Minireview

An amphibian with ambition: a new role for *Xenopus* in the 21st century

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Abstract

Much of our knowledge about the mechanisms of vertebrate early development comes from studies using *Xenopus laevis*. The recent development of a remarkably efficient method for generating transgenic embryos is now allowing study of late development and organogenesis in *Xenopus* embryos. Possibilities are also emerging for genomic studies using the closely related diploid frog *Xenopus tropicalis*.

Xenopus as a model organism

Amphibians have played a key role in the elucidation of the mechanisms of early development over the last century. The eggs and embryos are relatively large and robust, and they develop externally in a simple salt solution. Because of their suitability for micromanipulation, amphibian embryos provided many of the core results of experimental embryology. These include the provision of fate maps (stage-specific diagrams indicating where cells will end up and which tissues they will form), states of developmental commitment of cells to different lineages, and inductive signals that allow one cell type to influence the development of another. Since its introduction as a model organism in the 1950s, most of this work has focused on the African clawed frog *Xenopus laevis*. The eggs and embryos of *X. laevis* can be produced in large numbers by means of a simple hormone injection and are, like other amphibian embryos, easily manipulated, injected, grafted or labeled. In the molecular era, the ability of the *Xenopus* embryo to translate injected, synthetic mRNA has led to many important contributions to the study of early developmental events. In particular, the unique advantages of *Xenopus* described above have allowed the identification of the major classes of inducing factor used by all animals (the transforming growth factor β (TGFβ)/bone morphogenetic protein, fibroblast growth factor (FGF) and Wnt classes). In fact, *Xenopus* would seem to be the ideal all-purpose research organism, except for the fact that it is not ideal for experimental genetics given its pseudotetraploid genome and long generation time. Recently, however, the emergence of a potential ideal model organism, the closely related diploid species *Xenopus tropicalis*, along with the development of transgenic methods and the initiation of a community-wide initiative for the production of genomic and genetic resources for the future, has secured the future of *Xenopus* as a genetic model as well.

Xenopus ESTs in expression screening and microarray analyses

The sequencing of expressed sequence tags (ESTs) from *X. laevis* has lagged behind efforts on many other model organisms and humans, in part because of the pseudotetraploid nature of the *X. laevis* genome. Acquiring large numbers of ESTs has recently been recognized as a priority for future *Xenopus* research, and, as a result of several parallel efforts, the number of deposited ESTs has now exceeded 100,000. According to the National Center for Biotechnology Information (NCBI [1]), this now places *Xenopus* eleventh in the ranking of species in terms of available EST numbers. Information on *Xenopus* ESTs can also be obtained from the XEST website [2].

Although there is still a long way to go, the *Xenopus* EST collection is already starting to have an impact on research. A
large-scale gene-expression screen was performed by Gawan-
tka and colleagues [3] and has formed the basis for the search-
able Axeldb expression-pattern database [4,5]. In Gawan-
tka’s initial screen [3], 1,765 random Xenopus cDNAs were ana-
alyzed by whole-mount in situ hybridization to embryos. Of these, 273 unique, differentially expressed genes were identi-
fied, 204 of which had not been isolated previously from Xenopus. The relatively large size of the screen enabled Gawan-
tka and colleagues [3] to identify four putative ‘syn-
expression’ groups, or sets of genes with identical expression
patterns, two of which, named ‘delta’ and ‘BMP4’ after well-
known genes in the groups, were predicted to correspond to
molecular pathways involved in patterning and differen-
tiation. When you consider that the number of marker genes that
have been submitted by the entire research community to the
Xenopus Molecular Marker Resource (XMMR) [6] is less than
50, this is an impressive contribution from a single lab.

One of the forces driving the accumulation of ESTs is the wish
to produce Xenopus microarrays, which can be used for
genome-wide gene-expression analysis. Xenopus is ideal for
this kind of screening, as it is relatively easy to isolate specific
regions of the embryo in order to make tissue-specific RNA
probes. Traditionally, researchers attempting to characterize
a phenotype arising from their experiment would analyze
changes in the expression levels of a small panel of genes that
have well-documented expression patterns, known as
markers, by reverse-transcriptase-coupled (RT) PCR, RNase
protection assay, or Northern blot. Although this gives a good
idea of the molecular changes that have occurred, it might be
to possible to miss key events if markers are not chosen care-
fully or if appropriate markers are not available. In a recent
paper, Altmann and colleagues [7] have demonstrated the power of using Xenopus microarrays to address several key
aspects of early development. Their prototype microarray
contained 864 sequenced cDNAs prepared from embryos at the
gastrula stage of development, 107 of which were not re-
presented in the EST database and 96 of which were previ-
ously characterized markers (see the Xenopus microarray site
[8]). Three test experiments were performed using the array.
First, the authors searched for temporally regulated genes
using either maternal or zygotic mRNA to generate probes;
second, they compared mesoderm from dorsal and ventral
regions of early gastrula embryos; and third, they compared
untreated embryonic explants with those treated with the
known mesoderm inducer activin. Novel temporally regu-
lated, spatially restricted and activin-induced genes were iso-
lated using this screen, and the results were confirmed by
independent analysis using RT-PCR or in situ hybridization.
Importantly, probes could be generated from a quantity of
RNA equivalent to the amount in a single embryo, although
this was thought to be the absolute minimum requirement

Another use for the growing EST dataset is in the construc-
tion of UniGene sets for Xenopus. These are created auto-
matically from the NCBI database, which includes both ESTs
and well-characterized genes, by forming clusters of
sequences, each representing a unique gene (see NCBI’s
UniGene [9]). The usefulness of this type of information is
twofold. Firstly, the ideal microarray would contain a single,
representative copy of each expressed gene - in other words,
a non-redundant gene set. Secondly, a program has been
developed by the NCBI that makes use of the UniGene data
to enable ‘digital differential display’ (DDD) [10]. Each of
the libraries that have been used to generate ESTs can be com-
pared using DDD, to show the genes that are significantly
differentially represented between them. For example, a
library should have generated more EST hits for the
albumin gene than would a heart library, and this will be dis-
played as a darker spot on the DDD (see Figure 1). As the
number of ESTs increases and more libraries are used for
sequencing, this should become a very powerful tool.

Transgenic technology for studying later
development and organogenesis

Many of the experiments using Xenopus in the study of early
development have made use of injected mRNA, antibodies,
or ordinary or morpholino antisense oligonucleotides. These
methods are all essentially transient, however, as the
genome is not altered and the injected substance decays with
time. Arguably the biggest leap forward in the establish-
ment of Xenopus as a model organism beyond the limits of early

| A | Liver | B | Heart | Gene index | Gene description |
|---|-------|---|-------|------------|----------------|
| 0.16726 | 0.00000 | XI.395 | X. laevis 74 kDa serum albumin mRNA, complete cds |
| A>B | | | | |
| 0.00000 | 0.05747 | XI.3102 | ESTs, moderately similar to Actin 11 [Arabidopsis thaliana] |
| B>A | | | | |
| 0.04821 | 0.00000 | XI.5021 | X. laevis mRNA for transferrin |
| A>B | | | | |
| 0.00000 | 0.4807 | XI.1285 | X. laevis mRNA for myosin light chain |
| B>A | | | | |

Figure 1
Example of digital differential display (DDD [10]), comparing
gene expression between the liver (pool A) and heart (pool
B) by comparing the available EST data from two Xenopus
libraries. The top four results are shown. The numbers in
grey represent the proportion of sequences within each
pool that map to the UniGene cluster indicated in blue, for
which a description is given on the right; cds, coding
sequence. Dots are a visual representation of the numerical
values. Statistically significant results are shown by indication
of the relationship between two pools for a particular gene
cluster, for example A>B for albumin and transferrin in the
liver, B>A for myosin light chain and a probable actin
homolog in the heart library.
development has been the development of methods for generating transgenic embryos, by Enrique Amaya and Kristen Kroll [11,12]. Known as restriction enzyme mediated insertion (REMI) transgenesis, this method has been adopted across the Xenopus community with great enthusiasm. The key to REMI transgenesis involves mixing transgene DNA with purified and permeabilized sperm, along with a small quantity of restriction enzyme and an extract of egg cytoplasm generated by high-speed centrifugation (the key ingredient of which is thought to be nucleoplasmin, which serves to partially decondense the sperm chromatin, facilitating transgene insertion); this mixture is then injected into dejellied eggs at the rate equivalent to about one sperm per egg. Around a third of the recipient eggs usually cleave normally and a proportion (around 50%) of these integrate the transgene into the genome during the first cell cycle. This method produces large quantities of non-mosaic transgenics in the parent (Fo) generation, allowing experiments to be performed without the generation of stable transgenic lines. Importantly, several labs have shown that the transgene is reliably transferred through subsequent generations [13-15].

A key advantage of Xenopus transgenesis is the ability to study transgene expression in living embryos using green fluorescent protein (GFP) as the reporter; this allows quick and easy promoter analysis [15-19]. Furthermore, the advantages of Xenopus embryos allowing the later roles of genes involved in early patterning to be elucidated (these were previously accessible only using the far more difficult procedure of making conditional knockouts in mice). Already the technique has been used to observe the onset of differentiation of the lens [15], to study the role of bone morphogenetic proteins in heart looping [20], and to investigate several aspects of metamorphosis [21-24]. In our lab, we have been making use of tissue-specific promoters to study the formation of the gut organs [25] (Figure 2). With the adaptation of existing methods to Xenopus transgensics - for example, the Cre and FLP conditional mutagenesis systems [26] and tetracycline-inducible constructs [27] - it is becoming clear that the golden age for studying late development is fast approaching.

**A genomic basis for Xenopus research**

Although the establishment of a genomic base for Xenopus research is still in the planning stages, there are pilot mutagenesis studies already available. Amaya and colleagues [14] have shown that it is possible to adapt the transgenesis technique in order to trap genes with interesting, tissue-specific expression. Promoterless constructs containing GFP as the reporter, with or without splice-site acceptor sequences, were used to visualize ‘trapped’ genes (close to enhancers) in living Fo transgenics. Rapid PCR amplification of cDNA ends (5’ RACE PCR) was then used to amplify the disrupted genes from reverse-transcribed RNA extracted from the embryos, and the genes were identified by sequencing. Although no novel genes were identified in this pilot screen, a variety of tissue-specific expression patterns were generated, showing the potential power of the technique. In the future, it should be possible to use transposons to increase the rate of gene trapping (P. Mead, personal communication). It is also possible to isolate developmental mutants using inbreeding techniques, in which siblings are interbred for several generations to create ‘isogenic’ lines, or ‘gyngenesis’, in which haploid individuals are made homozygous by suppression of the first cleavage. In one study, 12 heritable developmental mutations were isolated from just eight wild-caught *X. laevis* females [28].

Despite the many advantages of Xenopus as a model organism, it does have one or two major drawbacks for genetic studies. Firstly, *X. laevis* is a pseudotetraploid, as a result of an additional genome duplication (relative to other vertebrates).

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**Figure 2**

Analysis of gut-specific promoters in live *X. laevis* tadpoles. Green fluorescence is seen where the promoter is active, and yellow is due to autofluorescence of the tadpole gut. Black patches are pigment cells. (a) The *Elastase* enhancer (rat) driving the expression of GFP in the pancreas of a 7-day-old tadpole; ventral view, anterior to the right. (b) The *Transthyretin* promoter (mouse) driving expression of GFP in the liver of a 6-day-old tadpole; ventral/left view, anterior to the left. (c) The *Intestinal fatty acid binding protein* promoter (rat) driving expression of GFP in the small intestine of a 7-day-old tadpole, ventral view, anterior to the left. Abbreviations: gb, gall bladder; li, liver; p, pancreas; si, small intestine; st, stomach. Reproduced with permission from [25].
about 30 million years ago [29,30]. This is also the case for zebrafish, another favorite model organism, as genome duplication also occurred in the teleost fish around 420 million years ago [31,32]. Aside from the larger genome that results from pseudotetraploidy, mutagenesis screens are less likely to be successful given the functional redundancy between closely related paralogous genes. The second disadvantage of *X. laevis* is the relatively long generation time, usually around 1-2 years, making the generation of stable transgenic lines a slow process. Fortunately, a closely related frog, *Xenopus* (formerly *Silurana*) *tropicalis*, has neither of these disadvantages, while retaining the many advantages of its larger relative. *X. tropicalis* has a diploid genome and a generation time of around 4-5 months [30,33]. *X. tropicalis* adults are also smaller, making them more practical for large-scale laboratory use, and they have a genome size of around half the size of that of the mouse (1.7 x 10⁹ base pairs per haploid nucleus [34]). Like *X. laevis*, they can be induced by hormone injection to lay around 1,000-3,000 eggs at a time. Although their eggs are around half the size of those of *X. laevis*, they are still big enough to be used for transgenesis and other manipulations [15]. Preliminary studies in Robert Grainger’s lab have shown that it is possible to use existing *X. laevis* probes for in situ hybridization studies on *X. tropicalis* (see the Grainger laboratory’s *X. tropicalis* site [35]), thus removing the need to reclone existing genes in *X. tropicalis*.

Driven by the recent advances in genetics, the *Xenopus* community is now working together to secure the large amounts of funding needed for development of genomic tools, resource centers and the *X. tropicalis* system. Details of the current proposals can be found on the NIH Xenopus Initiative website [36] and include both an expansion on the work discussed here and descriptions of new projects. For example, the planned creation of a genetic map for *Xenopus* will make use of the same methods as were used for zebrafish, such as radiation hybrid panels [37] and random amplified polymorphic DNAs (RAPDs) [38]. Peter Vize’s Xenbase [39], the main website resource for *Xenopus* researchers, is expanding to include sections on genetics and genomics, and further expansion of this facility is planned. As it becomes available, genomic technology, combined with the advantages of *Xenopus* as a model system, will allow *Xenopus* researchers to continue to make an extensive contribution to functional studies in the future.

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