Lineage-specific Gene Expression in the Sea Urchin Embryo

Division of Biology, California Institute of Technology, Pasadena, California 91125

Within a few days of fertilization, the sea urchin embryo develops into a small differentiated organism consisting of about 1800 cells and capable of feeding, swimming, and the further ontogenic transformations required in the succeeding weeks of larval growth. A number of distinct cell lineages that are clearly specialized at the morphological and functional levels can be discerned in the advanced embryo, and many of these can be traced back to particular sets of early blastomeres. Classical cell lineage and experimental studies (Hörstadius 1939; for review, see Angerer and Davidson 1984) have shown that certain of these lineages appear to be specified, at least in part, in consequence of the maternal components inherited in those regions of egg cytoplasm occupied by their progenitor cells. Specification of others among the early cell lineages clearly depends on inductive interactions that occur between blastomeres during cleavage. For the molecular biologist, as for his predecessors, this rapidly developing and simply constructed embryo offers the advantages of experimental accessibility. Thus, in respect to direct molecular-level analyses of gene activity in the embryo, for both specific genes and overall transcript populations and their protein products, the sea urchin is at present the best known embryonic system (e.g., reviews of Hentschel and Birnstiel 1981; Davidson et al. 1982; Angerer and Davidson 1984).

A focus of recent efforts in our laboratory has been the acquisition of a library of cloned genes that are expressed in the early embryo in a lineage-specific manner, for use in examination of the molecular processes by which these genes are differentially specified for activity early in development. This is, of course, the fundamental and general problem in understanding how the zygote gives rise to a functionally differentiated embryo, even given the well-established cytoplasmic anisotropy of most eggs (reviewed by Davidson 1976). It is a problem that is far from being solved for any embryo. Furthermore, diverse solutions may well be utilized in different modes of development. In the following, we review current progress on the isolation and characterization of lineage-specific sea urchin embryo genes, and then describe briefly recent studies that demonstrate apparently correct ontogenic expression of such genes after microinjection into the unfertilized egg.

The Sea Urchin Actin Genes

There are eight actin genes per haploid genome in Strongylocentrotus purpuratus, of which two are probably pseudogenes (Lee et al. 1984). All of the remaining six are expressed in the embryo. Of these, one is a muscle actin gene (M) and the others code for cytoskeletal (Cy) actin proteins (Durica et al. 1980; Scheller et al. 1981; Lee et al. 1984; Shott et al. 1984). Contrary to the implication of the cliché that cytoskeletal actin genes are "housekeeping" genes, our analysis of the patterns of expression of the individual members of this gene family has shown that each gene is expressed according to a specific ontogenic program. The genomic linkages of these genes, the nomenclature by which they are designated, and their patterns of activity in embryo and adult cell types, are summarized in Figure 1. Although their protein-coding regions are largely homologous, these genes differ greatly in the 3′ nontranslated trailer sequences of their mRNAs (Scheller et al. 1981; Lee et al. 1984), and this affords a means of preparing gene-specific molecular probes to identify their transcripts individually. Such probes have been utilized in RNA gel blot and in situ hybridization studies for determining the spatial and temporal patterns of actin gene expression in the embryo (Angerer and Davidson 1984; Shott et al. 1984; K.H. Cox et al., in prep.). As indicated in Figure 1 the expression of several of the actin genes is strictly lineage specific, and none is active ubiquitiously. For example, it is shown in Figure 2 that transcripts of the M actin gene are confined to two bilateral patches, each containing 10–20 cells that are associated with the newly formed coelomic pouches in the late embryo. These cells are of the secondary mesenchyme cell lineage, and their role is the construction of the pharyngeal muscle required for larval feeding. Expression of the CyIIIa and CyIIIB cytoskeletal actin genes is also confined to a single embryonic cell lineage, although CyIIIa messages are present as well in maternal RNA at the low level of about 1200 molecules per egg (J.J. Lee et al., unpubl.). They are not spatially localized in the egg (K.H. Cox et al., in prep.). After fertilization, newly synthesized transcripts of the CyIIIA and CyIIIB actin genes appear only in aboral ectoderm cells, although on different schedules. An in situ hybridization experiment carried out with a CyIIIA probe that demonstrates the early localization of these transcripts is shown in Figure 3 (Angerer and Davidson 1984; K.H. Cox et al., in prep.).

Expression of the CyIIA gene is restricted to mesenchyme cells, and in late embryos to parts of the gut. The CyI and CyIIIB genes are regulated similarly to one another, and their transcripts are more widely distributed than those of the other actin genes. These messages are also found in small quantities in the unfertil...
Figure 1. Organization and expression of actin gene family in the sea urchin *S. purpuratus*. Actin genes are named M (muscle) or Cy (cytoskeletal, i.e., expressed in nonmuscle cell types). Roman numerals designate the three nonhomologous 3' nontranslated trailer sequences found in the cytoskeletal actin genes, a, b, and c designate different though homologous trailer sequence variants. Linkage data are from analyses of cloned genes (Lee et al. 1984; R. Akhurst, F. Calzone, R. Britten, and E. Davidson, in prep.). Expression patterns in adult tissues were determined by RNA gel blot hybridizations, as reported by Shott et al. (1984). Expression of actin genes in embryonic cell types is summarized from the in situ hybridization study of K.H. Cox et al. (in prep.).

The number of mRNA molecules produced during embryonic development by five of the six functional actin genes has recently been measured by J.J. Lee et al. (unpubl.). These estimates were obtained by a single-strand probe excess titration method (Wallace et al. 1977; Scheller et al. 1978; LeV et al. 1980) using RNA transcripts synthesized in vitro from an Sp6 promoter (Butler and Chamberlin 1982; Melton et al. 1984). Since the cell types in which the various actin genes are expressed are known, the number of molecules of each mRNA species per active cell can be calculated from these data. These results are summarized in Table 1.

Figure 2. Expression of M actin gene in bilateral pharyngeal muscle anlage cells, visualized by in situ hybridization. Phase photomicrograph of embryo, an 82-hr pluteus, is shown at the left. (Reprinted, with permission, from K.H. Cox, L.M. Angerer, J.J. Lee, E.H. Davidson, and R.C. Angerer, in prep.)
Gene transfer experiments described briefly in a following section of this report have focused on the CyIIIa actin gene. This is the single most intensely expressed of all the actin genes in the embryo and its activity is confined to embryonic and larval stages. Thus, it is not utilized at all in the postmetamorphosis sea urchin (Shott et al. 1984). As shown in Table 1, late in embryogenesis there are about 200 molecules of CyIIIa mRNA per aboral ectoderm cell. The time course of CyIIIa gene expression during early development, as established by titration measurements, is shown in Figure 4. Within a few hours beginning at the very early blastula stage, new transcripts of this gene appear, and they accumulate dramatically to a level of over 80,000 molecules per embryo. Nuclear run-off experiments (S. Johnson, R. Britten, and E. Davidson, unpubl.) indicate that this accumulation is regulated primarily at the transcription level.

### Other Lineage-specific Cloned Genes Active in the Sea Urchin Embryo

Two other sea urchin genes that, like the CyIIIa and CyIIIb actin genes, are expressed specifically in aboral ectoderm have been characterized by Dr. William Klein and his associates (Bruskin et al. 1981, 1982; Lynn et al. 1983; Carpenter et al. 1984; S.L. Houck, C.D. Carpenter, P.E. Hardin, A.M. Bruskin, and W.H. Klein, unpubl.). These genes, called Spec1 and Spec2 (Spec, *S. purpuratus* ectoderm), code for intracellular Ca$^{2+}$-binding proteins related to troponin C. The aboral ec-

---

**Table 1 Transcripts per Cell for Five Actin Genes Expressed in the Embryo of S. purpuratus**

<table>
<thead>
<tr>
<th>Gene</th>
<th>Approximate number of cells transcribing actin mRNA per embryo</th>
<th>Number of actin transcripts per embryo</th>
<th>Number of actin transcripts per expressing cell</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cyl</td>
<td>1000</td>
<td>$9 \times 10^4$</td>
<td>90</td>
</tr>
<tr>
<td>CyIIa</td>
<td>160</td>
<td>$1.4 \times 10^4$</td>
<td>90</td>
</tr>
<tr>
<td>CyIIb</td>
<td>1000</td>
<td>$5.8 \times 10^4$</td>
<td>60</td>
</tr>
<tr>
<td>CyIIIa</td>
<td>470</td>
<td>$8.8 \times 10^4$</td>
<td>190</td>
</tr>
<tr>
<td>M</td>
<td>20–40</td>
<td>$2.5 \times 10^4$</td>
<td>650–1300</td>
</tr>
</tbody>
</table>

Data are from J.J. Lee, F.J. Calzone, R.J. Britten, R.C. Angerer, and E.H. Davidson (unpubl.); measurements refer to 65-hr pluteus-stage embryos.
toderm descends initially from one animal pole quadrant of the cleavage-stage blastomeres, and its differentiation thus involves the activation in this lineage of a whole battery of diverse genes. This battery may include many other genes in addition to the examples already known, i.e., Spec1, Spec2, CyIIa, and CyIIIB.

Another battery of genes active in the embryo of which diverse representatives are already in hand is that functional in the muscle precursor cells. In addition to the M actin genes (Fig. 2), we have isolated the muscle myosin heavy-chain gene, using a Drosophila probe kindly provided by Dr. Charles Rozek (Case Western University). A significant homology between a region of the Drosophila and the sea urchin gene sequences is of particular interest. Thus, it affords an opportunity to test experimentally the hypothesis that activation of this gene may be caused by interaction with maternal regulators, and hence the spicula matrix protein gene is of potential generality.

Figure 4. Quantities of CyIIa actin gene transcripts accumulated during embryonic development. Transcripts were measured by solution titration with excess single-strand complementary probes. Various quantities of whole-embryo RNA were reacted with a gene-specific RNA probe synthesized in vitro. The shape of the curve for the dashed line is derived from relative RNA gel blot data (Short et al. 1984) (open circles). Titration measurements are designated by closed circles.

Figure 5. Sequence homology between a region of the sea urchin and the Drosophila myosin heavy-chain genes. The region shown derives from a genomic S. purpuratus recombinant. Dots indicate identical nucleotides, and where changes result in coding differences the respective amino acids are indicated. In the sequence shown 23/32 amino acids are identical. Drosophila sequence kindly provided by Dr. Sanford Bernstein (pers. comm.). Data from S. Rose, M. Rosenberg, D. Chen, R. Britten, and E. Davidson (unpubl.).
be detected. By specification is meant the process by which the differentiative fate of the descendants of given blastomeres is first established (whether irreversibly or not is, for this argument, irrelevant). There are only a few "target" nuclei when specification takes place, while expression occurs after a number of divisions have intervened, and a meaningful number of properly situated blastomeres have been produced. Different kinds of mechanisms can be envisioned. For example, genes such as those described here might be activated secondarily by the products of pleiotropically active master regulator genes, which might be the only genes directly affected in the initial specification events. Alternatively, such upper-level hierarchy control genes might not be involved at all in the early development of this embryo, and immediate regulators for the lineage-specific structural genes could be stored in the egg or released by appropriate interblastomere contacts. Specification would then consist of the direct interaction of such regulators with cis-regulatory sequences of lineage-specific structural genes, such as those described in this article.

**Gene Transfer: Expression of Microinjected Cyllla Genes**

We recently described a method for introducing genes into the sea urchin egg by microinjection, and provided evidence that the exogenous DNA is replicated and stably retained during development (Flytzanis et al. 1985; McMahon et al. 1985). Unfertilized eggs are bound by electrostatic attraction to protamine-coated dishes, and several thousand linearized DNA molecules are injected into the cytoplasm of each. Sperm is added, and the eggs develop in situ until hatching, when they secrete an enzyme that dissolves the fertilization envelope, releasing the swimming blastulae. We have shown that within an hour of injection the exogenous DNA has formed random end-to-end concatenates, and during cleavage these replicate an average of 30- to 100-fold. Incorporation of the exogenous DNA into early blastomere nuclei apparently occurs with good efficiency, and the replicated sequences subsequently persist throughout the several weeks of larval life. In about 30% of larvae, the quantity of exogenous DNA continues to increase. Presumably in these larvae the DNA had originally been incorporated in nuclei of those cell lineages that remain mitotically active during larval growth. In several experiments (see Flytzanis et al. 1985), the genomic DNA of postmetamorphosis juveniles descendant from the injected eggs was examined, and 6-15% were found to bear integrated sequences. This fraction probably reflects the frequency with which the exogenous DNA is incorporated into cells that are ancestral to the imaginal rudiment from which the postmetamorphosis juvenile de-

*Figure 6. Localization of hyalin displayed by indirect immunofluorescence. Pluteus-stage (74-hr) embryos, fertilized eggs, and unfertilized eggs were fixed, embedded, and sectioned, then reacted with anti-hyalin rabbit antibody. Bound antibody was visualized with fluorescein isothiocyanate (FITC)-conjugated goat antirabbit IgG. (a) Unfertilized eggs. (b) Fertilized eggs. (c) Phase-contrast micrograph of 5-μm sections of pluteus-stage embryos. (d) Immunofluorescence from same pluteus-stage embryos as in c. (From J.J. Robinson, N. Ruiz-Bravo, H.M. Sucov, B.R. Hough-Evans, R.J. Britten, W.J. Lennarz, and E.H. Davidson, in prep.)*
The method described, and the developed embryos were derived from the DNA of their sperm when sexual maturity has been attained. Such a gene transfer system could thus be utilized to produce transgenic lines of sea urchins. In the experiments described here, however, we have focused on expression of the exogenous DNA in the embryos deriving immediately from the injected eggs.

Initial studies on the expression of DNA injected into sea urchin eggs were reported by McMahon et al. (1984). In this work a construct containing the regulatory elements of the gene coding for the 70-kD Drosophila heat shock protein fused to the bacterial structural gene for chloramphenicol acetyltransferase (CAT) (Di Nocera and Dawid 1983) was injected into eggs by the method described, and the developed embryos were subjected to heat stress. It was found that at 25°C, a temperature at which the endogenous heat shock response is elicited in this sea urchin species, CAT enzyme synthesis was induced in the host embryos. Since in Drosophila, from which the exogenous heat shock gene sequences derive, this gene is silent at 25°C, transcription of the injected construct must respond to the diffusible signals produced or activated in the heat-stressed sea urchin cells. A similar result has been obtained in other heterospecific studies on Drosophila heat shock gene expression (e.g., Corces et al. 1981). It follows that the exogenous DNA is present in the sea urchin embryo in an intracellular location, and in a form that permits regulated transcriptional expression.

We were thus encouraged to investigate the expression of sea urchin genes that in normal embryos display an easily identified ontogenic pattern of expression. An initial choice was the CyIIIa actin gene, which, as shown in Figures 3 and 4, is expressed in a spatially and temporally specific way. Figure 7 displays the structure of the in-frame fusion between the CyIIIa gene and the CAT gene that was utilized for the following experiments (C. Flytzanis, R. Britten, and E. Davidson, in prep.). The injected DNA contains several kilobases of upstream CyIIIa sequence, plus a large (2.2-kb) intron wholly included in the 5' leader of the CyIIIa transcript and coding sequence for only a few of the aminoterminus amino acids of the actin molecule. The remainder of the actin structural gene has been replaced by the CAT gene. Translation could start at either the actin or the CAT ATG codon. About 3000 molecules of this construct were injected per egg, the eggs were fertilized, and at various stages of development aliquots of 20–50 embryos were harvested and the CAT enzyme activity present in the embryos measured. Results from two such experiments are shown in Figure 8a and b. CAT activity appears essentially on the schedule expected for CyIIIa transcripts (cf. Fig. 4). Thus, by 20 hours the amount of CAT enzyme has attained maximum value. Comparison to an absolute standard for CAT enzyme activity (McMahon et al. 1984) suggests that about 10^9 molecules of CAT enzyme are produced by the CyIIIa-CAT genes per embryo, on the average, and assuming the usual translational parameters (see Davidson 1976) it may be roughly estimated that in the 20-hour embryo the quantity of fusion gene mRNA is on the order of about one-fourth of the natural quantity of CyIIIa message. An important additional observation is shown in Figure 8c. Here is presented a parallel series of experiments in which the upstream regulatory sequences of the S. purpuratus early H2a histone gene (for review of sea urchin histone gene regulation see Hentschel and Birnstiel 1981) have replaced those of the CyIIIa gene in a similar CAT

---

**Figure 7.** CyIIIa-CAT fusion constructs. (T.S.) Transcribed sequence; (U.S.) upstream sequence. A diagram of the linearized construct as injected into sea urchin eggs is shown in the top line. The plasmid was linearized at an SphI site 2.5 kb upstream of the 5' end of the CyIIIa transcript. Since the molecules form a random end-to-end concatenate after injection (McMahon et al. 1985), in half the cases the original 8-kb upstream sequence included in the plasmid will be regenerated. In the second line is a diagram of the CyIIIa gene, from data of R. J. Akhurst et al. (in prep.). Introns are denoted as thin lines and exons are numbered. The third and fourth lines show the sequence of the junction region (shown by the arrow in the second line) between the CyIIIa and the bacterial CAT genes, with the two in-frame ATG translation start codons, the first deriving from the CyIIIa and second from the CAT gene.
Figure 8. CAT expression from fusion genes injected into sea urchin eggs. The genes were injected into unfertilized eggs which were then allowed to develop until collected for assay of CAT activity, as described by McMahon et al. (1984, 1985; see text). (a and b) Two independent experiments using the CyIIIa-CAT fusion gene depicted in Fig. 7. (C) Control assay containing bacterial CAT enzyme. The arrow indicates the monoacetylated CAT product. Other samples contained sea urchin embryo extracts. (UN) Embryos derived from uninjected eggs; (I) embryos derived from injected eggs. The number of hours postfertilization and the number of embryos in the sample analyzed are indicated, respectively, in the bottom two rows of numerals. (c) CAT activity in embryos injected with an α-H2a-CAT construct. This contained all sequence upstream of the H2a gene to the preceding H3 histone gene, but only 30 nucleotides of coding sequence.

In embryos injected with this construct the CAT enzyme appears at 15 hours, just after the amount of endogenous H2a mRNA reaches its peak value in normal embryos (Mauron et al. 1982).

In preliminary experiments so far available, the CAT activity then disappears, just as does the endogenous α-histone message. This result would imply that within the sea urchin embryo the newly synthesized CAT protein is unstable, and also that the mRNA produced by the fusion gene is unstable. The same conclusion follows from the constant amount of CAT activity observed after the blastula stage in embryos injected with the CyIIIa-CAT fusion. Most significantly for the results shown in Figure 8a and b, it is clear from Figure 8c that the absence of CyIIIa-CAT expression at 15 hours cannot be due to insufficiency of exogenous DNA or inaccessibility of this DNA for transcriptional activation in 15-hour (as opposed to 20-hour) embryos. Some time is required for accumulation of the CAT protein once the message has appeared, and it must also be taken into consideration that injected embryos are delayed in their development by about one division cycle (McMahon et al. 1985). These factors account for the minor retardation, in regard to the respective endogenous transcripts in control embryos, in the appearance of CAT enzyme in both the CyIIIa-CAT and H2a-CAT experiments shown. We may conclude that both the exogenous CyIIIa gene sequences and the exogenous α-H2a sequences promote expression of their fusion constructs according to the predicted ontogenic schedule.

It should thus be possible to explore directly the mechanism by which lineage-specific genes are activated in the early sea urchin embryo. This is not necessarily the same problem as determining what activates genes in the terminal differentiation processes of later development, or in the physiologically induced genes of adult organisms. The early embryo utilizes spatial regulatory information that originates maternally; it relies to some extent on oriented interblastomere interactions for lineage determination; and, unlike the case in advanced developmental systems, its nuclei are equivalent and totipotent, i.e., until the initial events of lineage specification have taken place. Furthermore, the results summarized by Solter et al. elsewhere in this volume show that some genes destined to function early in mouse development may be "imprinted" at the chromatin level during gametogenesis. The qualitative results illustrated in Figure 8 are already instructive in this respect, however, in that they suggest that genes activated in the early sea urchin embryo respond to trans-activators that are at least initially present in excess. At the level of genomic function, the key to the mechanisms underlying the initial specification of embryonic cell lineages may lie in the nature, origins, and spatial disposition or release of such trans-acting embryonic regulatory molecules.

ACKNOWLEDGMENTS

This work was supported by National Institutes of Health grants HD-05753 and GM-20927. C.N.F. was...
supported by a Lieve Fellowship (S-11-83) from the American Cancer Society, California Division; J.I.R. by a Proctor & Gamble Fellowship; S.J.R. by a Muscular Dystrophy Association Fellowship; and H.M.S. and J.J.L. by a National Institutes of Health training grant (GM-07616).

REFERENCES


HERBST, C. 1900. Uber das Auseinandergehen von Forchhurs-
Lineage-specific Gene Expression in the Sea Urchin Embryo


Access the most recent version at doi:10.1101/SQB.1985.050.01.041

References
This article cites 28 articles, 12 of which can be accessed free at:
http://symposium.cshlp.org/content/50/321.refs.html

Article cited in:
http://symposium.cshlp.org/content/50/321#related-urls

Email alerting service
Receive free email alerts when new articles cite this article - sign up in the box at the top right corner of the article or click here

To subscribe to Cold Spring Harbor Symposia on Quantitative Biology go to:
http://symposium.cshlp.org/subscriptions

Copyright © 1985 Cold Spring Harbor Laboratory Press