



1990; Picard et al. 1978; Chaboissier et al. 1990, 1995). The dysgenesis caused by the *I* element in *D. melanogaster* is restricted to the female germline and has not been reported in males. Furthermore, the transposition rate correlates with the level of sterility in the SF females (Chaboissier et al. 1990; Seleme et al. 1999, 2006).

The number of *I* copies in *D. melanogaster* is approximately 30, but only five functional sequences are known to be dispersed on the chromosomal arms. The complete and functional copies are 5.4 kb long, possess two long ORFs with cysteine-rich motifs, apurinic–apyrimidic endonucleases, reverse transcriptase (RT), and RNase H domains, and the 3′-end has several TAA repeats. These copies are expressed in the nurse cells of dysgenic ovaries, and the transcripts are transported into the oocytes, where retrotransposition occurs (Pelisson and Picard 1979; Fawcett et al. 1986; Vaury et al. 1990; Seleme et al. 1999, 2005). In *Drosophila*, piwi-interacting RNAs (piRNAs) are important posttranscriptional regulators that are responsible for TE mRNA degradation (Khurana and Theurkauf 2010; Senti and Brennecke 2010; Kelleher et al. 2012; Akkouché et al. 2013; Dufourt et al. 2013). Specifically, the piRNAs are implicated in maternal *I* element control in *Drosophila* (Brennecke et al. 2008; Chambeyron et al. 2008).

The implication of TEs in hybrid dysgenesis is one of the examples in which transposition may drive population isolation and is often proposed as one of the first steps of speciation (Kidwell and Novy 1979; Fontdevila 2005). However, the TE dynamics of expression and transposition have not been well described during this process. Hence, several questions remain unanswered regarding this subject. If TEs act in the first steps of speciation, what happens when crossing very closely related species that still have incomplete reproductive isolation? In particular, are *I*-like elements associated with the incompatibility observed, and are they related to high rates of hybrid transposition as reported in other species (Ungerer et al. 2009; Cavallini et al. 2010; Moschetti et al. 2010; Kelleher et al. 2012; Vela et al. 2014)? In this report, we made a first investigation to check whether *I*-like elements could potentially be involved in hybrid sterility.

*Drosophila mojavensis* and its sister species analyzed in this study—*D. arizonae*—are a perfect pair of species with which to address these questions. They are found in the desert of the southwest United States and Mexico and share sympatric areas in southern Arizona and the State of Sonora (Mexico). These two species present three interesting features that facilitate the investigation of the above questions: 1) Their hybrids can be produced in the laboratory; 2) the genome of *D. mojavensis* has been sequenced, which allow us to analyze the sequences of its TEs; and 3) these species exhibit variable degrees of pre- and postzygotic isolation (Wasserman and Koepfer 1977; Reed and Markow 2004; Massie and Markow 2005). Moreover, the male offspring of *D. arizonae* females and *D. mojavensis* males is sterile, but in the reciprocal cross, the hybrids have motile sperm depending on the origin of the *D. mojavensis* population

that is used in the crosses (Reed and Markow 2004). Additionally, the strong prezygotic isolation is higher between flies from sympatric areas than between the allopatric flies, but the isolation level depends in part on the direction of the crosses and the geographic origin of the populations (Wasserman and Koepfer 1977; Ruiz et al. 1990; Reed and Markow 2004; Massie and Markow 2005).

We found four potential full-length copies of *I*-like elements and reconstructed their phylogenetic relationships with other *I* family members in *Drosophila* species, thus establishing the vertical and ancestral inheritance of the *I* element of *D. mojavensis* from a common ancestor shared with the *melanogaster* group. We also showed that the *D. mojavensis* *I*-like sequences are transcriptionally and transpositionally regulated in the hybrid female germline. However, a significant increase in transcription was detected in testes from hybrids coming from *D. arizonae* mothers. This is the first report that shows the activity of *I*-like elements in the male germline, and it suggests a link with the male-sterile phenotype observed in the male hybrids of the studied species.

## Materials and Methods

### Genomic Analyses

To investigate the occurrence of *I*-like elements in the *D. mojavensis* sequenced genome, we used the canonical *I* sequence from *D. melanogaster* (*I*<sub>DM</sub>), which is available in the Repbase library Rel.16.06, v.4 (Fawcett et al. 1986). This sequence was blasted against the scaffolds of the *D. mojavensis* genome (version-70, 13/dmoj\_caf1) on the Ensembl Metazoa platform using BLASTN *D. mojavensis* (<http://metazoa.ensembl.org/index.html>, last accessed July 4, 2014).

The hits were manually curated, and only the largest and putative complete copies (more than 2,000 bp) were selected. We then selected the *D. mojavensis* copies that were over 80% identical between them. The sequences with the above characteristics were selected and analyzed using the ORF-FINDER web tool (<http://www.ncbi.nlm.nih.gov/projects/gorf/>, last accessed July 4, 2014) to identify putative coding regions. The ORFs identified were compared and confirmed by BLAST search for protein sequences at the NCBI website, and the conserved protein domains were checked on the conserved domain database (<http://www.ncbi.nlm.nih.gov/Structure/cdd/cdd.shtml>, last accessed July 4, 2014). The sequence with the most complete structure was selected and hereafter referred to as “Reference Copy” or “RC” (fig. 1).

The RC was used to search for *I*-like element copies in the *D. mojavensis* genome with identity (ID) higher than 80% and length greater than 200 bp. This criterion was chosen based on the 80-80-80 rule for TE family determination, as proposed by Wicker et al. (2007) for the identification of TEs belonging to the same family. The position of each copy (start and end)

on the genome scaffolds, the length of each fragment aligned by BLAST, and the number of the unidentified nucleotides (*N*) were recorded (table 1). We performed multiple alignments of the gag and RT domains (nt) identified in the copies, and we determined the ID of each domain, based on those of the RC, using Bioedit version 7.0.4.1 (Hall 1999). A drawing of the copies is shown in figure 2 and represents the integrity of the putatively full-length copies compared with RC. To confirm that we recovered unique copies in the genome, the 5'- and 3'-end flanking sequences of each copy were retrieved, and their identities were determined compared with the RC flanking sequences. All copies corresponded to different insertions.

To check the functional organization of all putatively full-length copies of *D. mojavensis*, the 5'-end sequences were compared with the sequence of the *I* promoter of *D. melanogaster* (Minchiotti et al. 1997) using BLAST2seq.

The chromosomal location of the most complete *I* copies was inferred from the scaffold coordinates of each copy, and the correspondence of scaffolds with *D. mojavensis* polytene chromosome maps was performed using information found in Flybase.org (<http://flybase.org/maps/chromosomes/maps.html>, last accessed July 4, 2014) and in Schaeffer et al. (2008). For an intrachromosomal distribution of the copies, each *D. mojavensis* chromosome (X, 2, 3, 4, 5, and 6) was divided into three regions: Distal (10% of the sequence), central (80% of the sequence), and proximal (or centromeric, 10% of sequence) segments in relation to the position of the centromere, to test the distribution of the copies in these regions (Marzo et al. 2013).

The nomenclature used to identify each copy followed Wicker et al. (2007); for example, RII\_DMJA1 consists of RII (for the superfamily), DMJ (for *D. mojavensis*), and A to Z (for the clades formed by phylogenetic analyses, where the exception is X, which was used for the undetermined groups), followed by the number of each copy within each clade.

### Evolutionary Analyses of RT Sequences

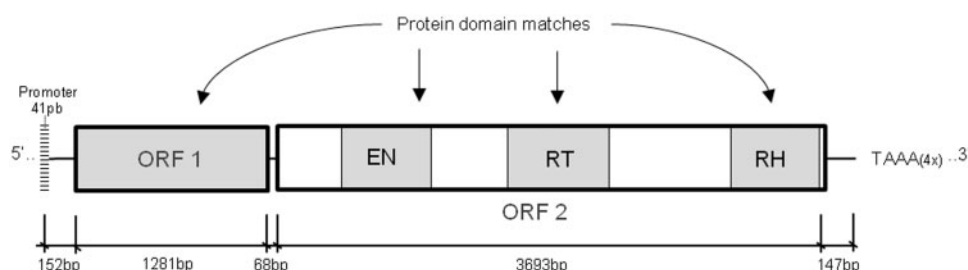
We reconstructed the evolutionary relationships between the *D. mojavensis*/*D. arizonae* *I*-like sequences with all the *I* superfamily members of *Drosophila* species deposited in the

Rebase library (<http://www.girinst.org/rebase/>, last accessed July 4, 2014). Each retrieved element was analyzed using ORF FINDER (<http://www.ncbi.nlm.nih.gov/gorf/gorf.html>, last accessed July 4, 2014), and only the amino acid sequences of the second ORF were used for further analysis. We identified the RT domain in each protein sequence using the structural domain bank from the Pfam platform (<http://pfam.sanger.ac.uk/>, last accessed July 4, 2014). The *Drosophila* Repbase RT domains were aligned with the *D. mojavensis* and *D. arizonae* RT domains using MEGA5 (Tamura et al. 2011). For *D. mojavensis* (strains #15081-1352.01 and #15081-132.26 from US San Diego *Drosophila* Stock Center) and *D. arizonae* (strains #15081-1271.17 and #15081-1271.18 from US San Diego *Drosophila* Stock Center), we amplified and sequenced the 627 bp RT sequence (RT\_If: 5'-GCCTAGTCATCCCTATC-3' and RT\_Ir: 5'-TGTTAGC GCCGGTTGTATT-3'), corresponding to the position 2,813 and 3,198 in the *D. mojavensis* RC sequence. Polymerase chain reaction (PCR) cycling parameters were: 94 °C for 3 min, 40 cycles of 94 °C for 45 s, 56 °C for 45 min, and 72 °C for 1 min, followed by a 72 °C for 7 min. The fragments obtained were purified directly from the PCR product, using the GFX PCR DNA and Gel Band Purification Kit (GE Healthcare), and cloned with the TOPO TA Cloning Kit (Invitrogen). Five randomly chosen clones were sequenced using the M13 universal primers.

We performed the amino acid phylogenetic reconstruction using the maximum-likelihood method, JTT model, and 500 bootstrap replicates as implemented in PHYML 3.0 (Guindon et al. 2010) at the ATGC: Montpellier Bioinformatics platform (<http://www.atgc-montpellier.fr/>, last accessed July 4, 2014). Two sequences of other LINE superfamilies were used as outgroups: The *D. melanogaster* *Fw* non-LTR retrotransposon (GenBank accession number: M17214.1) and the *D. yakuba* *Helena* non-LTR retrotransposon (GenBank accession number: AF012049.1).

### RNA Extraction and RT-quantitative PCR

The fly samples were obtained from the US San Diego *Drosophila* Stock Center. We used strains of *D. mojavensis*



**FIG. 1.**—The diagram represents the structure of the putative “RC” (5,382 bp) of the *I*-like non-LTR retrotransposon identified in the *Drosophila mojavensis* genome. The boxes represent the two ORFs, ORF1 and ORF2. EN, endonuclease domain; RT, reverse transcriptase domain; RH, RNase H domain; and crosshatch box, promoter region.

Summary of the *I* Element Copies in the *Drosophila mojavensis* Genome

<sup>a</sup>Sequence used in phylogeny (fig. 4).



(2009). The RT-qPCR experiments were performed with technical triplicates. Only RT-qPCR experiments with efficiencies greater than 1.9 were retained.

### Fluorescent In Situ Hybridization

*I* transcripts were detected by in situ hybridization in 5-day-old ovaries of parents and hybrids according to Akkouché et al. (2012). Ovaries were hybridized with a 952-bp riboprobe that corresponded to the RT and RNA-H domains in the RC and included a T3 promoter site in the reverse primer (forward: 5'-GCA ATA CAA CCG GCG CTA ACA-3', reverse: 5'-TGG CTG TGG ATT TGG CTG TGA ATT AAC CCT CAC TAA AGG GA-3'). For the in vitro transcription, a digoxigenin (DIG)-RNA labeling mix (Roche) was used. After hybridization and stringency washes, an anti-DIG POD antibody mix (ROCHE) was used for chemiluminescent detection, with a fluorescence amplification (the Tyramide Signal Amplification Kit; PerkinElmer). The DNA was labeled by SYTOX Green Nucleic Acid Stain (Life Technology), and the visualization was performed with a Zeiss LSM510 Meta confocal microscope. Tissues without anti-DIG treatment were used as a negative control, since no label is expected without this secondary antibody. Five-day-old testes hybridization was performed with the same *I* element probe according to Morris et al. (2009). To be able to localize the *I* transcripts in the testis, we used Cyclin B as a positive control. The Cyclin B gene is expressed at very high levels in primary spermatocytes, whereas the mitotic cells in the germinal proliferation center present lower levels of expression (Morris et al. 2009). We used a Cyclin B riboprobe of 730-bp in which primers were designed for the *D. mojavensis* gene and include a T3 promoter site (forward: 5'-CGA TGT CCT TGT CCA CCA AA-3', reverse: 5'-GCA GCC GTA TAA CGG GAA TAG ATT AAC CCT CAC TAA AGG GA-5') (supplementary fig. S1, Supplementary Material online). The testes images were processed in a Zeiss Axioskop 2 mot plus microscope and analyzed with Axiovision Release 4.8 software.

### TE Activity (Transposon Display)

Transposon display (TD) was performed with protocols adapted from several reports (Munroe et al. 1994; Esnault et al. 2008; Akkouché et al. 2012). Total genomic DNA was isolated from single individuals used in crosses between *D. mojavensis* females and *D. arizonae* males, and the F1 single hybrids were backcrossed with *D. arizonae* males to obtain the F2 generation (supplementary fig. S2, Supplementary Material online). We could only perform crosses in this direction because individual crosses between single *D. arizonae* females with single *D. mojavensis* males did not give any offspring. This is because flies will not mate under the low density condition of single pair crosses. From the six individual crosses performed with the parental lines, we were able to follow two independent families. Individuals

analyzed by TD are displayed in supplementary table S1, Supplementary Material online. The TD technique also allowed us to have an estimation of the copy number in each strain.

The genomic DNA (100 ng/μl) was digested with HindIII (10 U/μl, 3 h at 37 °C). The MSBE adaptor was linked with the HindIII enzyme adaptor site in one double-stranded reaction (HindIII-MSEB) using 100 μM of Hindlink, 100 μM of MSEB, SSC 20×, and 1 M Tris in total volume of 333 μl. The product was denatured for 5 min in 92 °C and cooled at room temperature. The linker HindIII-MSEB was ligated to digested DNA with a T4 DNA ligase (10 units), enzyme buffer (5×), and digested DNA in a 50-μl reaction for 3 h at 23 °C. DNA was amplified with a primer adaptor (LNP 5'-GAA TTC GTC AAC ATA GCA TTT CT-3') and a forward primer inside of the 3'-end of the *I* element (5'-TAA CTG TCC TGC AAC TTC CCA CCT-3'). The amplification conditions were as follows: denaturation at 94 °C, 2 min; followed by 94 °C, 30 s; 59 °C, 1 min; 72 °C, 1 min; 35 cycles; then, 72 °C, 10 min. Subsequently, the product amplification was run on a gel, and analyses were performed with QIAxcel ScreenGel 1.1.0 software.

The TD method was tested with *D. mojavensis* genomic DNA, extracted from a single flies, that was digested by HindIII and amplified by PCR with HindIII-MSEB and *I* primers. Then, two fragments (842 and 450 bp), identified in silico and observed in electrophoresis, were cloned (with five clones for each fragment), amplified by PCR with M13 universal primers, and sequenced. The alignment of these sequences against the *D. mojavensis* RC confirmed the veracity of the fragments obtained by enzyme digestion and amplification for the *I* element.

## Results

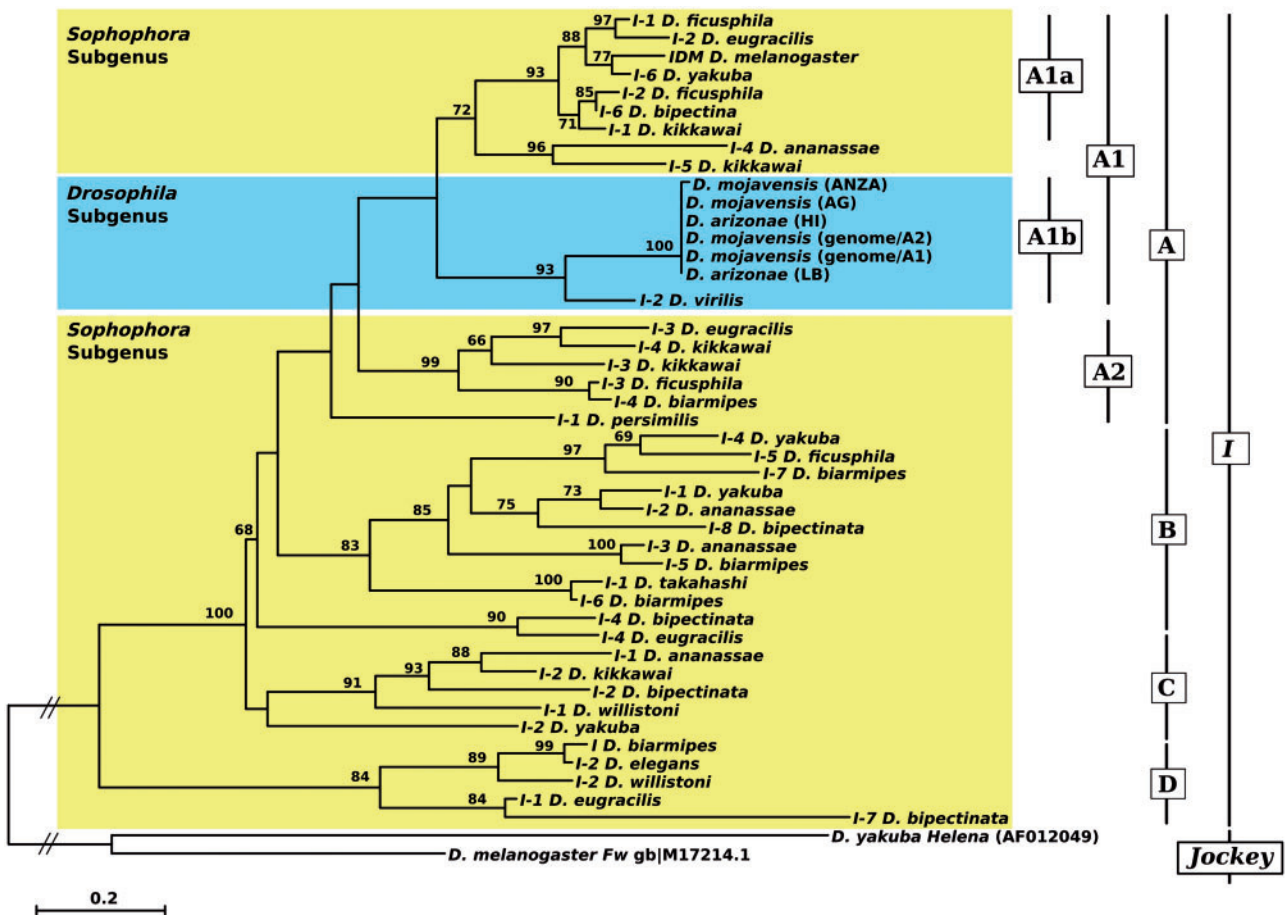
Our study revealed the presence of an *I*-like element in *D. mojavensis* and *D. arizonae* that is closely related to the *I* element found in the genomes of the *D. melanogaster* subgroup. In silico and in vivo analyses showed the occurrence of complete and transcriptionally active copies in both species and in their hybrids.

### The *I* Element of *D. mojavensis*

We detected 10 *I*-like copies in the *D. mojavensis* genome sequence (table 1). One potentially complete copy, called RC (RII\_DMJA1), is 5,382-bp long and contains two ORFs (ORF 1 and ORF 2) with no internal stop codons and with all the expected protein domains for this type of non-LTR retrotransposon (Malik et al. 1999; fig. 1). It has 62.42% nucleic ID with the *D. melanogaster I* element and is only 7 bp longer. We identified three other copies that are most likely complete: RII\_DMJA2, RII\_DMJA3, and RII\_DMJX1 (table 1 and fig. 2), which show high ID and conserved protein domains compared with the RC, but their sequences were interrupted with unidentified nucleotides (N). Similarly, the RII\_DMJX5

		Sites																																												
		1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40	41				
<i>D. melanogaster</i>		C	A	T	T	A	C	T	C	A	C	T	T	C	A	A	C	C	T	C	C	G	A	A	G	A	G	A	T	A	A	G	T	A	C	G	T	G	C	C	T	G	C	T	A	C
DMJA1 (rc)		.	.	G	.	.	T	.	.	.	.	.	.	.	.	.	.	.	.	T	C	C	.	G	.	.	.	.	.	.	.	A	.	.	.	.	.	.	T	G	.	A	T	.	.	
DMJA2		.	.	G	.	.	T	.	.	.	.	.	.	.	.	.	.	.	.	T	C	C	.	G	.	.	.	.	.	.	.	A	.	.	.	.	.	.	T	G	.	A	T	.	.	
DMJA3		.	.	G	.	.	T	.	.	.	.	.	.	.	.	.	.	.	.	T	C	C	.	G	.	.	.	.	.	.	.	A	.	.	.	.	.	.	T	G	.	A	T	.	.	
DMJX1		.	.	G	.	.	T	.	.	.	.	.	.	.	.	.	.	.	.	T	C	C	.	G	.	.	.	.	.	.	.	A	.	.	.	.	.	.	T	G	.	A	T	.	.	

**FIG. 3.**—Sequence alignment of the *Drosophila melanogaster* *I* promoter sequence with the putatively complete *D. mojavensis* *I* copies. Polymorphism sites are highlighted in gray and identical nucleotides are indicated by a dot.



**FIG. 4.**—Phylogenetic analysis of *I* elements based on the amino acid sequences of the reverse transcriptase domain. The tree was obtained using the maximum-likelihood method, JTT model, and 500 bootstrap replicates. Only bootstrap values greater than 50% are indicated. The tree has been rooted by the *F* element from *Drosophila melanogaster* and *Helena* from *D. yakuba*.

copy, with 1,629 bp aligned against RC, may also be complete because it has a long N stretch at its 5'-end. The other copies were small and fragmented by insertions and deletions. The identification of the relative chromosome position of these copies, as established by analyzing the polytene chromosome maps found in Flybase (Schaeffer et al. 2008), revealed that three of the four putative full-length copies, RII\_DMJA1, RII\_DMJA2, and RII\_DMJA3, were inserted in the central portion of different chromosome arms corresponding to chromosomes 2, 3, and 5, respectively.

Moreover, we looked for the *I* element promoter in the *D. mojavensis* copies by aligning the flanking sequences of each copy with the promoter sequence of the *I* element of *D. melanogaster* (Minchiotti et al. 1997; Han 2010). We identified a putative promoter in the RC that has the same length (41 bp) and is well conserved (76% ID) compared with that of the *D. melanogaster* *I* element. This promoter region is identical in the four putative full-length copies found in the *D. mojavensis* genome (fig. 3). The four longest *I* sequences have a poly-A tail in the 3'-UTR, with an average of four TAAA repeats.

### Phylogenetic Relationship between the *D. mojavensis* *I*-Like Element and the *I* Elements of Other *Drosophila* Species

The phylogenetic relationship of the different *I*-like elements was reconstructed using the RT protein sequences (see Materials and Methods for details) (fig. 4). It allowed us to distinguish four major clades of sequences (A–D). Different *I*-like sequences are present in species of the *melanogaster* group of the subgenus *Sophophora* (see [supplementary fig. S3, Supplementary Material](#) online, for the species phylogeny), which are included in all four clades (fig. 4). Clade A can be divided into two groups of sequences (Clade A1, bootstrap: 70% and Clade A2, bootstrap: 99%). Inside Clade A1, which branches in two smaller clades (A1a and A1b), all the *I*-like sequences of the *D. mojavensis* and of *D. arizonae* cluster in a highly supported clade (bootstrap: 100%), which clusters with the *D. virilis* sequence in a well-supported monophyletic group (A1b: 99%). This clade is closely related to the group of sequences from the *melanogaster* group (Clade A1a), which includes the canonical *I* element from *D. melanogaster* (*I*<sub>DM</sub>). These two groups of sequences (A1a and A1b) form a sister group of Clade A2. This clade includes sequences found in the same species that form Clade A1a with the exception of *D. biarmipes* and of *D. persimilis*, a species of the *obscura* group. The other three clades (B, C, and D) include more divergent *I*-like sequences from species belonging only to the *Sophophora* subgenus: Clade B includes only species from the *melanogaster* group, whereas Clades C and D also include sequences from *D. willistoni* (the *willistoni* species group). The phylogeny reveals that different *I*-like families are harbored by the same species, as is the case for *D. kikukwai*, whose sequences branch into three different clades (A1, A2, and C).

### Activity of the Complete *I* Copies

We observed *I* element mRNA in ovarian cells by immunofluorescence and detected the presence of transcripts in both parents and hybrids. The *I* transcripts had the same localization in both species, with an accumulation in the nurse cells and no labeling of the oocytes. We did not detect any signal in the follicle cells (fig. 5). In testes, we observed labeling in parental lines and both types of hybrids (fig. 6). Transcripts were specifically located in primary and meiotic spermatocyte cells (Fuller 1998). Since *in situ* hybridization is not a quantitative technique, to investigate whether the levels of expression were equal between the different lines (parents and hybrids), we performed expression analyses by RT–qPCR. The results showed transcription of the *I* elements in *D. mojavensis* and in *D. arizonae*, both in ovaries and testes (fig. 7). The levels of transcription between the parental lines and the reciprocal male and female hybrids were not significantly different between ovaries from parents and hybrids (two-way analysis of variance [ANOVA]). However, expression levels were

significantly higher in hybrid testes produced by crosses between *D. arizonae* females and *D. mojavensis* males than in the reciprocal cross ( $F(3, 9) = 84.78$ ,  $P < 0.0001$ ). We obtained the same results with crosses and expression analyses performed again 6 months later ([supplementary fig. S4, Supplementary Material](#) online).

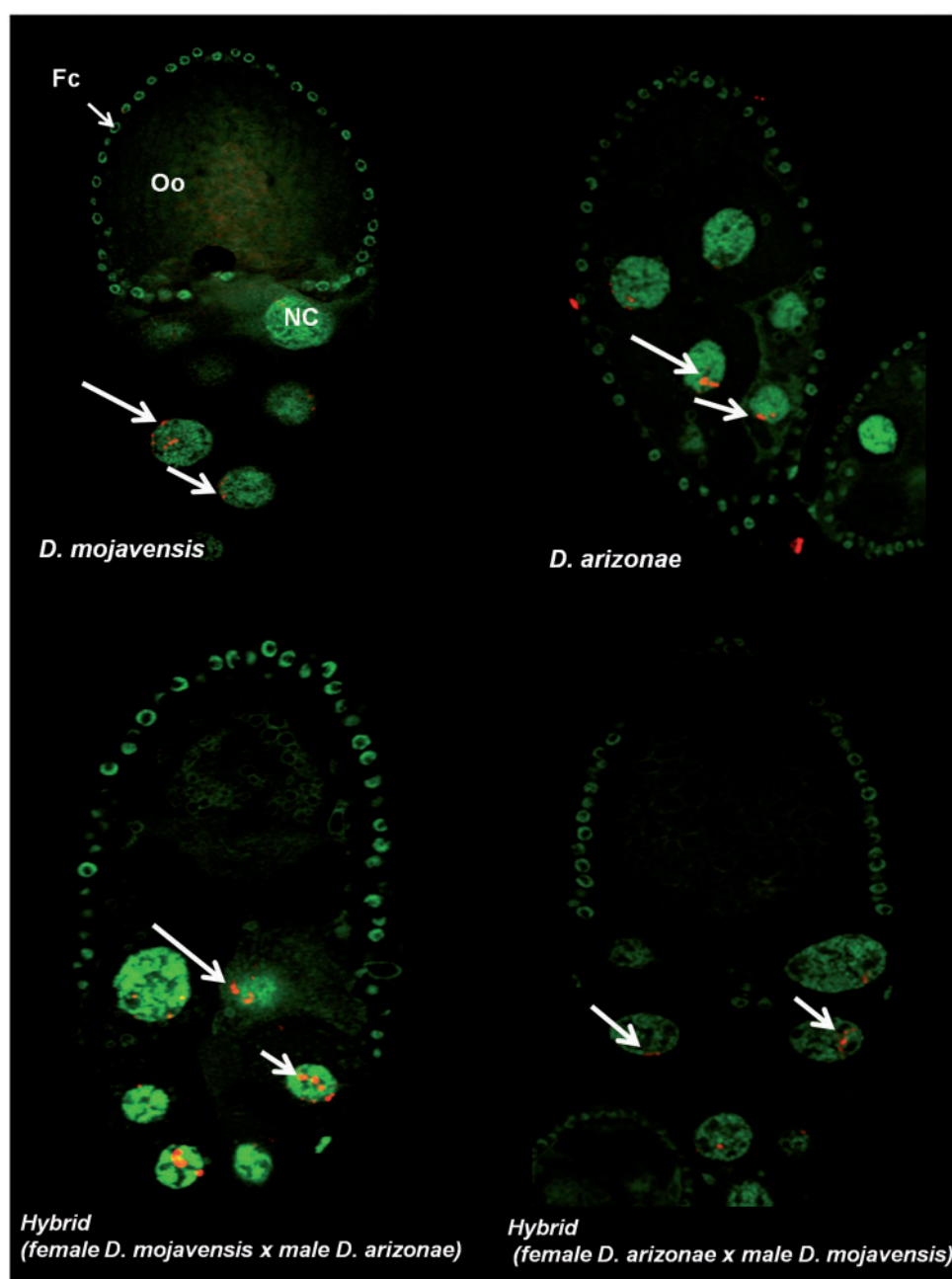
To detect transposition, we performed TD analyses. This experiment needs to be performed with individual crosses, in order to follow new transpositions that arrived in the germ-line cells, and that will be present in all the offspring of the two parents. Unfortunately, these experiments could not be performed for both reciprocal crosses because individual crosses between single *D. arizonae* females with single *D. mojavensis* males did not give any offspring. This cross is only possible with a mass of individuals. TD was thus only performed in one cross direction ([supplementary table S1 and fig. S1, Supplementary Material](#) online). TD also allows determining the copy number. In the individuals that we analyzed, *D. mojavensis* females had one copy of *I*-like element and *D. arizonae* males had from three to five copies of the element. In agreement with the RT–qPCR results, no transposition was detected in the F1 tissues, which indicates that the *I* element does not transpose in the parental gonads. Additionally, no transposition was detected in the F2 generation, indicating that neither the *I* sequences in F1 females nor in *D. arizonae* males induce transposition (data not shown). However, given the limitations of TD technique, we cannot exclude that somatic transposition occurred, but we were unable to measure this with confidence.

## Discussion

### Identification and Phylogenetic Analysis of *I*-Like Elements

In this report, we identified and characterized *I*-like sequences in the sequenced genome of *D. mojavensis*. In addition to several defective and divergent sequences, we identified four putatively complete copies in this species, but only one showed all typical structural domains of the *I* superfamily (Malik et al. 1999). This finding suggests that this copy may also have transposition ability. In *D. melanogaster*, two intact ORFs are a requisite for protein encoding and complete retrotransposition of the *I* factor (Seleme et al. 2005). The presence of unidentified nucleotides in some of the detected copies in *D. mojavensis* suggests the possibility that other sequences are potentially full-length copies. One argument favoring this hypothesis is the conservation of the promoter region that we identified, which is typically not conserved between species (Minchiotti et al. 1997). This promoter region is more conserved between the copies from *D. mojavensis* and *D. melanogaster* (76%), than the RT sequences (66%), which is puzzling. Further work should be done to understand if this could be related to the testis-specific expression in the hybrid male offspring.



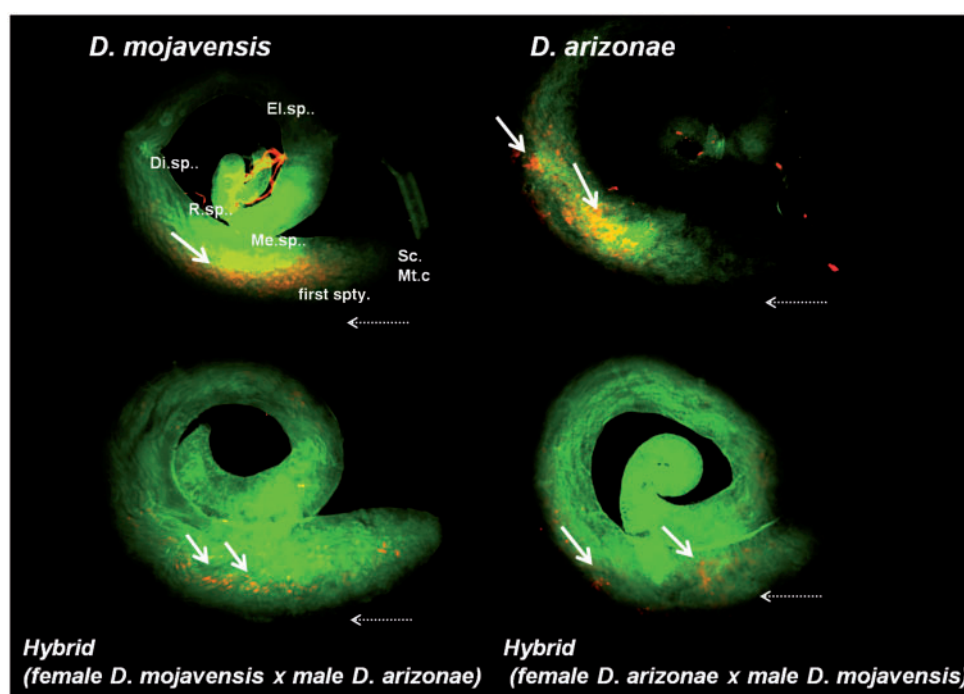


**Fig. 5.**—Immunofluorescence for the *I* element mRNA probe (RNA in red and DNA in green) in the egg chamber of the parental and hybrid crosses. Oo, oocyte; NC, nurse cells; Fc, follicle cells. The arrows indicate mRNA of the detected *I* transcripts.

Non-LTR retrotransposons are common elements in *Drosophila* genomes and are often described as vertically inherited in this group of species (Malik et al. 1999; Granzotto et al. 2011). *I*-like element sequences have been found in many species, and their distribution correlates with the phylogenetic relationships between species (Bucheton et al. 1992), which indicates that they are old components of *Drosophila* genomes. However, *I* elements with a structure strikingly similar to that of *D. melanogaster* (*I*<sub>DM</sub>) occur only in the species

of the *melanogaster* subgroup (*D. simulans*, *D. mauritiana*, and *D. sechellia*). The most widely supported hypothesis is that ancestral *I* elements were lost from the genome of *D. melanogaster* but were then reintroduced recently because older strains of *D. melanogaster* contain only defective and immobile *I* sequences, and all the strains collected after 1930 have both types of sequences (defective and active mobile *I* elements). This observation is in agreement with the hypothesis that the complete and active *I* factor progressively invaded





**FIG. 6.**—Immunofluorescence for the *I* element mRNA probe (RNA in red and DNA in green) in testes of the parental and hybrid lines. Yellow arrows indicate detected mRNA transcripts. Sc.Mt.c, sperm and mitotic cells; first spty, primary spermatocytes; Me.sp., meiotic spermatocytes; R.sp, round spermatids; Di.sp, differentiating spermatids; El.sp, elongating spermatids; horizontal arrow indicates the direction of differentiation (stem cells are located at the apical tip and germ cells move toward the basal end as they differentiate).

*D. melanogaster* after this period (Kidwell 1983). The process by which *I* elements have reinvaded the *D. melanogaster* genome is unknown. Horizontal transfers from another species (Simonel et al. 1988) is one of the possibilities, but more complex hypotheses, such as the reactivation of sequences sequestered in the heterochromatin, should also be considered.

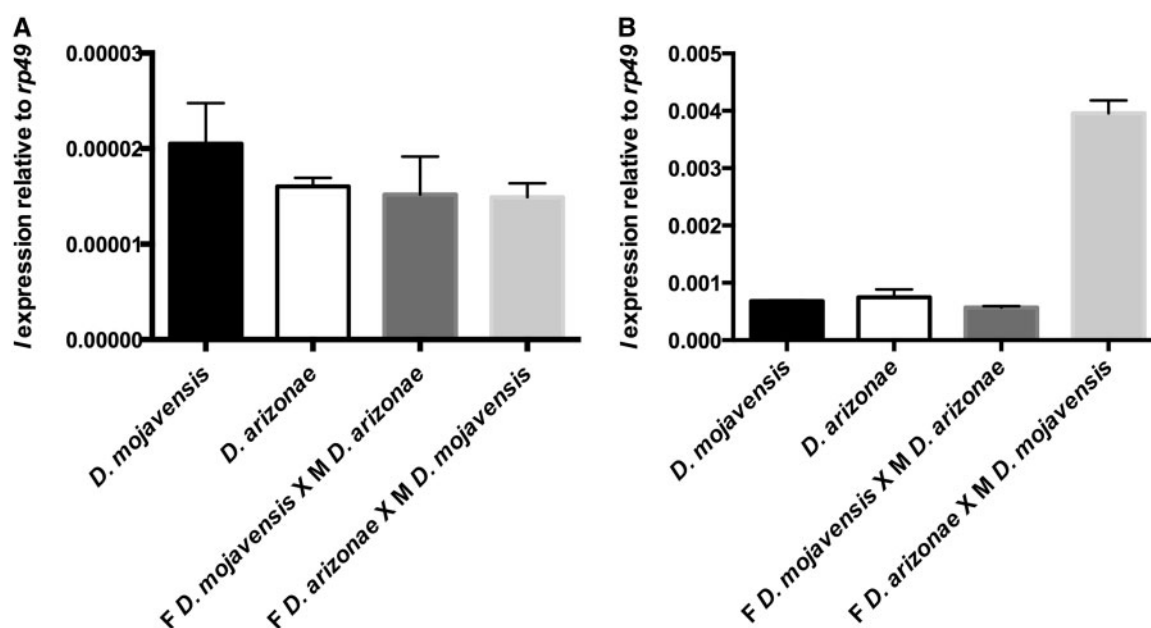
The phylogeny placement of the *I*-like elements from *D. mojavensis* is generally in agreement with the species phylogeny. All the *I*-like sequences of *D. mojavensis* and *D. arizonae* branch together with the sequence of *D. virilis*, as expected based on the species phylogeny because these species belong to the *virilis-repleta* radiation of the subgenus *Drosophila* suggesting a vertical inheritance of these sequences in this *Drosophila* radiation. However, when we analyze the global phylogeny of the *I*-like element, we cannot exclude that some horizontal transfer events may have occurred between other *Drosophila* species. For example, this could have been the case between *D. ficusphila* and *D. bipectinata* (see Clade A1a in fig. 4).

The *I*-like element superfamily is a very old component of the genome of *Drosophila*, and its diversification into several *I*-like families most likely occurred very early in the evolution of *Drosophila*, most likely before the separation of the *Sophophora* and *Drosophila* subgenera. More diversification

occurred in the *Sophophora* subgenus, which was accompanied by sequence loss and horizontal transfer events. In the *Drosophila* subgenus, we can only assume that random loss of the *I* families lead to only one remaining family, which is described in this article and which is closely related to the *I* factor from *D. melanogaster*.

### Activity of the *I* Element

We measured the transcriptional activity of the *I* elements by RT-qPCR and fluorescent in situ hybridization (FISH) experiments in the parental lines and their hybrids. The level of *I* element mRNA was low in ovaries and testes of the parental lines. In the parental ovaries, the *I* element transcripts accumulate in the nurse cells. This type of labeling is also observed for the *I* element in *Drosophila* ovaries (Seleme et al. 2005; Chambeyron et al. 2008). However, transcription of the *I* element was never reported in testes in this species. Our RT-qPCR analysis shows that the *I* element is transcribed in testes of both *D. mojavensis* and *D. arizonae*; this was confirmed by FISH, which showed labeling in primary and meiotic spermatocytes. In the reciprocal hybrids between the two species, the levels of expression were identical for the female germline tissues, with no difference between parents and F1 hybrids. FISH experiments also confirmed this result because no change in the localization of the transcripts and no labeling



**Fig. 7.**—Expression of *I*-like elements by RT-qPCR in ovaries (A) and testes (B) of *Drosophila mojavensis*, *D. arizonae*, and their hybrids (F for female and M for male). The transcript levels of the *I* element were estimated relative to those of the *rp49* gene. No significant differences were observed in ovaries (A). On the contrary, in testis, mRNA levels were significantly different (two-way ANOVA).

in the oocytes were observed. For the male germline, the results were quite different depending on the direction of the cross. F1 males from *D. arizonae* male parents showed no difference in the expression of the *I* element in testes. However, when *D. mojavensis* males were used as parents, a significant increase in the *I* element expression in testes was observed. The fact that F1 males in this particular cross are sterile (Ruiz et al. 1990) suggests a role for TEs in the male-sterile phenotype. Furthermore, it is important to determine whether the high expression in F1 hybrids observed in RT-qPCR is derived from complete copies of *D. mojavensis* males. Hybrid dysgenesis in *D. melanogaster* involves transposon-naïve females and active transposon-containing males in several different systems: The *I*-R system, the *P*-M system, and even the *H*-E system (Bucheton et al. 1976; Picard et al. 1978; Kidwell and Novy 1979; Kidwell 1983; Streck et al. 1986). However, at least in the *I*-R and *P*-M systems, it is the maternal line that is affected.

The presence of *I* transcripts only in the nurse cells in ovaries of *D. mojavensis*, *D. arizonae*, and their hybrids suggests that the *I* element in both species is most likely regulated by the piRNA pathway. In *Drosophila* ovaries, Chambeyron et al. (2008) showed that there is an association between the accumulation of piRNA and the regulation of the *I* element transcripts in the nurse cells, where the transcripts are processed, before being transported to the oocytes, where the retrotransposition occurs. Such findings suggest a posttranscriptional regulation of the *I* element described here before its

transportation to the oocytes, as reported in *D. melanogaster* (Brenneke et al. 2008; Chambeyron et al. 2008).

In testes, the *I* transcripts in parents and hybrids are located in primary spermatocytes, where high levels of transcription for genes encoding proteins for spermatogenesis and genes encoding male germline-specific isoforms are detected (Fuller 1998). The transcripts of many genes are stable and are present for days after meiosis but then decrease in the postmeiotic spermatids stage, during which proteins can be detected (Kuhn et al. 1988; Fuller 1998). The localization of the *I*-like transcripts observed in this study (fig. 7) corresponds to the transcriptional dynamics described for their genes. Other elements have transcripts detected in primary spermatocytes. Transcripts of 412 retrotransposon were not detectable in ovaries by in situ hybridization in *D. melanogaster* and *D. simulans* populations but were detected in the male germline in primary spermatocytes (Borie et al. 2002). Furthermore, no association has been found between expression in soma and in testes, which indicates independent regulation (Borie et al. 2002). The same localization patterns were observed for the retrotransposons 1731, *GATE*, *mdg1*, and *copia* (Haoudi et al. 1997; Kogan et al. 2003; Morozova et al. 2004; Kalmykova et al. 2005). The relatively high number of transcribed elements, including the *D. mojavensis I* element, in primary spermatocytes reveals the importance of these cells in the transcription of these TEs.

In this work, we have investigated the hypothesis that *I*-like elements (which are implicated in the hybrid dysgenic

incompatibility of *D. melanogaster*) could be activated in hybrids between *D. mojavensis* and *D. arizonae*, two recently diverged species from the *Drosophila* subgenus of the genus *Drosophila*. We showed that *I* elements are specifically activated in testes from a specific direction of the intercross, and we suggest that this activation may be associated with the observed male sterility. Moreover, we showed that this activation is sex specific because the female germline does not seem to be affected by this phenomenon. The impact of TEs in the male sterility observed in interspecific crosses is becoming clearer and needs further investigations. Studies using closely related species will be helpful to understand the first steps of the male sterility process and the role of TEs.

## Supplementary Material

Supplementary figures S1–S4 and table S1 are available at *Genome Biology and Evolution* online (<http://www.gbe.oxfordjournals.org/>).

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