Abstract. Although a major site of transcription in heat shock, the Drosophila hsrω gene does not encode any known heat shock proteins. Instead, studies of the hsrω transcripts suggest that the RNA molecules, rather than encoded proteins, are the active products of this gene. The cytoplasmic RNA, ω3, is spliced and polyadenylated and yet has only very small open reading frames (ORFs), and these are poorly conserved in different Drosophila species. Surprisingly, the work reported here leads us to conclude that one of the tiny ORFs in this RNA is translated. This ORF, designated ORF-ω, is notable in being the only ORF that shows sequence conservation in the three Drosophila species examined. However, translation of this ORF does not lead to detectable accumulation of the protein product. We suggest that ORF-ω may be an example of an unusual type of translated ORF. The act of translation itself may be important rather than the generation of a functional protein product. This nonproductive translation may play a role in regulation of cellular activities.

Activation of genes by temperature shifts, the “heat shock response,” was first identified in Drosophila by the cytological techniques that allowed visualization of gene activity as specific puffs on polytene chromosomes (Ritossa, 1962). As biochemical methods became available, the heat shock response was found to be inducible in cells of all tissues and to be common to all organisms, from bacteria to plants and animals (for review, see Craig, 1985; Lindquist, 1986). Furthermore, the set of heat shock genes was also found to be induced by a number of other stimuli that, like a heat shock, could be classified as cellular stress agents. Studies on many organisms have now suggested that the response is a general cellular homeostatic mechanism. Heat shock–induced proteins have amino acid sequences that have been extremely conserved during evolution. The evolutionary conservation of the heat shock response argues for its importance to cells of all types, yet little is known about the function of the heat shock proteins.

The first five Drosophila melanogaster heat shock loci to be cloned encoded all of the major heat shock proteins, raising questions about the role of the last major heat shock–induced locus, in polytene region 93D (Craig, 1985). In addition to its lack of identification with a protein product, the 93D puff has other unusual properties (for review, see Lakhotia, 1987; Bendena et al., 1989a). A major portion of the transcripts from this gene are found in the nucleus (Lengyel et al., 1980). In polytene chromosomes, the 93D puff contains large RNP granules not seen in other puffs (Dangl et al., 1983). The 93D puff is induced independently of the rest of the heat shock set with agents such as benzamide and colchicine, suggesting that its activity is more sensitive to other stimuli than the rest of the heat shock genes (Lakhotia and Singh, 1982).

Recently, the 93D gene has been cloned from D. melanogaster, as have the equivalent genes from D. hydei (2-48B) and D. pseudoobscura (58C) (Walldorf et al., 1984; Garbe and Pardue, 1986; Ryseck et al., 1987; Garbe et al., 1986; Garbe et al., 1989). The DNA sequences of these three loci are highly diverged, and yet the loci share a unique molecular structure and have small regions of conserved sequence at several places in the transcription unit. In addition, in each species, this is a major heat shock puff, and it is also induced by the other agents that induce 93D (e.g., benzamide and colchicine), binds antibodies that bind 93D uniquely, and contains large RNP granules (for review, see Lakhota, 1987). Each of the loci is the single locus in the species that shares these phenotypic and molecular characteristics, giving strong evidence that 93D, 2-48B, and 58C are homologous. Because of this homology each of the loci has been named the hsrω locus for its species. (Although it is now known that the hsrω locus is constitutively expressed, the name hsrω reflects its original identification as the locus encoding the ω set of heat shock RNAs.)

The hsrω gene, like the gene for hsp82, is a constitutive gene whose expression is elevated, rather than newly induced, by heat shock. In both normal and stressed cells, the locus produces three transcripts, all from the same start site.

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The largest transcript of hsrω, ω3, which encompasses the entire transcription unit of ~10 kb, is marked by some 7–8 kb of a short tandem repeat at the 3’ end. This is the transcript that accumulates to very high levels in the nucleus. A smaller nuclear transcript (~1.9 kb), ω2, appears to be the result of alternate termination at a polyadenylation site just upstream of the tandem repeats. The 1.9-kb RNA species is not as abundant as the ω1 transcript and apparently acts as the splicing precursor to an abundant cytoplasmic RNA, ω3. The ω3 transcript is 1.2–1.3 kb (depending on the Drosophila species) and is formed from ω2 by removal of a single intron (Garbe et al., 1986).

The cytoplasmic RNA, ω3, has unusual properties. All previously known RNAs of this size that are spliced and polyadenylated act as messages and encode proteins. Surprisingly, the sequences of the hsrω loci from D. melanogaster, D. hydei, and D. pseudoobscura do not show any long open reading frames (ORFs). In fact, the short ORFs in this transcript are all within the range that might occur by chance alone and are shorter than some of the ORFs that are found in the nontranscribed strand of the hsrω sequence. Furthermore, comparison of the sequences from the three Drosophila species show no conservation of either the positions or the amino acid sequences of all but one of these ORFs. The single exception would yield only a small polypeptide of 23–27 amino acids, depending on the Drosophila species (Garbe et al., 1986; Garbe et al., 1989). Therefore, it appears likely that the cytoplasmic RNA, ω3, does not serve simply to provide the information for a protein but, instead, has some other function in and of itself.

To look for clues to the function of hsrω, we investigated the properties of the hsrω cytoplasmic RNA in cultured cells. Contrary to expectation, these studies, as reported here, have led us to conclude that the ω3 RNA has an ORF that is translated. The ORF, which is translated and now called ORF-ω, is the only one showing sequence conservation in the three Drosophila species examined. However, translation of this ORF does not lead to detectable accumulation of the protein product. We suggest that ORF-ω may be an example of an unusual type of translated ORF. The act of translation itself, rather than the generation of a functional protein product, may be of importance in this case. Other studies have shown that ω3 transcripts turn over rapidly in control cells but accumulate to high levels whenever protein synthesis is inhibited (Bendena et al., 1989b). We suggest that nonproductive translation of ORF-ω allows the level of ω3 to reflect the level of protein synthesis in the cell and thus provides information that is used in coordinating intracellular activities.

Materials and Methods

Cell Growth and Preparation of Cell Lysates

D. melanogaster cells of the Schneider line 2-L (Storti et al., 1980) were grown in DME supplemented with 10% FCS, 0.5% lactalbumin hydrolysate, and MEM nonessential amino acids. Schneider line 3 (S3) cells (D. melanogaster) (Schneider and Blumenthal, 1978) and D. hydei cells of the KUN-DH-15 line (Sondheimer et al., 1980) were grown in Schneider’s medium supplemented with 10% FCS, 50 U/ml penicillin, and 50 µg/ml streptomycin. (All cell growth materials from Gibco Laboratories, Grand Island, NY) Melanogaster line 2-L cells were maintained in spinner culture at 3–15 × 10^6 cells/ml by dilution every 1–3 d. D. melanogaster S3 cells and hydei cells were maintained at similar levels by 5–10-fold dilutions every 3–7 d. For use in an experiment, line 2-L cells were maintained at 3–6 × 10^6 cells/ml by daily dilution for at least 3 d. Line 3 cells and hydei cells were diluted 5–10-fold, allowed to grow to 6 × 10^6 cells/ml, or 1.5 × 10^7, respectively, then diluted a second time, and used when they had grown to 3–6 × 10^6/ml or 1–1.5 × 10^7, respectively (mid to late log phase of growth).

Sucrose Gradient Sedimentation

A cytoplasmic fraction of cell lysates was prepared and analyzed on 15–50% sucrose gradients exactly as described by Ballinger and Pardue (1983), except that lysates were prepared using 0.1% Triton to prevent nuclear leakage and lysis. However, high salt conditions (0.5 M KCl) were maintained to minimize nonspecific molecular associations and to dissociate “vacant couples” (monosomes not associated with mRNA). Identical conditions were used for preparation and centrifugation of cell lysates on 5–20% sucrose gradients. For each gradient, a cell lysate prepared from 1–3 × 10^6 cells of the 2-L line or 0.5–1 × 10^6 of line 3 cells or D. hydei cells was used for a gradient of 27 ml volume. In some experiments, cleared cell lysates were treated for 15 min with 25 mM EDTA on ice before sedimentation on sucrose gradients supplemented with 10 mM EDTA. For studies requiring inhibition of translation, cells were treated at 25°C with cycloheximide at 10^{-4} M for 2 h (Lodish, 1971), with pactamycin at 10^{-7} M for 2 h (Stewart-Blair et al., 1971), or with 120 µg/ml 1-t-lysylamino-2-phenylethyl chloromethyl ketone (Pong et al., 1975) for 15 min at 25°C before preparation of cell lysates.

Metrizamide Gradient Sedimentation

Cell lysates of Schneider line 2-L cells were analyzed by equilibrium gradient centrifugation on metrizamide exactly as described by Ballinger and Pardue (1983). Gradients were 5 ml in volume and 20% of the amount of cell lysate was loaded on metrizamide gradients as compared to the amount used for sucrose gradients.

Cell Labeling and Protein Gel Electrophoresis

One milliliter of cells was concentrated into 25 µl of Shields and Sang salts (Schneider and Blumenthal, 1978) and 2 µl of 10^{-4} M 35S-methionine (15 µCi/µl, sp act, 800 Ci/mM) for 10 or 30 min. In some experiments, cells were lysed in 100 µl of lysis buffer (100 mM Tris, pH 6.8, 1% SDS) and boiled immediately for 5 min. Proteins were separated from free amino acids by precipitation in 10 vol of acetone. Proteins were then dissolved in sample buffer containing 0.1% SDS and boiled for 3 min with or without the addition of β-mercaptoethanol to 5%. In other experiments, cells were lysed in a sample buffer containing 1% SDS and loaded on gels directly after boiling for 3 min with or without β-mercaptoethanol added to 5%. Samples were electrophoresed on acrylamide gels containing 7.5 M urea, 0.1% SDS (Garbe and Pardue, 1986) or on standard Laemmli gels (Laemmli, 1970).

Cloning Vector

A derivative of pSVOSna-CAT obtained from M. Gilman (Massachusetts Institute of Technology), was used as the cloning vector for assay of hsrω gene fragments. This vector was constructed by replacement of the pBR322 polylinker directly 5' to the bacterial gene for chloramphenicol acetyltransferase (CAT). About 150 bases around the polylinker of this vector were replaced for assay of transient CAT expression.

Establishment of Transformed Cell Lines and CAT Assay

S3 cells were transformed with construct DNA and heterogeneous cell lines were established as described (Bourouis and Jarry, 1983). Transformed cell lines were sufficiently amplified for use in an experiment within 4–5 wk.

The level of CAT activity was analyzed in cell extracts prepared as described (Gorman, 1986), but extracts from these established, transformed cell lines needed to be diluted ~100-fold over amounts required for assay of transient CAT expression.
Figure 1. Size of cytoplasmic particles containing ω3 RNA as determined by sedimentation gradient centrifugation. In both heat-shocked and control cells the ω3 transcript is found in the monosome and disome fractions. (A) Northern analysis of a 15-50% sucrose gradient probed with cDNA for (a) ω3 RNA, (b) hsp82 RNA, and (c) rp49 RNA. Lysates were prepared from cells heat shocked for 1 h at 36°C. The gradient profile of light absorbance at 254 nm is drawn above the Northern blots and the portions of the gradient from which each fraction was drawn are indicated. Fractions 4-10 were scanned at four times increased sensitivity. The gradient profile is typical of that from lysates of heat-shocked cells in which a majority of the polysomes have broken down as the cells stop translating many non-heat shock RNAs. The very small monosome peak reflects the fact that, under the high salt and triton conditions used to prepare these lysates, vacant couples (complete ribosomes that are not actually translating on mRNA) are dissociated into their subunit components. The UV absorbance peaks as well as gel analysis indicated that fraction 1 contains free cell proteins; fraction 2 contains free 40S ribosomal subunits; fraction 3 contains free 60S ribosomal subunits and translating monosomes; fraction 4 contains translating disomes; and the rest of the fractions contain translating polysomes of increasing size. The positions of additional mRNAs in the gradients were determined to help assess the accuracy of our determination of polysome localization. The hsp82 mRNA is on larger polysomes while the ribosomal protein mRNAs, rp49, which is not translated under these conditions (Ballinger and Pardue, 1983), is not in the polysome region. (B) Northern analysis of a 5-20% sucrose gradient probed with cDNA for ω3 RNA. Lysates were prepared from cells grown at 25°C. The gradient profile of light absorbance at 280 nm is drawn above the Northern blot and shows the ribosomal contents of each gradient fraction. (Fractions 6-10 were scanned at four times increased sensitivity.) Peaks in fractions 1 and 2 correspond to free cell proteins; fraction 3 contains free 40S subunits; fraction 4 contains free 60S subunits; fractions 5 and 6 contain translating monosomes; fractions 7 and 8 contain translating disomes; and fraction 9 contains translating trisomes. The localization of histone mRNA helped confirm these fraction assignments (data not shown).
Protein Gel Blotting and Antibody Probing

A crude protein preparation was made from the pelleted material obtained from 1 ml of tissue culture cells by lysis in 100 μl of Laemmli sample buffer. β-mercaptoethanol was then added to some samples to 5%, the samples were boiled for 5 min, and 25 μl of the lysate was loaded onto a single lane of a standard 15% acrylamide gel. Gels were electroblotted to nitrocellulose (electroblot) and probed with a rabbit antibody against bacterial CAT (5 Prime → 3 Prime, Westchester, PA) at an antibody dilution of 1:50,000.

Results

Localization of hsrω Cytoplasmic RNP, ω3, to a Distinct Cytoplasmic Fraction

The ORFs in the hsrω transcript would encode only small polypeptides (Garbe et al., 1986; 1989). Searches for proteins of the appropriate size, in cells containing abundant hsrω transcripts, have been unsuccessful (Garbe and Pardue, 1986; Lakhota, 1987; see below). These results support the hypothesis, discussed above, that the cytoplasmic transcript of hsrω acts as a functional molecule in itself. To provide clues to possible functions, the hsrω cytoplasmic RNA, ω3, was examined in its native RNP form. Tissue culture cells of the D. melanogaster line Schneider 2-L or of the Drosophila hydei cell line, KUN-DH-15, were lysed by treatment with 0.1% Triton X-100 in the presence of an RNase inhibitor and fractionated by differential centrifugation. These initial crude cell fractionation experiments demonstrated that much of the ω3 RNA was located in the postmitochondrial supernatant. To examine the size of native ω3 RNA-containing particles, a postmitochondrial supernatant was prepared and sedimented on 15–50% or 5–20% sucrose gradients. Gradients were collected in 10 fractions and the position of ω3 RNA in the gradient was determined by Northern blotting of total RNA extracted from each fraction. A tracing of the UV absorbance profile, showing ribosome sedimentation, (the predominant particle in these lysates), was used to obtain the approximate sedimentation coefficient of each fraction. Fig. 1 shows an example of one such experiment. On 15–50% sucrose gradients (Fig. 1 A), the ω3 transcript exhibited a wide distribution across the top of the gradient appearing strongly in the second, third, and fourth fractions. This distribution indicated sedimentation with particles between 40S and 120S. Expansion of the area between these sedimentation values by sedimentation of cell lysates on 5–20% gradients (Fig. 1 B) showed ω3 RNA distributed most strongly in particles of a size between 80S and 120S, precisely in the fractions corresponding with single complete ribosomes (monosomes) and disomes. Identical results were obtained whether lysates were from cells grown at ambient temperatures or cells that had been heat shocked for 1 h at 36°C. Also, no differences were seen in sedimentation behavior between the cell lines of the two Drosophila species examined.

The position of ω3 sedimentation is that expected if this RNA were being translated on polysomes of monosome and disome size, consistent with the small size of ω3 ORFs. Because translation of ω3 had not been expected, a comparison of the gradient positions of several other translatable RNAs was made to ensure the accuracy of our polysomal localization. The comparisons were accomplished by stripping the Northern blot of ω3 probe and rehybridizing with additional probes. An example of this type of experiment is shown in Fig. 1 A. For the experiment shown, the cells were heat shocked before lysis. The RNA for hsp82, which is translated under heat shock conditions, was distributed across the gradient beginning at about 40S, but was most prevalent in fractions containing larger polysomes, consistent with the fairly sizable ORF of this message. Conversely, the RNA for the ribosomal protein, rp49, is found only in the first two fractions, containing only particles smaller than monosomes. This is the result expected since rp49 RNA is not translated during heat shock and has been shown to leave the polysomes completely (Ballinger and Pardue, 1983). Therefore, the conclusion that particles containing ω3 RNA colocalize with monosomes and disomes, that had been based on the identification of polysome peaks by UV light absorbance, is supported by localization of other RNAs known to be associated with polysomes.

The hsrω Cytoplasmic Transcript ω3 Is Associated with Polysomes of a Size Consistent with Translation of Its Small ORF

Polysomes of monosome and disome size are fully consistent with those expected for translation of the small ORFs found in the ω3 transcript, and thus ω3 RNP could colocalize with these particles because it is translated. However, untranslated, free cytoplasmic RNP particles of a size as large as monosomes and disomes have been described (McCarthy et al., 1983). To determine whether the ω3 cytoplasmic RNA was actually associated with polysomes or in a free RNP particle, we have tested other attributes of bona fide polysomes. The detergent sodium deoxycholate, which is known to disrupt the structure of some free RNPs, but not polysomes (Faiferman et al., 1971), did not disturb the colocalization of ω3 RNA and monosomes/disomes in the Schneider 2-L cells. In addition, increasing monovalent cation concentration (K+ or Na+) in the gradient over a broad range of 10–500 mM (which would be expected to strip loosely associated proteins from RNPs differentially according to their individual structures) had no effect on this colocalization (data not shown).

Density gradient centrifugation fractionates RNPs on the basis of their protein/RNA ratio, while sedimentation gradient centrifugation fractionates on the basis of molecular size. Therefore, cell lysates were banded on metrizamide density gradients as an alternate method of testing the colocalization of ω3 RNA-containing particles with translating polysomes. As shown in Fig. 2, the ω3 cytoplasmic RNP was found in the region of the gradient that contained polysomes. Lysates of Schneider 2-L cells that had been heat shocked at 36°C for 1 h were used for the experiment shown; the ω3 RNA had the same density when obtained from cells held at ambient temperature. An example of a translating RNA, coding for hsp82, was found near the bottom of the gradient (fraction 3), consistent with polysomal localization (Ballinger and Pardue, 1983). In contrast, the RNA coding for the ribosomal protein, rp49, which becomes released from polysomes during heat shock (Ballinger and Pardue, 1983), was found distributed much more broadly in the gradient, at densities that might be expected for free RNPs (McCarthy et al., 1983). The ω3 RNA was found near the bottom of the gradient in the same fractions as hsp82 RNA, suggesting again that ω3 is on polysomes.

Another test for the polysomal association of ω3 is the
Figure 2. Density of cytoplasmic particles containing ω3 RNA. The ω3 RNP has the buoyant density of a polysome. Northern analysis of fractions from metrizamide density gradients probed with cDNA for (A) ω3 RNA, (B) hsp82 RNA, and (C) rp49 RNA. Lysates were prepared from cells heat shocked for 1 h at 36°C. The density of gradient fractions is indicated below the gel lane. Fraction 1 is from the bottom of the gradient. The ω3 RNA and the hsp82 RNA band at the density of polysomes. The rp49 RNA, most of which sediments as free RNP in sucrose (Fig. 1), is found at densities appropriate for free RNP.

If ω3 RNA is associated with polysomes, then, when cell lysates are treated with EDTA before sedimentation on sucrose gradients, the position of ω3 RNA should show a shift out of the polysome region of sucrose gradients to the free RNP region at the top of the gradient. In these experiments, ω3 RNA from untreated lysates colocalized with monosomes and disomes as expected; after EDTA treatment, it shifted to the free RNP region. To demonstrate complete polysome disruption with EDTA, the gradients were probed for several mRNAs. Fig. 3 shows results for ω3, hsp 82 mRNA, and histone mRNA (a message that codes for a small protein and thus is found on small polysomes). The ω3 cytoplasmic RNP behaved the same way as the known messenger RNAs: it shifted to the top of the gradient with EDTA treatment, suggesting that it indeed is associated with polysomes (although EDTA can also dissociate other types of RNPs).

Another way to test for polysome association is to determine the localization of the RNA in cells treated with either an inhibitor of translation elongation or an inhibitor of translational initiation. A typical experiment is shown in Fig. 4. Polysome disruption was monitored, as before, in two ways: by the UV absorbance profile generated from sucrose gradients during collection, and by analyzing the shift in gradient position of RNAs known to be translating on polysomes under normal conditions. Sucrose gradients of 5–20% were used to allow visualization of polysomes containing up to three ribosomes.

For elongation inhibition, cells were treated with cycloheximide (Lodish, 1971), at 10^{-4} M for 2 h at 25°C, a treatment that we have found blocks up to 95% of protein synthesis. Cycloheximide treatment freezes polysomes by preventing the continuation of translations that had already begun, but it does not disrupt polysomes. Fig. 4 shows that, after cells have been treated with cycloheximide, ω3 RNA continues to colocalize with polysomes in much the same way as seen in untreated cells. This distribution is similar to that of representative, translating mRNAs, the histone transcripts.

In contrast to the effects of elongation inhibitors, inhibitors of initiation will cause mRNA to accumulate in the mono-
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The region of the polysomes after cycloheximide treatment, which pactamycin at 10^-7 M for 2 h at 25°C. Both RNAs accumulate in a single gradient made from a cell lysate that had been treated with cycloheximide at 10^-4 M for 2 h at 25°C. C and D represent RNA from a single gradient made from a cell lysate that had been treated with pactamycin, at 10^-7 M for 2 h at 25°C. Both RNAs accumulate in the region of the polysomes after pactamycin treatment, which disrupts translation. Pactamycin inhibits initiation, thus disrupting those fractions, as expected of RNAs that are being translated.

The Conserved ORF of hsro·, ORF-ω, Is Translated

Collectively, our data leads to the conclusion that the ω3 RNA is associated with polysomes. Furthermore, in both D. melanogaster and D. hydei, this RNA associates only with monosomes and disomes, consistent with translation of one of the small ORFs in the ω3 sequence. Comparison of the ω3 sequences in these two species show only one ORF that shows any conservation in sequence or position (Fig. 5). It is important to note that this ORF, which we now call ORF-ω, is the first ORF in either species that is in a sequence context of the type that has been shown to be necessary for efficient initiation, at least for mammalian cells (Kozak, 1986). More recently the sequence of ω3 from D. pseudoobscura has shown that, in this species also, ORF-ω is the only ORF showing any sequence conservation (Garbe et al., 1989). Even though the sequence conservation does not extend through the entire ORF, the absence of other conserved ORFs make ORF-ω a strong candidate for translation.

Synthetic gene constructs were prepared to test whether ORF-ω is the ORF that is translated in vivo. Two recombinant genes were constructed for the initial experiments (Fig. 5). ORF-ω-CAT contains the entire hsro· promoter with its heat shock–inducible element followed by the transcribed portion of the gene to a point 30 bases past the end of ORF-ω (base 229 of the hsro· sequence). This base is linked directly to the bacterial gene for CAT, which has its own translational start. The transcript from this construct contains 5' to 3': a leader sequence, a nine codon ORF in an unfavorable context, the complete ORF-ω, and then the CAT coding region. A second construct, ω-leader-CAT, was made by generating a deletion in ORF-ω-CAT that completely removes ORF-ω but leaves the sequence 5' to ORF-ω, including the first, unfavorable ORF, intact. Each construct was introduced into D. melanogaster S3 cells, along with the bacterial gene for dihydrofolate reductase which allows selection of transformed cells with the drug, methotrexate. Both of the resulting, drug-resistant cell lines were shown by Southern blotting to contain an average of 10 copies/haploid genome of the transforming construct in unrearranged form.

Since both constructs have the hsro· promoter and its 5' transcribed sequences, both would be expected to confer on the transcript the ability to be induced and translated during...
The ORF-ω-CAT construct was made by ligation of a 1.5-kb Xho I fragment from the cloned hsrw gene of D. melanogaster into the Sal I site of the polylinker of the vector, p06 (see Materials and Methods). This directly apposes the D. melanogaster DNA to the gene for CAT. The Drosophila Xho fragment contains, in 5' to 3' order: the 5' flanking portion of the hsrw gene with the promoter sequences for gene expression; the start site for transcription of the hsrw gene (base 1 of the sequence); a short ORF (beginning at base 9 of the sequence and not in an optimal context for translation according to the Kozak theory); ORF-ω with its translation start and stop codons (extending between bases 118 and 201 of the sequence and colored in black on the diagram); and 29 additional bases of sequence. The CAT gene translational leader is colored in gray on the diagram and the start codon for CAT translation is indicated. ω-leader-CAT is a deletion of ORF-ω-CAT from the Sma I site in the polylinker (just 3' to the Xho I site) to the Rsa I site at base 67 of the hsrw gene fragment. This deletion removes ORF-ω, but leaves the first ORF. The sequence of the resulting transcripts from these constructs up to the beginning of the portion of the construct contributed by the CAT gene is shown below the diagrams. The start site for transcription of the hsrw gene of D. melanogaster has been previously determined (Garbe et al., 1986; Ryseck et al., 1987). (B) The ORF-ω/CAT fusion construct was made by ligation of a 1.2-kb Eco RI/Sal I fragment of the hsrw gene from D. hydei into the Sma I site of the polylinker in the vector, p06. The Drosophila fragment was blunt-ended by a fill-in reaction using T4 DNA polymerase. The Drosophila fragment contains 5' to 3': the 5' flanking portion of the hsrw gene with the promoter sequences for gene expression; the start site for transcription of the hsrw gene (base 1 of the sequence previously determined for the D. hydei gene [Garbe et al., 1986; Ryseck et al., 1987]); and the majority of ORF-ω (21 out of 23 heat shock. The ω-leader-CAT construct should further allow the translation of the CAT polypeptide since the CAT coding region is the first ORF that begins in a favorable context for translation. According to our hypothesis, the first tiny ORF should not be translated because of the unfavorable sequence context. In contrast, if ORF-ω were translated, the mRNA produced from the ORF-ω-CAT construct should begin translation in ORF-ω and terminate at its stop codon. In this case, ribosomes should never reach the CAT start codon to translate CAT protein. An assay of CAT activity in the two transformed cell lines supports these predictions (Fig. 6). A moderate amount of CAT activity was seen in the cell line S3-ω-leader-CAT at ambient temperature (as expected because hsrw is also a constitutive gene). A significant increase in the activity was induced by heat shock for 1 h at 36°C. In contrast, in the cell line S3-ORF-ω-CAT, only a very low level of CAT activity was found at any temperature. Examination of the level of CAT protein by gel electrophoresis provided further confirmation of the results of enzymatic activity assay. The cells carrying the ORF-ω construct indeed had very little CAT protein (data not shown). The low level of CAT protein in the S3-ORF-ω-CAT cell line was not because of a low expression of CAT RNA. In fact, Northern blotting showed that the steady-state level of ORF-ω CAT mRNA was 7.3 times greater than that found in the line S3-ω-leader-CAT at 25°C. RNA from the S3-ORF-ω-CAT cell line was found to be induced 8.4-fold at 33°C and 12.6-fold at 36°C over constitutive amounts compared to 10.3-fold at 33°C and 16.2-fold at 36°C for the RNA from S3-ω-leader-CAT cells. Thus, in spite of the consistently greater amounts of the transcript carrying the ORF-ω, very little CAT protein was detected at any temperature in cells with the ORF-ω-CAT. These results are consistent with the hypothesis that ORF-ω is translated.

The ORF-ω just in front of the CAT gene should impede CAT translation because the translation of ORF-ω itself should inhibit reinitiation on the CAT sequence. (The results above could also be obtained if the placement of ORF-ω adjacent to the CAT gene causes some other problem for translation, possibly because of structural considerations such as hairpin formation.) If translation of the ORF-ω were inhibiting translation of the CAT gene, we would expect to find the ORF-ω-CAT RNA on monosomes and disomes and the ω-leader-CAT (the construct lacking ORF-ω), on polysomes of the size appropriate for the 24.1-kD CAT protein. Fig. 7 shows that this was the result obtained. In cells of the S3-ORF-ω-CAT line, ORF-ω-CAT RNA was found on polysomes of monosome and disome size, while polysomes translating the ω-leader CAT RNA were slightly larger, as predicted by codons), including its translational start codon but missing its last two amino acid codons and its translational stop codon. Base 187 of the sequence marks the end of the Drosophila DNA and base 188 is the beginning of the CAT leader (italicized in the sequence below the diagram), and is followed by the translational start codon for CAT (base 225). Translation that begins at ORF-ω should continue through the CAT leader (italicized amino acid sequence) and arrive at the CAT translational start in frame for continued translation through the CAT gene. It is noteworthy that there are no methionine codons in any reading frame in the CAT leader, so that any larger CAT protein should start within the hsrw sequence.
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Figure 6. Assay of CAT activity in cell lines transformed with hsrw-CAT constructs. Thin layer chromatogram demonstrating the level of CAT activity found in equivalent amounts of extract from cell lines carrying different CAT constructs. The origin of migration is at the bottom of the photo; the autoradiographic densities in order of increasing distance from the origin represent [14C]chloramphenicol substrate followed by the acetylated products. (Lanes 1, 3, and 5) CAT activity at 25°C. (Lanes 2, 4, and 6) CAT activity at 36°C. (Lanes 1 and 2) CAT activity in the cell line S3-0-leader-CAT. (Lanes 3 and 4) CAT activity in the cell line S3-ORF-0-CAT. (Lanes 5 and 6) CAT activity in the cell line S3-hsp22-leader-CAT. This last construct was made by linkage of the promoter/leader of the D. melanogaster gene for hsp22 to the CAT gene in the vector, p106 (Bendena, unpublished data). Cell lines carrying both the 0-leader-CAT and the hsp-22-leader-CAT show significant CAT activity at 25°C, and this activity is increased after heat shock induction of the CAT transcript. Cells carrying the ORF-0-CAT construct show very little CAT activity at any temperature.

Translation of the Cytoplasmic RNA o3 Cannot Be Linked to a Small Protein Product

The evidence presented above for translation of ORF-0 seems to contradict the failure of previous work to link a protein product with the hsrw gene (Garbe and Pardue, 1986; Lakhota, 1987). Therefore, an additional effort was made to search for the translation product of ORF-0 by examining small cellular proteins under diverse conditions of stress (when o3 is abundant). Conceptual translation of the D. melanogaster ORF-0 yields a protein product of ~3 kD containing two methionine residues. Fig. 8 shows an example of our experiments to search for such a protein. The autoradiogram of the 10% polyacrylamide gel in A shows the spectrum of the peptides synthesized in each culture. The autoradiogram of the 20% gel shown in B displays the same samples in a way that maximizes resolution of the small peptides. No protein of the size predicted from the ORF-0 sequence was visualized after any of the cell treatments, even when the gel is exposed for times much longer than required to easily demonstrate the 20-kD family of heat shock proteins. In contrast, another small protein, ubiquitin, which is the product of a minor heat shock puff (Arribas et al., 1986), can be clearly visualized on these gels. Ubiquitin migrates on these gels with the 6-kD size standard and was identified by immunoprecipitation. A shorter labeling time (10 min) as well as a different label (complete 3H-labeled amino acids) was tried, but in no case was there a detectable heat shock synthesis of any small polypeptide other than ubiquitin. These results suggest that, if ORF-0 is translated, the protein is either turning over rapidly or being sequestered.

If the product of ORF-0 is turned over very rapidly, it might still be possible to visualize the resulting peptide by examining labeled proteins from a cell line that is overexpressing RNA containing ORF-0. The cell line S3-ORF-0-CAT synthesizes at least ten times more transcripts containing ORF-0 (in the construct that also has the CAT gene) than do untransformed cells. The additional transcripts are on monosomes and disomes and apparently translated, and yet, even in these cells, no new proteins of the appropriate size were found.

Visualization of the hsrw Translation Product as Part of a Fusion Protein

Our inability to find a small protein product of the hsrw cytoplasmic transcript suggests that the protein may be considerably more labile than the other heat shock proteins. If the protein is, in fact, labile, it might be possible to stabilize it by covalent linkage to another protein and thus visualize it directly. One way to obtain such a covalent linkage is by a translational fusion between ORF-0 and the CAT ORF. We therefore created a construct similar to ORF-0-CAT but with the 21st codon of ORF-0 linked directly to the CAT ORF, in frame. For this construct we used the D. hydei hsrw gene since it has a convenient restriction site for fusing the ORF-0 to the CAT gene in frame. (As stated earlier, the D. hydei o3 shows the same monosome/disome association as the D. melanogaster o3.) The D. hydei ORF-0 has 23 codons; thus, the ORF-0 in the construct is complete except for its last two codons. In this new gene construct, the ORF-0/CAT fusion, transcription of the RNA is still dependent on the Drosophila promoter, as it was with all the other constructs. If the ORF-0 is translated in vivo, we would expect that, in this fusion construct, synthesis of the fusion protein would begin at the ORF-0 start codon and proceed through the ORF-0 into the CAT sequence, producing a CAT protein that is larger by the size of the linked ORF-0 protein.

The ORF-0/CAT construct does direct the translation of the expected fusion protein. The fusion protein encoded by the ORF-0/CAT gene was identified and visualized by binding to anti-CAT antibody on Western blots (Fig. 9). Verification that the correct fusion protein was made is based on the apparent molecular size of the protein as compared to the unaltered CAT gene product. The resulting ORF-0/CAT fusion protein would be 253 amino acids in length, which is 34 amino acids longer than unaltered CAT protein. This would cause an approximate increase in apparent molecular weight of CAT protein from 24.1 kD to 27.8 kD. The putative fusion protein was indeed larger than the unaltered CAT protein by the amount expected if translation began at the methionine start codon in ORF-0, using the natural Drosophila sequence signals for initiation, and continued through the CAT sequence.

The translation of this fusion protein in the Drosophila cell under natural conditions provides strong evidence that ORF-0 is indeed translated in vivo. Translation of the fusion protein also indicates that the ORF-0 does not impede the translation...
Figure 7. Position of particles containing *hsrω*-CAT transcripts in polysome gradients. (A) Northern analysis of 15–50% sucrose gradients generated from lysates of the cell lines; (a, c, and d) *S3-ω*-leader-CAT; and (b) *S3-ORF*-ω-CAT. The Northern blots were probed with cDNAs for (a and b) CAT RNA; (c) hsp82 RNA; and (d) ω3 RNA. Lysates from equal numbers of cells grown at 25°C were loaded on each gradient in (a and b) and Northern analysis on both was performed at the same time. Above the blots is the UV absorbance profile of the gradient determined as it was collected with four times increase in sensitivity after fraction 3. The portions of the gradient in each Northern blot fraction are indicated. Fraction 3 contains translating monosomes; fraction 4 contains disomes and trisomes; fraction 5 contains trisomes and quadrisomes; and higher number fractions contain larger polysomes. In the cell line that is making CAT protein (a) ω-leader-CAT RNA is on polysomes containing up to 4–5 ribosomes, although ω3 RNA is in the monosome and disome region (d). In the cell line that is not making CAT protein the ORF-ω-CAT transcripts are in the monosome and disome fractions, as are the transcript for endogenous ω3 RNA. (B) Northern analysis of a 5–20% sucrose gradient of *S3-ORF*-ω-CAT cells probed with cDNA for ω3 RNA, showing the monosome and disome regions at greater resolution. This lysate was prepared from the same number of cells and analyzed at the same time as a and b above. Drawn over the blot is the UV absorbance profile that was determined as the gradient fractions were collected. The portions of the gradient from which each fraction was taken are indicated. Fractions 1–4 represent prepolysomal RNP particles; fractions 5 and 6 contain translating monosomes; and fractions 7 and 8 contain translating disomes.

Discussion

The *hsrω* locus is distinctly different from the other major heat shock loci (for review, see Pardue et al., 1987; Bendena et al., 1989a). It does not encode a known heat shock protein, and yet the size of the puff in polytene cells and the level of *hsrω* transcripts induced in diploid cells suggest that *hsrω* is an important locus. The *hsrω* locus has heat shock promoter elements and is induced by heat shock just as are the other
heat shock genes, but hsro\textsubscript{2}, like the gene for hsp82, is also constitutively transcribed. In addition, studies of polytene puffs and, more recently, analyses of transcripts from other cell types (Bendena et al., 1989b) have shown that levels of hsro\textsubscript{2} transcripts can be modulated by a number of agents that do not affect other heat shock loci, suggesting that hsro\textsubscript{2} is especially sensitive to the environment. This raises the question of what roles the hsro\textsubscript{2} transcripts play. In this paper, the role of the cytoplasmic transcript, o\textsubscript{3}, has been considered.

Some clues as to the function of o\textsubscript{3} come from consideration of the sequences conserved in different Drosophila species. Each Drosophila species has one heat shock puff with the phenotypic peculiarities of the hsro\textsubscript{2} locus. Sequences of the phenotypically equivalent puff from three distantly related species, D. melanogaster, D. hydei, and D. pseudoobscura, have been cloned. Although the hsro\textsubscript{2} sequences in these species have diverged sharply enough to almost eliminate crosshybridization, the very distinctive structure of the three transcripts is conserved. In addition, sequence alignments show a few short regions of conservation. The conserved regions include the heat shock–inducible transcriptional promoter, the start site of transcription, the 5' and 3' splice sites and the polyadenylation sites. Taken together, these conservations argue that the cell is preserving the capacity to express a cytoplasmic RNA of this size. The sequence comparisons give little evidence that there is conservation of the ability to encode a protein; all ORFs are small, and larger ORFs can be found on the nontranscribed DNA strand. Only one ORF shows any sequence conservation between any of the Drosophila species studied (Garbe et al., 1989).

The conserved ORF in o\textsubscript{3} is also the first ORF in each species in what is thought to be a favorable sequence context for translation (Garbe et al., 1989). It is this ORF that we have named ORF-ω. The first four amino acids of ORF-ω are conserved in all three species. Conservation is at the amino acid level; the glutamic acid codon is specified by its alternative codon in D. hydei. In each species, the polypeptide specified by ORF-ω is small, 23 amino acids in D. hydei, 24 in D. pseudoobscura, and 27 in D. melanogaster. Beyond the first four amino acids there is little obvious conservation of sequence. Only two more amino acids are conserved in all three species and six more are conserved in two species. We can see no strong evidence that the amino acid replacements would conserve any sort of a structure in the peptide.

In spite of the lack of a long ORF, there is strong evidence that ω3 is involved in translation. It is associated with the polysomes as judged by sedimentation and by buoyant density. It is moved out of the polysome region only by agents that dissociate translating RNAs from polysomes. Furthermore, the ω3 transcript is on monosomes and disomes, as expected from the sizes of all of its potential ORFs. Had the RNA associated with the entire spectrum of polysome sizes, it would have opened the possibility that ω3 was an accessory, rather than a translated, RNA. Other work has shown that any inhibitors of protein synthesis, whether acting at initiation or elongation, stabilize the ω3 transcript, which normally turns over within minutes in nonstressed cells (Bendena et al., 1989b). This stabilization is another piece of evidence that ω3 is closely associated with translation.

Because earlier attempts to find a protein product from the hsro\textsubscript{2} locus had been negative, we tried a less direct assay for translation. The ORF that we tested, ORF-ω, was chosen because it is the only one conserved in its position in all three Drosophila species. In addition, although it is not the first ORF that a ribosome would encounter in scanning the hsro\textsubscript{2} transcript (in two species), in all three species ORF-ω is the
first ORF that has its initiation codon in a favorable sequence context, as determined by Kozak (1986). (A) A survey of sequence conservation around start codons in Drosophila messages (Cavener, 1987) suggests that translation initiation in this genus requires a similar sequence context.) There is a 5' ORF in an unfavorable context in D. melanogaster and D. pseudoobscura; the lack of conservation of the first ORF in D. hydei also argues against its importance to the cell. Our experiments support the idea that this first ORF is untranslated.

Our indirect assay for translation was based on the hypothesis that if ORF-ω were translated, most ribosomes would be impeded from progressing on to the translation start site for the CAT protein. The results of our experiments are entirely consistent with the hypothesis. Although the ORF-ω-CAT construct differs from the ω-leader-CAT construct only by having ORF-ω inserted in front of the CAT gene, the difference in the level of CAT activity and CAT protein produced by the two constructs was dramatic. Cells with ORF-ω made almost no CAT protein although they contained even more CAT mRNA than cells with the other construct. As expected of an RNA in which the ORF-ω was being translated, the ORF-ω-CAT transcript was on monosomes and disomes. This eliminates the alternative possibility, that structural considerations in the ORF portion of the transcript inhibited translational initiation. A more direct demonstration of the translation of ORF-ω comes from the construct in which ORF-ω was fused in frame to the CAT ORF. Cells with this construct produced CAT protein that was larger by the amount predicted from the extended coding region, showing that the ORF-ω initiation site was functional in vivo.

The evidence that ω3 was translated led us to extend the earlier search for a small polypeptide that might be encoded by one of the ω3 ORFs. We have looked throughout the period of maximal heat shock–induced increase in the ω3 levels and have seen no polypeptides of the appropriate size (3 kD) in either heat shock or control cells. Thus, if ω3 is translated, the product must be either rapidly turned over or rapidly sequestered, perhaps by linking to other proteins. Although ORF-ω has a conserved cysteine, we do not think that its linkage to other proteins by disulfide bonds has prevented its identification here, because the conditions of electrophoresis should have reduced such bonds. It is possible that the hsrω polypeptide is normally covalently linked to other cellular proteins, much like ubiquitin, so that there is little of the unmodified form to be found (Finley and Varshavsky, 1985). In fact, one of the conserved residues of ORF-ω is a lysine, the amino acid by which ubiquitin forms bonds to other proteins; however, such posttranslational linkage also seems unlikely. We are easily able to identify the unmodified form of ubiquitin among the 35S-labeled heat shock proteins from S3 cells; if the hsrω polypeptide is linked to other proteins, we should detect some unlinked form unless that attachment is significantly more efficient than the ubiquitin linkage.

Although we cannot rule out the possibility that the hsrω translation product is rapidly sequestered, there are several reasons to think that it is rapidly turned over, possibly even before leaving the ribosome. Our searches have failed to turn up any free polypeptide, even in cells carrying the ORF-ω-CAT construct. If the amount of hsrω transcript on polycytes is any measure of its translation, these cells should be making much more of the hsrω polypeptide than is normal for the cultured cells. The extra peptide might be expected to overload even an efficient sequestration system, and yet we detect no 3-kD polypeptide in these cells. We also note that, in the three Drosophila species, the conservation of the polypeptide seems almost entirely limited to the first four amino acids and the small size (although the size varies slightly). If what is required is the act of translation itself and then rapid degradation of the product, the conserved features seem adequate. If the polypeptide has a more complex role, it might be expected to show more conserved features.

The ORF-ω-CAT fusion protein is easily detected and has given no insight into the fate of the authentic ORF-ω product. The fusion protein is not found in linkage with other proteins or with itself either on reduced or unreduced gels (Fini, unpublished data); however, it is possible that the ability to form such bonds is blocked by CAT linkage. Similarly, if the peptide is normally turned over very rapidly, linkage to CAT protein may stabilize the peptide, either by blocking the carboxy terminus, or by embedding other vulnerable portions of the peptide within the CAT protein structure, and thus protecting it from intracellular proteases.

It is important to note that efficient translation of the ORF-ω can be observed under heat shock temperatures (>33°C in our cell lines) at which only heat shock messages and a few 25° messages can be translated. This suggests an additional evolutionary conservation, that of sequences that are essential to translation at high temperatures. Recently, Hultmark et al. (1986) have given evidence that the first 26 nucleotides of the hsp 22 transcript have elements affecting both heat shock translation and efficient transcription. The 5' ends of hsp 70, 68, 27, 26, 23, and 22 show identical nucleotides in positions −1, +1, 7, 12, 15, and 20. The hsrω genes in the three Drosophila species that we have examined also have these nucleotides in all but position 20 (Garbe et al., 1986; 1989). These sequences could have been retained during
evolution to allow for efficient translation of the conserved ORF under heat shock conditions. Such conservation along with the sequence conservation in the conserved ORF provides a substantial argument for the importance of this translation to the cell.

In what way could translation of ORF-ω affect the cell? Since we cannot find detectable levels of the peptide, it is possible that translation of the conserved ORF is the important factor; the protein product may have no consequence and be rapidly degraded by the cell. Such a role has been suggested for other upstream ORFs in several mRNAs. The first ORF of Rous sarcoma virus has only 7 codons. This ORF is evolutionarily conserved but can not be linked to a protein product in vivo, although a very short-lived protein product has been detected by in vitro translation (Hackett et al., 1986). It has been proposed that translation of this ORF acts to place the transcript in an appropriate configuration for viral coat assembly. For SV40 early mRNA (Kahlili et al., 1987) or yeast GCN4 mRNA (Mueller and Hinnebusch, 1986; Fink, 1986) the translation of small, upstream ORFs appears to play a role in regulation of the translation of protein-coding ORFs located more 3' on the message. For a number of different mRNAs, the machinery for regulation of mRNA half-life appears to be associated with the polysomes, requiring a message to be actively translated in order to be degraded (Hunt, 1988). A similar mechanism may well be operating with the ω3 RNA since we have found that its half-life is significantly and specifically increased when cells are treated with inhibitors of protein synthesis (Bendena et al., 1989b). If translation serves only to regulate the turnover of ω3 RNA, the RNA must have some additional function. We should note that ORF-ω represents only a small portion of this 1.2-1.3 kb RNA, and yet the size of the transcript is conserved in all the Drosophila species. There are several small islands of conserved sequence in the transcripts that may well have functional significance which we do not yet understand.

Our results do not eliminate the possibility that ORF-ω could encode a protein that performs a function. If so, the function might be expected to be a regulatory one since the protein must be short lived, if it exists at all. Peptide hormones offer examples of small regulatory molecules with minimally conserved sequences and very short half-lives (Tager and Steiner, 1974). We note that three of the six amino acids conserved in ORF-ω in all three Drosophila species are conserved at the level of the amino acid rather than the nucleotide sequence (Garbe et al., 1989). This might suggest conservation of the ability to make a functional protein. On the other hand, the conserved amino acids are clustered at the amino-terminal end, reminiscent of the four conserved amino acids in β-tubulin that are required to assure the regulated turnover of the mRNA while it is on the polysomes (Yen et al., 1988). If the ω3 transcript is indeed serving to monitor the translational state of the cell, a similar polysome-related turnover might obtain.

How could translation of the ORF-ω contribute to cell function at ambient and heat shock temperatures? We have been impressed with the sensitive way in which the level of all the hsrω transcripts respond to various external conditions. If, as we propose, the levels of ω3 RNA are indeed controlled by translation on polysomes, then this may be a finely tuned mechanism to link the levels of ω3 to protein synthesis. Thus, one intriguing possibility is that the level of the ω3 transcript serves to communicate, to the nuclear compartment of the cell, information about the status of cellular protein synthesis at any particular time.

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