

Original Article

Drosophila melanogaster Laboratory Rearing for *Wolbachia*-Based Control Programs, a Component of Dengue Control

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Abstract

Background: *Drosophila melanogaster* flies are smooth, low upkeep and safe model organisms, they can be effortlessly used in different fields of life sciences like genomics, biotechnology, genetics, disease model, and *Wolbachia*-based approaches to fight vectors and the pathogens they transmit.

Methods: Fruit fly specimens were collected in 25 districts (14 provinces) of Iran and their morphological recognition was proven by molecular analysis based on sequence homology of mitochondrial COI barcode region. Essential information and specific requirements were provided for laboratory rearing of *D. melanogaster*.

Results: *Drosophila melanogaster* colonies were found in 23 out of 25 districts. Also, five related species coincident with *D. melanogaster* were reported in this study including *D. ananassae/D. parapallidosa*, *D. hydei*, *D. repleta*, *Zaprionus indianus* (Diptera: Drosophilidae), and *Megaselia scalaris* (Diptera: Phoridae). The Iranian *D. melanogaster* molecular signature and their rearing techniques have been described here. The complete life cycle, from (egg to adult), takes approximately 8 days at 25 °C. Some biological points have been presented with highlighting capturing, rearing, culturing, and embryo collection along with primitive recognition and segregation between females and males have been presented. A recipe for culture media and the quantity of various ingredients have been provided.

Conclusion: This is the first report on the *D. repleta* and *D. ananassae/D. parapallidosa* species for the country. Results of this study provide efficient and effective rearing procedures which are requirement for both small-scale for facilitating entomological research and large-scale use in justifiable vector control management such as disease model or Dengue control.

Keywords: Fruit fly; Mass rearing; *Wolbachia*; Arboviruses; Iran

Introduction

The fruit fly *Drosophila melanogaster* (Meigen), a miniature insect nearby three mm long, was used broadly as a model organism biology through the last era and for genetics research (1). Fruit flies act for a model investigative device for exploring genetics and biology (2). Morgan received 1933 Nobel Prize for a successions of genetic studies of *Drosophila* that directed them to make the chromosome model of heredity (1).

Drosophila melanogaster fruit flies are cosmopolitan in distribution, easy to capture, easy

to produce and maintenance, low-cost, have a little life length (15–20 days), are relaxed to sex, and need very little area or special tools (3, 4). *Drosophila melanogaster* is utilized as a typical organism to study fields varying from essential genetics and disease models to the improvement of tissues and organs. There are about sixty percent homology between *Drosophila* genome and that of humans. *Drosophila* genome has few redundant region, and approximately more than 70 percent of the genes accountable for human diseases or disorders have similar

gene in fruit flies (5). These traits, together with a short production time, small maintenance costs, and the readiness of powerful genetic apparatuses, permit the fruit fly to be desirable to search difficult pathways associated in biomedical research, containing cancer (6). The full genome of *D. melanogaster* was sequenced in 2000 (7). The likelihood to subordinate the supremacy of traditional genetics with an extensive variation of molecular and cellular methods has fascinated many scientists to study on *Drosophila* in different disciplines, including cell biology, neurobiology, regulation of aging, behavior, gene expression, development, and the physiopathology of human disorders or diseases (1).

Wolbachia bacteria are best known for their capability to guard their hosts against and to inhibit the transmission of pathogens. *Wolbachia* protection against pathogens was first reported in *Drosophila*, where host flies containing *Wolbachia* had higher persistence contrasted to uninfected flies when confronted with RNA viruses (8–10) or the entomopathogenic fungus *Beauveria bassiana* (11). In mosquitoes, *Wolbachia* inhibits or interferes the reproduction and spread of human pathogens particularly arboviruses (e.g., dengue virus) and insect-specific viruses (12), rodent and human malaria parasites (13–16), and non-human animal pathogens containing the filarial nematode *Brugia pahangi* (17, 18). The model organism *D. melanogaster* naturally harboring the wMel *Wolbachia* strain which is well recognized for cytoplasmic incompatibility (CI)-induction and preventing the dengue transmission. Syncytial embryo microinjection has been employed broadly for transferring *Wolbachia* in different insect classes (19). A *Wolbachia* strain separated from *D. melanogaster* (wMel) and artificially introduced in the mosquito vector *Aedes aegypti*, is currently being used as a bio-agent to control the transmission of mosquito-borne diseases such as Zika, Dengue, and Chikungunya. wMel and wMelPop strains strongly reduce Dengue virus transmission from infected mosquitoes (20). Literature indicates

that several mechanisms might have been involved in the suppression of pathogens within vector body including struggle for an inadequate resource such as cholesterol, production of reactive oxygen species (ROS), instruction of immune genes, and subsidiary host gene regulation over other cellular machinery (siRNA, RNAi) (21).

Recently, the discovery of *Ae. aegypti* in the south of Iran caused great concern to prevent its expansion in that area and probable transmission of the diseases for which it is a vector. Approaches using *Wolbachia* for the prevention of diseases transmitted by normal and uninfected insects depend on the effective founding of constant *Wolbachia* infections, typically by embryonic microinjection of *Wolbachia*-infected cytoplasm or purified *Wolbachia* from infected insect hosts, including *D. melanogaster* (22). The success of embryonic microinjection operation requires continuous purification of *Wolbachia* from donor *D. melanogaster* and inject purified *Wolbachia* into *Ae. aegypti* recipient embryos to determine the optimal number of *Wolbachia*-infected individuals that need to be introduced into the natural population.

Transfection techniques normally employed embryonic microinjection to relocate *Wolbachia*-infected embryo homogenate or embryo cytoplasm. The embryo cytoplasm or homogenate ought to be infused at 25 °C in less than five hours (23) and it is unbearable to use embryo cytoplasm or homogenate in following injection. Therefore, laboratory rearing of *D. melanogaster* flies to take permanent access to many *Drosophila* embryos (early embryos will be collected every 30 min) for the purification of *Wolbachia* bacteria or direct transfer of *Wolbachia*-infected embryo cytoplasm in *Aedes* embryonic microinjection operation is highly needed.

For these reasons, providing laboratory rearing of fruit flies of *D. melanogaster* makes a perfect model organism for rearing in our research laboratory. This study was conducted to

provide basic biological information and rearing in laboratory conditions associated *D. melanogaster* flies from Iran.

Materials and Methods

Study area: Field collections of *Drosophila*

We have tried to collect fruit fly specimens from a wide range of geographical regions (25 districts, 14 provinces) throughout the country. Adult fruit flies were collected from different habitats: human dwellings, jungle, and riverside, using the methods described in Hoffmann et al. (24).

Fruit flies were collected by retaining containers with banana in the stands and netting the adults afterwards a few hours, or by collecting adult fruit flies immediately from fallen fruit. The collected specimens were transferred to the Insect Molecular Biology Laboratory in the School of Public Health, Tehran University of Medical Sciences, Tehran.

Species Identification

Adult fruit fly specimens were recognized to species level using a standard morphological key (25). Moreover, PCR-directed sequencing of mitochondrial DNA cytochrome c oxidase subunit 1 (barcoding region) was used to confirm morphological identification for a subset of specimens, including the unknown or suspected specimens that had lost crucial identifying morphological characteristics. Collins et al. (1987) method was used to extract genomic DNA (26). A 710 bp barcoding region of the mtDNA-COI gene was amplified by polymerase chain reaction (PCR), using the following primers: LCO 1490 (5'-GGTCAACAAATCA TAAAGATATTGG-3') and HCO 2198 (5'-AACTTCAGGGTGACCAAAAATCA-3') (27). The PCR reaction was performed in a overall volume of 25 μ L using the Taq DNA Polymerase 2xMaster Mix RED, Ampliqon (Denmark), with the following reagents: 12.5 μ L of Master mix, 1 μ L of each primer (10 mM), 10–20 ng (1–2 μ L) of DNA extract, and 8.5–9.5 μ L of double distilled water. The PCR ther-

mal program was 5 minutes at 94 °C, followed by 30 cycles of 45s at 94 °C, 30s at 50 °C, and 1 minute at 72 °C, with a final extension at 72 °C for 10 minutes. PCR amplicons were assessed by 1% agarose gel electrophoresis, followed by GreenViewer staining and visualizing using a UV transilluminator. Sharp and cleaned amplicons were sequenced bidirectionally by the Genetic Codon Company, Tehran, Iran, using the same primers as for PCR amplification. The quality of raw sequences was enhanced using the Chromas 2.6.5 program, by deleting regions with poor quality at both ends of the sequences. The consensus of sure sequences was inspected using the NCBI (nucleotide collection) database package (28).

Life Cycle and some bionomic characters of *D. melanogaster*

Drosophila belongs to dipteran insects with holometabolous metamorphosis and has distinct stages of development including egg, larva, pupa and adult. Duration of these stages can be affected by the temperature (2). *Drosophila melanogaster* flies were reared in 500- and 1000 mL plastic culture bottles with cotton plugs. They were maintained at room temperature of about 25 °C during the entire experiment. Flies were monitored for development events, and average durations of different stages in their life.

Sexing of fruit fly specimens at pupation and adult stages was performed using morphological characters (29). The criteria for identification virgin specimens were explained according to the meconium and or the time passed after adult emergence.

Culture medium preparation:

The materials and recipe to prepare culture medium for fruit flies was provided in this study. Generally, agar powder, wheat flour, sugar, yeast (dried), and propionic acid are necessary for the medium.

Results

Drosophila melanogaster distribution in Iran

In total, 1273 fruit flies were collected from the study areas (25 districts of the country) and identified to species level using morphological characteristics and or COI barcoding sequences, through comparison with Genbank published sequences. Most of the flies found in this study belonged to *D. melanogaster* (21 out of 25 colonies) (Table 1). In addition to *D. melanogaster*, five other drosophilae or related species including *D. ananassae* (Doleschall) or *D. parapallidosa* (Tobari), *D. hydei* (Sturtevant), *D. repleta* (Wollaston), *Zaprionus indianus* (Gupta), and *Megaselia scalaris* (Loew) were found respectively from Chalous-Pakdasht, Tabriz, Chabahar, Kerman, Mashhad, and Ramsar locations (Fig. 1). This is the first report of the species *D. repleta* and *D. ananassae* (or *D. parapallidosa*) in the country.

D. melanogaster sexing

Sexing according to the adult feature

Drosophila melanogaster is sexually dimorphous and the sexes can be differentiated by the following morphological characters as already explained by (29): males are slightly smaller than females. Sexing of fruit flies is performed based on the variations in the magnitude of the females and males (Fig. 2A). In females, tip of the abdomen is pointed whereas it is somewhat rounded in males (Fig. 2A). There are eight segments in females' abdomen, with seven being obviously noticeable whereas the final three abdominal segments are merged in males, so that only five segments are visibly detectable (Fig. 2A). In *D. melanogaster* fruit fly, and numerous other *Drosophila* species, the males' abdominal tip is opaquer than the ones in females (Fig. 2A). However, the variation in abdominal pigmentation is not true in the newly emerged flies. The female and male genitalia, evident on the ventral face of abdominal tip, are clearly different, with the male genital plate showing darker and harder. The males possess a sex comb (Fig. 2B) as a row of around ten black, stable

bristles on the first tarsal segment of front legs. In disparity, female specimens have rather pointed posterior ends with no sex combs. Moreover, females and males can also be recognized in the pupal stage by inspecting their forelegs. In the last pupal phase, male pupae can be recognized by the sex-comb which are black hairy tufts exhibit on the first tarsal segment of foreleg and obvious over the pupal case (Fig. 2C).

D. melanogaster Virginity

Virginity according to adult eclosion time

In fruit fly experiments particularly genetic studies, it is important that female be a virgin to determine genetic background of progeny. A female *Drosophila* can store and use the sperm from a single insemination (33). Thus, it is necessary to select virgin females for genetic crosses (34). Female fruit fly does not couple within the first 8–12 h after eclosion from the pupae (1). In this study female flies mate 4–6 h after eclosion at 25 °C. Therefore, we suggest females to be separated from the males before 4 h post emerging and kept separately to use once needed.

Virginity according to meconium

In the initial hours after adult emergence, young virgin adults are distinguishing from older adults as their bodies will be very light in coloration and they are very light with a unique dark greenish spot obvious underneath the abdomen cuticle. This spot is known as meconium, which is the remnants of their last food prior to pupating. The adult flies expel meconium from the body in a few hours after eclosion. Adults having meconium may be consistently considered as virgins yet in a collection that has not been separated within the 6 h gap (Fig. 3). We found the freshly emerged adult fruit flies, sooner less than 6 h post eclosion.

Selecting virgin in pupal stage

In addition to the methods mentioned above, researcher can separate males and females in pupal stage. For this purpose, we used a fine brush to select and separate mature and dark-

ened pupae. Mature pupae are reserved on a glass slide followed by examining their ventral side under a stereo-binocular microscope. Their legs can be seen via semi-transparent pupal case. Female's pupa does not have the sex comb on their front legs (Fig. 2C) and can be transferred to a separate food vial.

Culture media for fruit fly maintenance in laboratory

Rearing media preparation

An exclusive recipe for preparation of fly food has been prepared in our laboratory. The necessary materials include agar powder 40–60 gr, wheat flour 300 gr, sugar 300 gr, yeast (dried) 200 g, propionic acid 25 mL, and distilled water 5 liter. Firstly, the yeast, sugar and wheat flour were mixed in cold water and then boiled on heater, then slowly added the agar powder to the medium till agar is completely dissolved. The mixture was then cooked on heater for 7–10 minutes. Once the medium temperature reaches close to 50 degrees centigrade, propionic acid is added and mixed well. Throughout the preparation, the culture should be continually blended. The cooked food was poured in bottles (around 30–40 mL each) and allowed it to harden as it cools down. Afterwards 4–5 h, any damp deposited on the bottle's walls wiped dry with fiber and plugged with cotton. The food is ready to use after one day. The bottles could furthermore be kept in a refrigerator for 1–2 weeks for forthcoming use.

Media for *Drosophila* embryo collection

Wolbachia-infected *D. melanogaster* is being used in transfection studies for Dengue control program. Flies will kept at 25 °C using normal *Drosophila* rearing settings (35). Microinjection experiments (Transfection of *Wolbachia* bacteria to *Aedes* embryos), require collection of relatively large number of *D. melanogaster* (donor of *Wolbachia*) early embryos in a limited time (every 30 min) (23). In general, embryos are gathered in Petri dishes encompassing *Drosophila* culture, because of the assist with which they can be collected from the food

surface. For this reason, the media has been poured (10 mL) in Petri dishes, then about 150 adult flies that already were numbered in freezer for few minutes were released in Petri dish (Fig. 4A). The lid of Petri dishes was sealed with adhesive tape to avoid escaping flies. After adults' egg laying (Fig. 4B), *Drosophila* embryo (Fig. 4C).

Drosophila melanogaster Life Cycle

Four discrete stages exist in the life cycle of *D. melanogaster* fruit fly: egg (Fig. 5A), larva (Fig. 5B), pupa (Fig. 5C) and adult (Fig. 5D). Eggs of *D. melanogaster* are white in color and are approximately 0.5 mm in length with two dorsal appendages (Fig. 5A). This two egg threads or dorsal projections develops from its anterior end to avoid egg from sinking into soft breeding places. A fresh culture of *D. melanogaster* can produce new adults in 9 days at 25 °C: 5 days in the egg and larval phases, and 4 days in the pupal period. Principally it should be noted that higher temperature resulted in a shorter life cycle. In this study condition at 25 °C, eggs hatched after 24 h, and the 1st tiny instar larva appeared. The larval phases comprise of three mobile steps or instar larvae, the first (L1), second (L2) and third (L3) instars. After 24 h, the 2nd instar larvae appeared and turned into 3rd instar larvae after 24 hours. The first, second and initial third instar larvae mostly persist inside the food media, aggressively feed, and raise their body mass. Late third instar larvae get out of food and scramble in seek of a dry position for pupation, which is commonly the wall bottle or the upper cover of Petri dish, where they are sustained in laboratory.

Late third instar larvae stick at dry positions using the adhesive protein produced by their salivary glands. For four days pupa completed their development cycle from white pupa (which in the commencement is white and soft), brown pupa (harder cuticle and duskier in color because of sclerotization and chitinization), mid pupa and late pupa with eye pigments which they experience the final molt to begin the pupal phase.

At the end of pupal stage, the imago opens by driving its way over the anterior end of the pupal case (Fig. 6A). The newly emerged *D. melanogaster* fly is lengthened, igniter in color, and with folded wings with visible meco-

nium (Fig 6B and 6C). In the next 2–3 h, the wings open out; the body blackens and reaches the typical adult body form while the meconium pass away (Fig. 6D). Female flies get ready to couple in 4–6 h after emergence.

Table 1. The distribution of *Drosophila* spp. and related species in Iran, 2021

No	Origin, location	Direction	Fly species	GenBank ID Number	Closest Entry in Genbank, (Country, Similarity rate)	Reference
1	Amol	North	<i>D. melanogaster</i>	NA	NA	NA
2	Chalus	North	<i>D. melanogaster</i> <i>D. hydei</i>	NA OR077700	NA KJ463782 (USA, 100) KJ671603 (New Zealand, 100)	NA DS DS
3	Ramsar	North	<i>D. ananassae</i> or <i>D. parapallidosa</i>	OR077701	MK659805 (NI, 99.85) MK659836 (NI, 99.85)	DS DS
4	Salmanshahr	North	<i>D. melanogaster</i>	NA	NA	NA
5	Qaemshahr	North	<i>D. melanogaster</i>	NA	NA	NA
6	Tehran	Center	<i>D. melanogaster</i>	NA	NA	NA
7	karaj	Center	<i>D. melanogaster</i>	NA	NA	NA
8	Pakdasht	Center	<i>D. melanogaster</i> <i>D. hydei</i>	NA OR077699	NA KJ671603 (New Zealand 100) KJ463782 (USA, 100)	NA (30) (31) NA
9	Isfahan	Center	<i>D. melanogaster</i>	NA	NA	NA
10	Kashan	Center	<i>D. melanogaster</i>	NA	NA	NA
11	Shahrekourd	Center	<i>D. melanogaster</i>	NA	NA	NA
12	Kerman	Center	<i>Zaprionus indianus</i>	OR077697	EF632366 (India, 99.69) EF632355 (Benin, 99.69)	DS DS
13	Bojnurd	Northeast	<i>D. melanogaster</i>	NA	NA	NA
14	Mashhad	Northeast	<i>D. melanogaster</i> <i>Megaselia scalaris</i> (KM88*) <i>Megaselia scalaris</i> (PM33*)	NA OR077703 OR077704	NA MT396285 (S.Korea, 99.55) AB907181 (India, 99.39) KU949579 (Iran, 99.18) JQ941745 (China, 99.70) MT396354 (S.Korea, 99.54)	NA (32) DS (32) DS (32)
15	Urmia	Northwest	<i>D. melanogaster</i>	NA	NA	NA
16	Darreh-ye Qasemlu	Northwest	<i>D. melanogaster</i>	NA	NA	NA
17	Miandoab	Northwest	<i>D. melanogaster</i>	NA	NA	NA
18	Tabriz	Northwest	<i>D. repleta</i>	OR077702	MG605140 (USA, 98.78) MG605139 (USA, 98.78)	DS DS
19	Sanandaj	West	<i>D. melanogaster</i>	NA	NA	NA
20	lorestan	West	<i>D. melanogaster</i>	NA	NA	NA
21	Shiraz	South	<i>D. melanogaster</i>	NA	NA	NA
22	Ahvaz	Southwest	<i>D. melanogaster</i>	NA	NA	NA
23	Dezful	Southwest	<i>D. melanogaster</i>	OR077696	KY559390 (Finland, 99.69) KJ767245 (Uzbekistan, 99.69)	DS DS
24	Zahedan	Southeast	<i>D. melanogaster</i>	OR077695	KY559390 (Finland, 100) KJ767245 (Uzbekistan, 100)	DS DS
25	Chabahar	Southeast	<i>Zaprionus indianus</i>	OR077698	EF632366 (India, 99.54) EF632355 (Benin, 99.54)	DS DS

NA: not indicated, DS: direct submission, *: Voucher specimen



Fig. 1. Adult specimens of five fruit flies or related species to *Drosophila melanogaster* found in the study, Iran, 2021
 a: *Zaprionus indianus*, b: *Drosophila ananassae*, c: *D. heydei*, d: *D. repleta*, e: *Megaselia scalaris* (Original photos)

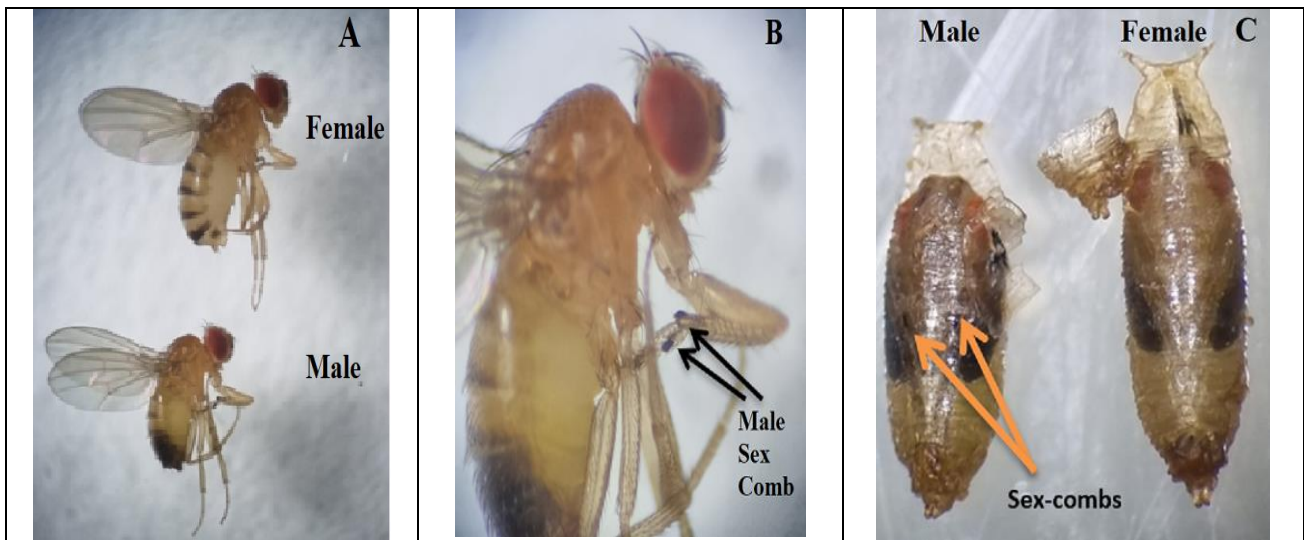


Fig. 2. Sexual dimorphism in *Drosophila melanogaster* adults in size, shape, segments, and color of abdomen (A), presence of sex comb in adult male prolegs (B) and pupal stage (C) (Original photos)

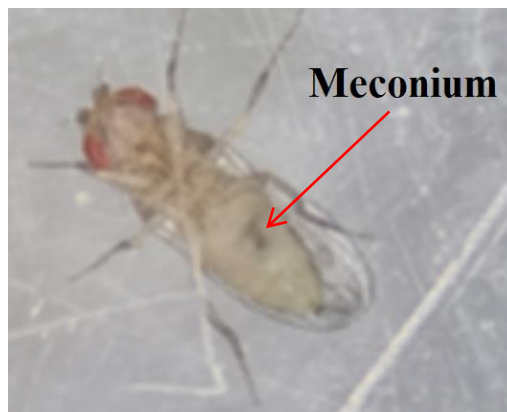


Fig. 3. Meconium on virgin adult *Drosophila melanogaster* (Original photo)

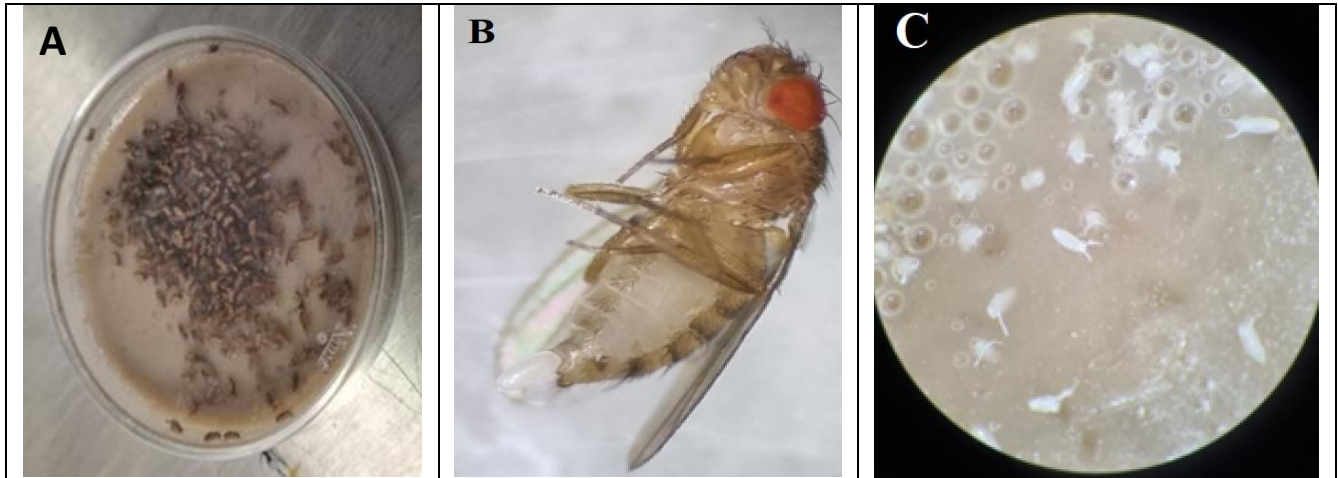


Fig. 4. *Drosophila melanogaster* culturing in Petri dish (A), female' egg laying (B) and *D. melanogaster*'s eggs (embryos) in media culture (Original photos)

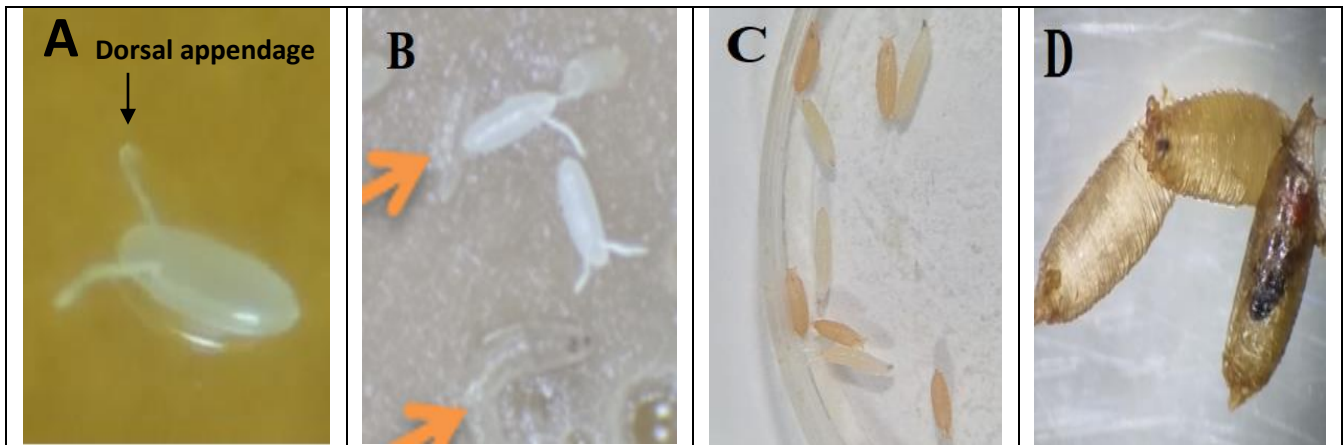


Fig. 5. *Drosophila melanogaster* egg with dorsal appendages (A), first instar larva (B), third instar larva and brown pupal stage (C) and white, brown and late pupa (D) (Original photos)

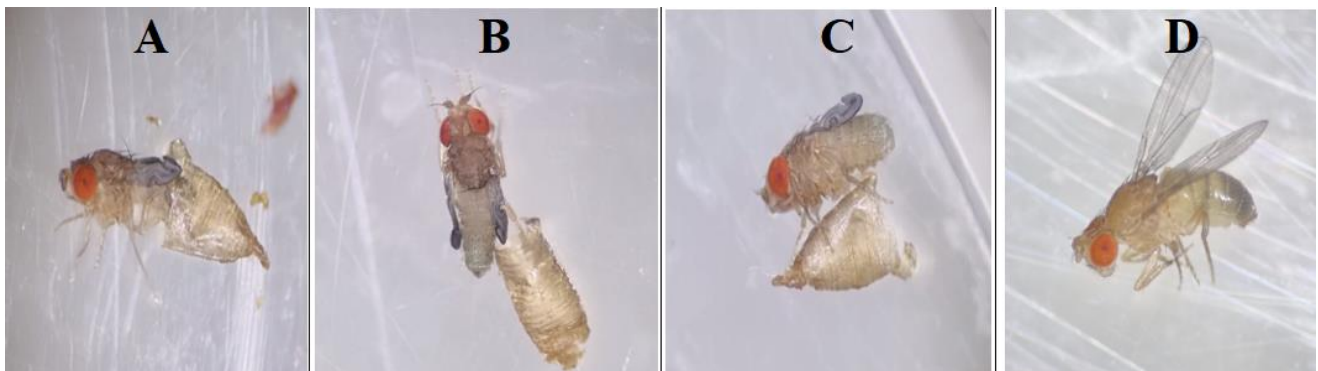


Fig. 6. *Drosophila melanogaster* adult eclosion stages: pupal case (imago) ecloses by imposing its way over the anterior end of the imago (A) freshly emerged adult, with light color and folded wings (B and C), adult with expanded wings, darkened body, and attained the typical adult body shape (D) (Original photos)

Discussin

Despite the extensive use of *Drosophila* in biological and genetic research, the fruit fly fauna of Iran is inadequately studied. The family Drosophilidae is an extremely diverse, universal family of flies that contains fruit flies and pomace or vinegar flies (36). This is the first survey to investigate fruit fly specimens' distribution from a wide range of geographical regions (25 districts, 14 provinces) throughout the country.

Previous research on *Drosophila* species in the province of West Azarbaijan recorded ten drosophilid species (37), which were all new to the west of Iran (except for *D. melanogaster*). Interestingly, one drosophilid genus *Lordiphosa* Basden and three species *Scaptodrosophila lebanonensis* (Wheeler), *Lordiphosa andalusiaca* (Strobl), and *D. hydei* (Sturtevant), were reported for the first time in Iran. In addition, the species *Zaprionus indianus* (Gupta), *D. simulans* Sturtevant, *D. phalerata* Meigen, *D. subobscura* Collin, *Scaptomyza flava* (Fallén), and *S. pallida* (Zetterstedt), were recorded in the study area.

In the current study most of the fruit flies were *D. melanogaster* (21 colonies), while *D. ananassae* (or *D. parapallidosa*) specimens were found in only one location, Ramsar in the North of the country. Although we also found specimens belonging to species like *M. scalaris*, *D. hydei*, *D. repleta*, and *Z. indianus* (Fig. 1) respectively from Mashhad, Chalous-Pakdasht, Tabriz, Chabahar-Kerman. In 2015 the first report of urogenital myiasis due to *M. scalaris* in Iran was published (38). Also, this species was found on human cadavers in Tehran District (33).

In this study we used the mtDNA-COI gene for species identification of fruit flies. This gene has been displayed to be a powerful molecular marker to identify different insects at species or even population level. Several studies have assessed the effectiveness of COI gene and proved to be successful for many groups of invertebrates, such as, flies, mosquitoes, cockroaches, and black flies (33, 39–47).

This is the first report on the described *D. repleta* (Patterson) and *D. ananassae* (or *D. parapallidosa*) species for the country. *Drosophila repleta*, frequently referred to as dark-eyed fruit fly or dark-eyed vinegar fly, is a synanthropic species of Nearctic-Neotropical origin (48). The larvae of this species forage on yeast in putrefying and fermenting organic matter (49). Because of this ecological niche, these fruit flies have turned into pests in various settings where food debris exists. In the agricultural setting, *D. repleta* fruit fly can be an annoyance pest in animal and poultry facilities (50). *Drosophila repleta* has been shown to scatter from pit toilets into nearby residences from around 305 m away (51). Spreading of flies from animal facilities to local houses may be of some trouble because fruit fly species can transfer pathogens (52). Black et al. (2018) showed that *D. repleta* is a carrier of food-borne pathogens containing *Salmonella* Saint Paul, *Escherichia coli*, and *Listeria innocua* onto human food (53).

Drosophila hydei belongs to subgroup of the repleta species group (54). Strangely, this fruit fly is known to have a long sperm proximately 23 mm, ten times the size of the male's body (55). *Drosophila hydei* are frequently observed on dung piles globally and are one of the more usual flies used as feeders in the domestic animal market. Flightless forms of this species are reported (56).

The COI sequence analysis of fruit fly specimens in Ramsar, northern Iran, revealed a similar sequence homology (99.85%) to either *D. ananassae* or *D. parapallidosa* species. These two sister taxa are new for Iran, belong to the subgenus *Sophophora*, melanogaster group, ananassae subgroup, and the *D. ananassae* species complex. Like *D. melanogaster*, *D. ananassae* is a useful model organism for genetic studies and is among the 12 fruit fly species that its genome was sequenced (57). *Drosophila ananassae* is a human-commensal species scattered in subtropical and tropical regions worldwide,

and it contains a number of sibling or cryptic species in Asia-Oceania (58–61). This is suggested that *D. parapallidosa* has recently moved northward from a tropical (hot and humid) to a subtropical region and has happened to sympatric with *D. ananassae* there. It is proposed that *D. parapallidosa* is constantly sympatric with *D. ananassae*, but not the other way around (62).

Zaprionus indianus was another fruit fly species reported in this study. *Zaprionus indianus* was reported for the first time from western Iran (Zirtang-e Siab in Lorestan Province) (63) according to BLAST analysis of COI sequences. This species is known as an invasive species, also recognized as the striped vinegar fly in the United State of America and the African fig fly in South America. This is a fig pest in the new- and old-world countries, and it could cause economic lose in the areas they are present (64). The distribution and its possible stable establishing in Iran especially in fig farms should thus be monitored in the future to evaluate its damages.

Drosophila fruit flies are holometabolous insects and four development stages: egg, larva, pupa, and adult can be seen in their life cycle. In laboratories, *D. melanogaster* is usually maintained at 25 or 18 °C (the lower temperature is used principally for keeping stocks) (1); we prepare 25 °C for all the timing. The generation time in 25 °C took roughly 9 days from fertilized egg to freshly emerge adult. At 25 °C a new and fresh culture of *D. melanogaster* produced new adults in nine days: five days in the egg plus larval phases, and four days in the pupal stage. Flagg reported that at 21°C a fresh culture of *D. melanogaster* could create new adults in two weeks; eight days in the prepupal stages, and six days in the pupal phase (34). It is known that development time of immature stages of insects is affected by the temperature (65–67).

Maintenance of *D. melanogaster* fly colonies in laboratory needs permanent supervision and attention to guarantee that flies are fit and

there is an optimal condition inside the culture. Density is one of the important biotic factors in laboratory rearing of insects. Over-crowding is a frequent trouble as fruit flies continue laying eggs and increase their amounts within the restrained area and resources accessible in the bottle/vial (29), younger larvae consume aggressively and mainly stay interior the diet until they prime for pupation. Nevertheless, overpopulation imposes much the young larvae (1st and 2nd instar larvae) to scramble out of the food stuff. The *D. melanogaster* flies must be relocated repeatedly to fresh bottles/vials with diet because rotting food and fly remains charm infecting pathogens like mites, fungi, and bacteria. In the existing work we relocated *D. melanogaster* flies repeatedly to new containers within 25–30 days. Establishing a new medium with around 10–20 flies in containers and less than fifty fruit flies in jugs or bottles prevents overcrowding and asserts a vigorous culture.

To date, the ectoparasitic mite *Macrocheles subbadius* and Allantonematid nematodes linked with mycophagous fruit flies are known as parasite interaction with drosophilid colonies (68). We have maintained a *D. melanogaster* colony in the laboratory since 2017. Mites are a main trouble in any insect laboratory colonies. These tiny animals can be simply noticed in aged colonies as mites walk surrounding blank pupal cases. They put a pearly feature of 10–20 eggs near pupae. Mites cause a significant risk to the developing colonies. In addition to feeding lifeless flies, mites similarly damage vital pupae and embryos, causing devastation of the fly colony. During these few years, except for some cases of nematode contamination we didn't have any other kinds of major infections in *Drosophila* cultures. The source of contamination was discarded in those cases.

Historically, fruit fly colonies in laboratories have been kept on bananas. However, since banana converts very dim and hard to process, agar-based culturing medium using wheat cream or maize powder, has suited universal. Several

culture media have been introduced, all built on simple ingredients such as sucrose, propionic acid, yeast (not yeast extract), and agar (1). Some media encompassing a diverse additives comprising rice, raisins, oat hulls, pears, and molasses have been recorded (4). A few of the procedures applied for grounding of fruit fly diet in diverse laboratories are (28): 1: Agar-Maize powder-sugar food, 2: Corn meal-Agar medium, 3: Wheat Cream-Jaggery Medium, 4: Corn flour-Jaggery Medium, 5: Banana- Barley-jaggery medium. In current study culture media based on Agar powder, wheat flour, sugar, yeast (dried), propionic acid, and distilled water were used for maintenance of flies in laboratory. All fly cultures in the food media which are prepared in this way can be stored in the freezer for up to 3 months in sterilized bottles.

Conclusion

This is the first report of the species *D. repleta* and *D. ananassae/D. parapallidosa* in the country. Further monitoring and investigation is suggested for the fruit fly species that can carry pathogens, food-borne diseases, or act as agricultural pests. Results of this study provide effective and efficient rearing information and recipe which are important for both minor entomological investigation and large-scale exploitation in maintainable vector control management such as disease model or Dengue control.

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Ethical consideration

The protocols were conducted in this study followed the guidelines of the institutional ethical committee (Tehran University of Medical Sciences, TUMS). The protocols were approved by TUMS ethical committee under registry IR.TUMS.SPH.REC.1399.176.

Conflict of interest statement

The authors declare there is no conflict of interests.

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