| Xro3                                 | 4.4             | AF027175          | XI.4522.1.S1_at      |
| Gravin-like                          | 4.4             | AF308810          | XI.3468.1.S1_at      |
| Alkaline phosphatase                 | 4.3             | BC043760          | XI.1299.1.S1_at      |
| Apobec2                              | 4.2             | AW766385          | XI.5876.1.A1_at      |
| p75-like fullback receptor           | 4.2             | AF131890          | XI.3540.1.S1_at      |
| Wnt8                                 | 4.0             | X57234            | XI.49.1.S1_at        |
| Fructokinase-related protein         | 3.9             | CB756273          | XI.15623.1.A1_at     |
| Crescent                             | 3.9             | AF260729          | XI.619.1.S1_at       |
| Pinhead                              | 3.8             | BJ056268          | XI.3529.1.A1_at      |
| Wnt5b                                | 3.6             | AW148258          | XI.11619.1.S1_at     |
| Unknown                              | 3.6             | BJ092401          | XI.5479.1.A1_at      |
| Retrotransposon protein 1a11         | 3.6             | L11263            | XI.3352.1.S1_at      |
| FoxA4                                | 3.4             | S93559            | XI.1082.1.S1_at      |
| Mitotic phosphoprotein 67            | 3.2             | BJ077239          | XI.20772.1.A1_at     |
| Cdx1                                 | 3.2             | CB564190          | XI.23739.1.A1_at     |
| Sprouty2                             | 3.1             | AF331825          | XI.11965.1.S1_at     |
| DUSP5                                | 3.0             | BJ077463          | XI.15374.1.A1_at     |
| Chordin                              | 2.8             | BF610870          | XI.3549.1.S1_at      |
| MKP1                                 | 2.7             | AJ320159          | XI.2803.1.S1_at      |
| Unknown                              | 2.7             | BI32705           | XI.18179.1.S1_at     |
| Xor                                  | 2.7             | X98454            | XI.37.1.S1_at        |
| Putative nucleolar GTP binding protein | 2.7           | BM179370          | XI.14776.1.A1_at     |
| Lin28a homologue                     | 2.7             | BJ047699          | XI.34181.1.A1_at     |
| Glut1 transporter                     | 2.7             | BJ049047          | XI.24121.1.A1_at     |
| Unknown                              | 2.6             | BJ056692          | XI.15382.1.A1_at     |
| Dkk1                                 | 2.6             | AF030434          | XI.251.1.S1_at       |
| Unknown                              | 2.5             | AW460550          | XI.11594.1.A1_at     |
| RALDH2                               | 2.5             | B449483           | XI.18999.1.A1_at     |
| Prickle                              | 2.5             | AF387815          | XI.7556.1.S1_at      |
SMAD1/5/8 (p-SMAD). In keeping with a role for FGF signaling in regulating dorsally expressed secreted BMP inhibitors, such chordin and noggin, levels of p-SMAD1/5/8 are elevated in response to FGF inhibition. Our data indicate a widespread but not ubiquitous requirement for FGF signaling in the regulation of organizer gene expression during gastrula stages.

Expression profiling and cluster analysis of FGF target genes

In order to generate temporal expression profiles for individual FGF target genes from pre-MBT stages until early neurula stages we carried out Affymetrix microarray analysis on normally developing sibling embryos at hourly time points from 5 hours pf (stage 8) to 16 hours pf (stage 14) at 23°C. Figure 5A shows the relative expression profiles of FGF8 and 5 known FGF target genes. The initial rise in FGF8 expression is first detected at stage 9 (7 hours pf), indicating that FGF8 expression is activated very rapidly post-MBT. As mentioned earlier, this increase in FGF8 expression corresponds closely with the detected rapid elevation of dp-ERK levels in the embryo between stage 8.5 and stage 9 (Figure 1A). We note that although normal expression of these genes requires FGF signaling, the timing of gene expression is likely to have an impact on the outcome of the signaling pathway.

Table 1. cont.

| Gene                  | Fold inhibition | GenBank accession | Affymetrix probe set |
|-----------------------|-----------------|-------------------|----------------------|
| ADMP                  | 2.4             | BF231842          | XL.3809.1.A1_at      |
| Unknown               | 2.3             | BJ085271          | XL.1521.1.A1_at      |
| Cytochrome B561       | 2.3             | U16364            | XL.11917.1.S1_at     |
| Goosecoid             | 2.3             | BJ056432          | XL.801.1.S1_at       |
| FoxC1                 | 2.3             | AF116844          | XL.180.1.S1_at       |
| Noggin                | 2.2             | M96807            | XL.834.1.S1_at       |
| Sprouty1              | 2.2             | BG022481          | XL.10087.1.A1_Fat    |
| Oct1                  | 2.2             | BG022051          | XL.1265.1.S1_at      |
| Rexp52                | 2.1             | BG555868          | XL.3023.1.A1_at      |
| Grb10 interacting protein2 | 2.1            | BJ098841          | XL.14208.1.A1_at     |
| Putative methyltransferase | 2.1          | BJ100128          | XL.20056.1.S1_a_at   |
| Connexin 29           | 2.1             | BJ076720          | XL.8924.1.A1_at      |
| SMCT                  | 2.1             | BJ047668          | XL.6392.1.A1_at      |
| Weakly similar to Rab1| 2.1             | BJ079872          | XL.3365.1.A1_at      |
| Unknown               | 2.1             | CA972457          | XL.19961.1.S1_at     |
| Moderately similar to Brain protein 44 | 2.1  | BJ088835          | XL.15887.1.S1_x_at   |
| Ephrin receptor A2    | 2.0             | BF025525          | XL.14496.1.A1_at     |

Table 2. Genes negatively regulated by FGF signaling.

| Gene                  | Fold activation | GenBank accession | Affymetrix probe set |
|-----------------------|-----------------|-------------------|----------------------|
| XIRG protein          | 13.5            | AJ278067          | XL.4965.1.S1_at      |
| PDGF A chain          | 5.6             | M23238            | XL.841.3.S1_a_at     |
| WIG-related           | 5.3             | BJ044287          | XL.23988.1.S1_at     |
| CP2-like transcription factor | 4.3          | BJ046394          | XL.16094.1.A1_at     |
| Glucocorticoid inducible leucine zipper | 4.1  | BC043841          | XL.12378.1.S1_at     |
| Unknown               | 3.8             | AW147865          | XL.2077.1.A1_at      |
| WIG                   | 3.7             | AF310008          | XL.736.1.S1_at       |
| XANF1                 | 3.1             | X60099            | XL.131.1.S1_at       |
| HES-related 1B        | 3.0             | AB071434          | XL.12126.1.S1_at     |
| Darmin                | 2.9             | CD324819          | XL.6024.1.S1_at      |
| ODC2                  | 2.8             | AF217544          | XL.8949.1.S1_at      |
| Unknown               | 2.4             | AW460608          | XL.11598.1.A1_at     |
| Thioredoxin binding protein 2 | 2.3  | BQ399899          | XL.24749.1.A1_at     |
| Unknown               | 2.2             | BM192746          | XL.25985.1.A1_at     |
| Selenophosphate synthetase 1 | 2.1  | BJ091471          | XL.65222.1.A1_at     |
| Adenosine deaminase   | 2.1             | BJ090126          | XL.24155.1.A1_at     |
activation relative to the initiation of FGF signaling in the embryo can be quite different. For example, *sprouty2* is expressed at low levels maternally and the initial rise in levels of zygotic expression is detected at stage 9.5 which is 1 hour at 23°C after the activation of FGF8 expression. Subsequently, *sprouty2* expression continues to closely follow that of FGF8 during late blastula stages (stage 9 to 10). The initiation of *brachyury* and *Iro3* expression is somewhat delayed relative to initiation of FGF signaling, with a slight rise in expression by stage 9.5 and a more significant increase in expression by stage 10. Expression from *Cdx4* and *marginal coil* are even more delayed and their expression levels only begin to rise steeply from stage 10 onwards.

The differing dynamics of expression from known target genes indicate that there are different classes of FGF response genes. We

### Figure 3. Validation of FGF target genes.

(A) shows whole mount in situ hybridizations for genes positively regulated by FGF signaling at gastrula stage 10.5 in control embryos and embryos injected with 2 ng of dnFGFR1 mRNA. (B) shows the expression gene negatively regulated by FGF signaling in control embryos and embryos injected with 2 ng dnFGFR1 mRNA. All embryos are vegetal view with dorsal to the top. Non-uniform down regulation around the circumference of the blastopore is apparent in some embryos and is likely due to variability in the diffusion of injected dnFGFR mRNA. (C) is an RPA showing the expression at gastrula stage 10.5 of a number of genes in control animal cap explants and explants treated with FGF4 protein. 5 μg total RNA was used per hybridization. ODC is a loading control. doi:10.1371/journal.pone.0004951.g003

| Gene                        | Control | dnFGFR |
|-----------------------------|---------|---------|
| purine phosphorylase        | ![image](image1.jpg) | ![image](image2.jpg) |
| glycogen phosphorylase      | ![image](image3.jpg) | ![image](image4.jpg) |
| ephrin receptor A4           | ![image](image5.jpg) | ![image](image6.jpg) |
| spocb2                      | ![image](image7.jpg) | ![image](image8.jpg) |
| unknown xl.5479             | ![image](image9.jpg) | ![image](image10.jpg) |
| DUSP5                       | ![image](image11.jpg) | ![image](image12.jpg) |
| MKP1                        | ![image](image13.jpg) | ![image](image14.jpg) |
| methyltransferase           | ![image](image15.jpg) | ![image](image16.jpg) |

| Gene                        | Control | dnFGFR |
|-----------------------------|---------|---------|
| CP2-like                    | ![image](image17.jpg) | ![image](image18.jpg) |
| glucocorticoid inducible leucine zipper | ![image](image19.jpg) | ![image](image20.jpg) |

| Gene                        | Control | FG4 |
|-----------------------------|---------|-----|
| Cdx4                        | ![image](image21.jpg) | ![image](image22.jpg) |
| Esr5                        | ![image](image23.jpg) | ![image](image24.jpg) |
| purine phosphorylase        | ![image](image25.jpg) | ![image](image26.jpg) |
| glycogen phosphorylase      | ![image](image27.jpg) | ![image](image28.jpg) |
| ephrin receptor A4           | ![image](image29.jpg) | ![image](image30.jpg) |
| unknown xl.5479             | ![image](image31.jpg) | ![image](image32.jpg) |
| DUSP5                       | ![image](image33.jpg) | ![image](image34.jpg) |
| MKP1                        | ![image](image35.jpg) | ![image](image36.jpg) |
| methyltransferase           | ![image](image37.jpg) | ![image](image38.jpg) |
| ODC                         | ![image](image39.jpg) | ![image](image40.jpg) |
investigated this further by undertaking a cluster analysis of FGF dependent genes based upon their temporal expression profiles from early blastula to early neurula stages. For the generation of clusters of genes positively regulated by FGF signaling a correlation value of \( p \geq 0.85 \) was used. The cluster analysis of genes negatively regulated by FGF is not presented because this group contains considerably fewer genes leading to multiple clusters containing single genes.

The genes in each of the clusters used for further analysis are shown in Table 4. The dendrogram generated during cluster analysis is shown in Figure 5B. Figure 5C shows a heat map of the relative expression of each of the generated clusters from stage 8 to stage 14. The expression profiles of the clusters positively regulated by FGF signaling, together with that of FGF8, are shown in Figure 5D. In keeping with our findings for individual known FGF targets, the initiation of expression from each of the clusters relative to the activation of FGF signaling varies considerably. For example, activation of expression from genes in clusters #11 and #9 rapidly follows the activation of FGF8 expression. However, at 23°C activation of expression from genes in cluster #1 is delayed 2–3 hours relative to that of FGF8. Activation of expression from genes in clusters #7 and #8 occurs at an intermediate time point with a 1–2 hour lag relative FGF8 and the activation of FGF signaling.

Identification of a novel negative regulator of FGF signaling

The earliest activation of zygotic transcription from putative FGF target genes occurs between blastula stage 8.5 to 9. During this period a number of genes undergo \( >10 \)-fold increase in expression. Amongst these rapid responders Dual specificity phosphatase 5 (DUSP5) has the highest fold increase during this period (>35). *Xenopus laevis DUSP5* codes for a putative MAP kinase phosphatase with 61% peptide sequence identity to human DUSP5 (Figure S1).

Expression of another MAP kinase phosphatase, *MKP1/3* [33], is also significantly down regulated in response to
FGF inhibition (Table 1 and Figure 3). Interestingly, another Xenopus MAP kinase phosphatase, MKP3, has been shown to inhibit FGF dependent mesoderm induction [21] and is implicated as a component of a negative feedback loop regulating FGF activity in the embryo [34]. Given the critical role that ERK/MAP kinase activity has in mediating responses to FGF signaling in the early embryo we investigated the potential role of DUSP5 as determined by microarray analysis. Expression of all three MKPs rises rapidly during late blastula stages is as a regulator of gene transcription and it is this latter function which is the focus of the present study.

Figure 6A is a chart showing the temporal expression profiles of MKP1, MKP3 and DUSP5 as determined by microarray analysis. Expression of either dnFGFR1 or dnFGFR4a leads to very similar effects on gene expression at the start of gastrulation.

**Table 3. Effects of FGF inhibition on organizer gene expression.**

| Gene                | Ratio of expression in dnFGFR and control groups |
|---------------------|-----------------------------------------------|
| Egr1                | 12.4                                          |
| FoxD5A              | 10.8                                          |
| SIP                 | 9.8                                           |
| Paraxial P-C        | 7.3                                           |
| FoxD3A              | 6                                             |
| Frzb1               | 5.1                                           |
| Zic3A               | 4.5                                           |
| Xiro3               | 4.5                                           |
| Gravin-like         | 4.4                                           |
| Crescent            | 3.9                                           |
| Xspr2               | 3.7                                           |
| Protein 1a11        | 3.6                                           |
| FoxA4               | 3.4                                           |
| Chordin             | 2.9                                           |
| Dkk1                | 2.6                                           |
| ADMP                | 2.4                                           |
| Goosecoid           | 2.3                                           |
| Noggin              | 2.2                                           |
| Xlim1               | 1.9                                           |
| Xnr3                | 1.2                                           |
| Cerberus            | 1.2                                           |
| Er81                | 1.0                                           |
| Xnr4                | 0.9                                           |
| Siamois             | 0.9                                           |
| Otx5                | 0.8                                           |
| Xiro1               | 0.8                                           |
| Otx2                | 0.8                                           |

Figure 6B shows the degree to which expression of the three MKPs in the early gastrula is down regulated in response to FGF inhibition with dnFGFR1 and dnFGFR4a. Consistent with this, expression of MKP1 and DUSP5 in the circum-blastoporal region during gastrula stages is dependent on FGF signaling (Figure 3A). Similar FGF dependence has been reported for MKP3 [34]. We also show that FGF signaling is sufficient for MKP1 and DUSP5 expression; treatment of animal cap explants with FGF proteins leads to marked up regulation of both genes (Figure 3C).

Identification of transcriptional targets of FGF signaling

Our study was designed to identify early transcriptional responses to FGF signalling. The reagents that we used to inhibit FGF signalling were dominant negative mutants of FGF receptor 1 (dnFGFR1) and FGF receptor 4a (dnFGFR4a) [24,25]. Over expression of either dnFGFR1 or dnFGFR4a leads to very similar effects on gene expression at the start of gastrulation.

Using the strict criteria outlined, we have identified 67 genes which are down regulated and 16 genes which are up regulated in response to FGF signaling. The target validation undertaken indicates that these FGF targets lists are well supported and should provide the basis for further studies into FGF dependent transcriptional regulation.

Different waves of FGF dependent gene regulation

As part of this study we carried out a time course analysis of gene expression from mid-blastula to early neurula stages. Our
cluster analysis based on these expression profiles reveals that activation of expression from FGF dependent genes occurs in a number of waves following the mid-blastula transition (MBT). Some genes are activated very rapidly following the MBT and closely follow the expression profile of FGF8. Expression of other genes, including Brachyury and Cdx4, which are known immediate early response genes, activated by FGF signaling in the absence of protein synthesis, occurs in later waves [26,37]. At present it is unclear why some immediate early responses are more rapid than others. However, we speculate that the presence of identifiable clusters of FGF response genes indicates that similar upstream mechanisms are involved in regulating the expression of genes within the same cluster. The identification of such clusters of co-expressed transcriptional targets of FGF signaling will allow the analysis of these genes for shared regulatory elements required to drive their common modes of expression.

It is important to note that our analysis does not rule out the involvement of other signaling pathways in the regulation of the identified FGF target genes. Indeed this is to be expected, given that FGF signaling has been shown to interact with a number of pathways regulating gene expression in early development, including the activin/nodal and Wnt signaling pathways [38–41].

Patterns of FGF target gene expression

The initial zygotic expression of several FGF ligand genes, including FGF3, FGF4, FGF8 [42–44] and FGF20 (Figure S2) is restricted to the early mesoderm during late blastula stages. In keeping with this we find that many target genes positively regulated by FGF signaling are also expressed in the mesoderm. Our analysis of FGF dependent gene expression in later development shows that there is diversity in their later expression. However, we note that the posterior mesoderm, the paraxial
Table 4. Gene clusters positively regulated by FGF signaling.

| Cluster | Gene |
|---------|------|
| #1      | Alkaline phosphatase |
|         | Cdx4 |
|         | Cytochrome B561 |
|         | FoxC1 |
|         | Glycogen phosphorylase |
|         | Gravin-like |
|         | Lin28a homologue |
|         | Marginal coil |
|         | mitotic phosphoprotein 67 |
|         | p75-ike fullback receptor |
|         | Purine phosphorylase |
|         | Putative methyltransferase |
|         | Unknown |
|         | SIP1 |
|         | SMCT |
|         | Uncharacterised protein C2orf32- xl25136 |
|         | Unknown-xl.15382 |
| #7      | Dkk1 |
|         | FoxA4 |
|         | FoxD5A |
|         | Frzb1 |
|         | NADH dehydrogenase sub-unit |
|         | Pinhead |
|         | Unknown-xl.3023 |
|         | XSpr2 |
|         | Zic3A |
| #8      | Brachyury |
|         | Chordin |
|         | Ephrin receptor A4 |
|         | FoxD3A |
|         | Glut1 transporter |
|         | Paraxial protocadherin |
|         | Retrotransposon protein 1a11 |
|         | Unknw-xl.18179 |
|         | unknown-xl.5479 |
| #9      | Egr1 |
|         | FoxA4 |
|         | Goosecoid |
|         | Prickle |
|         | Related to DC-STAMP domain receptor |
| #11     | ADMP |
|         | DUSP5 |
|         | Ets5 |
|         | Noggin |
|         | Sprouty2 |
|         | Wnt8 |
|         | Xom |

FGF regulation of organizer gene expression

Early studies of FGF function in early amphibian development focused on their potential role as regulators of gene expression in the ventro-lateral mesoderm [35, 45, 46]. However, more recent studies have provided evidence that FGF signaling is also required for the expression of genes within the dorsal organizer region of the amphibian embryo [47–49]. The large scale analysis of gene expression provided by our microarray experiments show that FGF signaling is required for the normal expression of a large number of organizer genes, including goosecoid, chordin, noggin, dkk1 and Frzb, and a number of genes, such as Sipt1 and FoxD5, which are expressed in the dorsal neuroectoderm (See Table S14). Such a role is very much in keeping with the observed activity of MAP kinase signalling in the dorsal marginal zone and dorsal neuroectoderm, and supports the view that FGF signaling is required during late blastula and early gastrula stages for the establishment of both the Spemann organizer and the presumptive neuroectoderm.

There are also indications that FGF signaling is required to negatively regulate and therefore restrict gene expression in the dorsal region of the embryo. For example, the transcription factors Hes1b is up regulated in response to FGF inhibition. We note that Hes1b is expressed in the dorsal region of the gastrula but at some distance from the highest levels of FGF activity in the blastopore region [50]. It is also seems likely that negative regulation by FGF restricts XANF1 expression to the deeper layers of the organizer region in the early gastrula [51].

Gene function downstream of FGF signaling

A detailed description of the putative function for each of the identified target genes is beyond the scope of this discussion and we refer the reader to the extensive annotation and literature resources provided in Tables S2, S3, S4, S5, S6, S7, S8, S9, S10 and S11. A number of the genes identified in the screen are of unknown function either because they are orthologs of genes with poorly characterized function or because we were unable to identify orthologous genes in the databases and may therefore represent novel *Xenopus* genes. However, analysis of the annotated genes identified in the screen reveals that a large proportion of the genes regulated by FGF signaling, are themselves also involved in gene regulation, either directly, as in the case of transcription factors, or via involvement in other signaling pathways. This indicates the key position of FGFs as upstream regulators of genetic pathways leading to germ layer specification during the late blastula to early gastrula stage of amphibian development. In addition, FGF signaling is required for the normal expression of genes such as Prickle, marginal coil and Ephrin receptor A4 (pagliaccio) which are involved in regulating cell movements and adhesion during gastrulation [52–54].

We also note that the *purine phosphorylase* and *glycogen phosphorylase* genes, which code for enzymes involved in nucleotide and carbohydrate metabolism respectively, are dependent on FGF
signaling and are expressed in highly restricted, FGF associated domains in the early mesoderm. Further studies will be required to determine if these genes have previously unsuspected roles in the regulation of developmental mechanisms.

Previous studies have identified a number of FGF inducible inhibitors of the FGF signaling pathway, including the Sprouty genes, which we find are significantly down regulated in our screen [36,55]. In the present study we identify the MAP kinase phosphatase genes MKP1 and DUSP5 as FGF targets which are activated rapidly following the mid-blastula transition and show that they are able to inhibit FGF signalling in *Xenopus* tissues. DUSP5 is a novel *Xenopus* MKP which is expressed in the early mesoderm and neural plate in a pattern which is remarkably similar to that of *Xenopus* FGF3 [44]. Similar correlation with sites of FGF activity has been reported for MKP3, which has also been shown to act as a feedback inhibitor of FGF signaling [34,56].

The picture that emerges is that activation of FGF signaling induces the production of multiple inhibitors which act to moderate and limit the response to FGF signaling. A number of positive feedback mechanisms also impact on the FGF pathway, including the transcriptional activation the FLRT3 and brachyury genes [6,57,58]. FLRT3 is a transmembrane protein that

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**Figure 6. MKPs and FGF signalling.** (A) shows the temporal expression profiles of MKP1, MKP3 and DUSP5 from blastula stage 8 until early neurula stage 14 (5 to 16 hours pf at 23°C). Profiles are derived from normalised microarray expression levels. Relative expression values are represented as percentages of the maximum expression value for each gene. (B) is a chart showing the expression of MKP1, MKP3 and DUSP5 in control embryos and embryos injected with 4 ng dnFGFR1 mRNA or 4 ng dnFGFR4 mRNA. Microarray derived expression values are based on the average of three biological replicates. Relative expression is calculated a percentage of the expression in control embryos. Experiments in (C, D and E) were carried out on animal cap explants removed from blastula stage 8 embryos. In all cases control explants are from un.injected embryos, FGF4 treatment was with 10 units of recombinant protein and mRNA injections were with 10 ng MKP1, MKP3 or DUSP5. (C) shows the morphology of animal cap explants at tailbud stage 41. (D) shows 10 μm histological sections of animal cap explants at stage 41. (E) is a Western blot showing levels of dp-ERK and the loading control GAPDH in animal cap explants at stage 10.5.

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potentiates FGF signal transduction and brachyury is a T-box transcription factor which has been shown to be a component of a positive feedback loop that leads to increased transcription of FGF ligand genes in the early mesoderm [5,6,30]. The presence of positive and negative feedback loops which modulate FGF signaling at multiple levels highlights the critical importance for fine tuning the overall levels of FGF signaling during development. An interesting observation is that expression of the FoxD3 and Lin28 genes are down regulated in response to FGF inhibition. The FoxD3 transcription factor is linked to amphibian organizer function [59] but has also been implicated as a component of the pluripotency circuit of mammalian embryonic stem cells via regulation of the nanog gene [17,18]. The Lin28 gene codes for an RNA binding protein, which together with the Okt4, Sox2 and Nanog genes, can convert somatic cells to an embryonic stem cell phenotype [14].

Previous studies have indicated that FGF signaling is required as a competence factor necessary for the response of embryonic amphibian cells to activin-like signals during development [39,40,60]. We also note a recent study which showed that FGF signaling in murine ES cells is necessary to allow differentiation into multiple lineages, including mesoderm [12]. These observations raise the intriguing possibility that FGF signaling, acting via downstream targets such as Lin28 and FoxD3, might have a general role in regulating pluripotency or the competence of embryonic cells to respond to signals which direct differentiation both during normal development and in culture.

Materials and Methods

Ethics statement

All animal work was undertaken under a licence from the UK Home Office.

Embryological methods

Embryos were staged according to Nieuwkoop and Faber [61]. Normal embryos were cultured in NAM/10. Injection of embryos with synthetic mRNAs was carried out in 33% NAM+5% ficoll (Sigma) at the 2 or 4-cell stage. Animal caps were explanted from embryos in 50% NAM Recombinant FGF4 [43] treatment was in 50% NAM+ 5 mg/ml BSA.

Identification of Xenopus tropicalis full length clones

Clones containing full length coding region of X.tropicalis MKP1 (DUSP1/XCL100) (accesion number AL697333) and DUSP5 (accesion number AL648624) were identified using the peptide sequence of X.laevis orthologues (accession numbers NM_001088684 and BJ067398 respectively) and BLASTP on the Sanger Institute X.tropicalis EST database (www.sanger.ac.uk/Projects/X_tropicalis/).

mRNA synthesis

Capped mRNA was synthesised using the SP6 Megascript kit (Ambion) and a modified protocol using a 1:10 ratio of GTP to m7G(5')ppp(5')G cap. All CDNAs used for mRNA were in either pSP64t or CS2+ or CS107 and were transcribed using SP6 polymerase. The dominant negative X.laevis FGFR1 (dnFGFR1) plasmid was a gift from Enrique Amaya [24]. The dominant negative X.laevis FGFR4a (dnFGFR4) plasmid was a gift from Harumasa Okamoto [25]. The X.laevis MKP3 (DUSP6/X17C) plasmid was a gift from Bob Old [62].

In situ hybridisation

Embryos were fixed in MEMFA and in situ hybridizations were carried out as per [63] with the modifications described in [64].

Details for in situ probe plasmids are shown in Table S15. The sources of the plasmids, including Geneservice (www.geneservice.co.uk) and the NIBB/NIG/NBRP X. laevis EST project (xenopus.nibb.ac.jp) are indicated.

RNAase protection analysis

RNA extraction and RNAase protection analysis were carried out according to the methods of Pownall et al. [1996]. Data relating to RNAase protection plasmids are shown in Table S16.

Whole-mount immunohistochemistry and Western blotting

dp-ERK immunohistochemistry was carried out according the methods of [22]. Western blot samples were homogenized in PhoshoSafe (Novagen) extraction buffer per embryo. Following centrifugation supernatants were subjected to SDS-PAGE. Gels were blotted onto Immunoblot-P (Millipore) transfer membrane. Antibody concentrations were mouse anti-dp-ERK (Sigma), 1:8000, anti-phosphoSnad1/5/8 (NEB), 1:500, anti-GAPDH (HyTest), 1:3000. 1:3,000, anti-GAPDH (HyTest), 1:1,000,000, anti-mouse POD (Amersham), 1:3000 and anti-rabbit POD (Amersham), 1:2000. Peroxidase activity was detected using BM chemiluminescence blotting substrate (Roche) and Hyperfilm (Amersham).

Embryos for microarray experiments

Embryos were injected with 4 ng of dnFGFR1 or dnFGFR4 mRNA and were collected with sibling controls at early gastrula stage 10.5 (11 hours post-fertilization at 23°C). In order to enable statistical analysis three biological replicates were produced by in vitro fertilization from different pairs of male and female frogs. Each replicate set comprised control, dnFGFR1 injected and dnFGFR4 injected embryos. Before processing for microarray analysis each replicate set was assessed for the effectiveness of FGF signaling inhibition by analysing levels of dp-ERK and levels of the known FGF targets Xbra, Cdx4 and myoD expression in sibling embryos.

For the early developmental timecourse embryos from a single mating were cultured at 23°C and collected at 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15 and 16 hours post fertilization (NF stage 8, 8.5, 9, 9.5, 10, 10+, 10.5, 11, 12, 12.5, 13 and 14) for microarray analysis. Sibling embryos were also collected at the same time points for Western blot analysis of dp-ERK.

Preparation of total RNA for microarray analysis

Batches of 10 embryos were extracted in Tri-reagent according manufacturer’s protocol (Sigma). RNA was precipitated using isopropanol and cleaned up using the Qiagen RNaseq kit followed by a lithium chloride precipitation. Quality of RNA was assessed using the Agilent 2100 Bioanalyzer.

Preparation of labelled cRNA and chip hybridization

2 μg of total RNA was processed for the microarray by using the Affymetrix GeneChip one-cycle target labelling kit (Affymetrix) according to the manufacturer’s recommended protocols. The quality and quantity of the resulting bioinylated cRNA was determined by using NanoDrop ND 1000 (NanoDrop Technologies) and Agilent Bioanalyzer 2100 (Agilent Technologies). Biotin-labelled cRNA samples were fragmented randomly to 35–200 bp at 94°C in fragmentation buffer (Affymetrix) according to the manufacturer’s recommended protocols and aliquots of the fragmented cRNA were run on the Agilent 2100 Bioanalyzer to assess the quality of the generated cRNA.
Biotin-labeled fragmented cRNA samples were combined with hybridization buffer containing hybridization Control cRNA and Control Oligo B2 (Affymetrix), before hybridization to GeneChip® Xenopus laevis Genome Array for 16 h at 45 ºC. The arrays were washed, stained, and scanned using the Affymetrix Model 450 Fluidics Station and Affymetrix Model 3000 scanner using the manufacturer’s recommended protocols.

Microarray data analysis

Affymetrix microarray experiments were conducted in accordance with the MIAME standards requirements [65]. Raw data processing was performed by using the Affymetrix GCOS 1.2 software. After hybridization and scanning, probe cell intensities were calculated and summarized for the respective probe sets by means of the MAS5 algorithm. To compare the expression values of the genes from chip to chip, global scaling was performed, which resulted in the normalization of the trimmed mean of each chip to a target intensity (TGT value) of 500 as per manufacturers documentation. Each sample and hybridization underwent a quality control evaluation checking for adequate scaling factors (1–3 for all samples), percentage of probe sets reliably detected (between 40–60% present call), and optimal 3'/5' hybridization ratios for the housekeeping genes (e.g., GAPDH, poly(A) spike-in controls, and the proaryctotic controls (bioB, bioC, bioD and cre).

Data were imported into BRB ArrayTools software version 3.6.0 (http://linus.nci.nih.gov/BRB-ArrayTools.html). Imported array data were filtered using the following criteria.

1) **Spot filters**- Threshold minimum value if spot intensity below 5.
2) **Normalization**- Normalize (center) each array using median over entire array.
3) **Exclude a gene under any of the following conditions**- Percent of data missing or filtered out exceeds 50%. Percent of absent (i.e., Detection Call = A) data exceeds 50%.

Scatterplots were generated using the phenotype averages tool of BRB ArrayTools. Gene lists of FGF targets were generated using the BRB between groups of arrays class comparison tool (unpaired, two sample t-test with random variance model and nominal significance level p = 0.01). An additional filter excluded genes with less than 2-fold difference from controls.

Temporal expression profiles for a given gene were generated in Microsoft Excel by plotting relative expression at each time point as a percentage of the maximum expression level within the time course. Cluster analysis was undertaken using the BRB gene cluster analysis tool (complete linkage and centred correlation). The Affymetrix Cel files for all microarray experiments are available at EMBL ArrayExpress, accession numbers E-MEXP-2058 and E-MEXP-2059.

Target gene annotation

Target gene annotation was accomplished using a combination of existing Affymetrix Gene array annotation and BLAST searching of target sequences against Genbank, Swiss-prot (www.ncbi.nlm.nih.gov/BLAST/) and NIBB/NIG/NBRP Xenopus laevis EST project (xenopus.nibb.ac.jp) databases.

Supporting Information

**Figure S1** Alignment of amphibian and human DUSP5 peptide sequences. Alignment of the peptide sequences for human and Xenopus tropicalis DUSP5 produced by the Clustal W method. Identical residues are boxed in red.
Table S11 Genes negatively regulated by FGF signaling of unknown function
Found at: doi:10.1371/journal.pone.0004951.s013 (0.03 MB DOC)

Table S12 GO terms for genes positively regulated by FGF signaling
Found at: doi:10.1371/journal.pone.0004951.s014 (0.10 MB DOC)

Table S13 GO terms for genes negatively regulated by FGF signaling
Found at: doi:10.1371/journal.pone.0004951.s015 (0.04 MB DOC)

Table S14 Expression of genes positively regulated by FGF signaling
Found at: doi:10.1371/journal.pone.0004951.s016 (0.38 MB DOC)

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