**Declines In Chill Coma Recovery and Negative Geotaxis Due To an Entamopathogen in *Drosophila melanogaster* the Fruit Flies**

Usha Bhocal ¹, Bijender Singh Sabharwal ² and Veer Bhan ³

¹Research Scholar, Department of Biotechnology, University Institute of Engineering and Technology, Maharshi Dayanand University, Rohtak, Haryana, India
²P.G. Student, Department of Law, Maharshi Dayanand University, Rohtak, Haryana, India
³Assistant Professor, Department of Biotechnology, University Institute of Engineering and Technology, Maharshi Dayanand University, Rohtak-124001 Haryana, India

**Abstract:** Infection related studies in *Drosophila melanogaster* provide insight into both mechanisms of host resistance and pathogens tolerance. The advantages of the fly as an experimental system include its genetic tractability, short life span and the observable possibility and to quantitatively analyse behavioral responses. At present, research has been limited by the relatively few metrics that can be used to measure health and illness during the course of infection. Here by, we explain the pleiotropic measurements of infection-related declines in flies performance on two different assays. *D. melanogaster* are somewhat slower to recover from an induced chill coma after fungal infection with *Beauveria bassiana*. Due to fungal infection, flies performance also gets affected during a negative geotaxis assay, revealing a decrease in their behavioral response. In addition, to provide new advancements for assessing health, these non-lethal assays also suggest many pathological consequences and also up and downs of metabolic that may occur over the entire course of an infection.

**Keywords:** Chill coma, *Drosophila*, geotaxis, infection, resistance

**I. Introduction**

All organisms need to defend themselves against pathogens, parasites and other natural enemies, and investment in defense is a significant component of the life history strategy of most animals and plants [1,2]. Infection can impact the health of an organism in complex ways. Direct damage to the host caused by pathogenic toxins, damage to host tissue resulting from immune effectors, and the energetic expense of responding to an infection are all known to affect the health of a host [3,4,5,6]. Specifically, *D. melanogaster* possess thermo sensitivity (the ability to detect temperature change), allowing individuals to seek out desirable habitat [7,8] note that the strength of thermo sensitivity a species elicits is dependent on the species surrounding environment. Surviving an infection requires both resistance, the ability to limit pathogen burden, and tolerance, the ability to minimize the impact a pathogen has on fitness [3,9,10]. Also, insects have evolved a range of molecular adaptations to cope with seasonal exposure to stressful (high or low) temperatures [11].

Understanding the various pathways mediating resistance and tolerance allows for better development of interventions that focus on the maintenance of health throughout an infection. However, dissecting the mechanisms that determine the balance of eliminating pathogens and the damage and energetic cost of mounting that immunological response requires metrics that move beyond survival and allow assessment of health during infection.

Work with the model organism *Drosophila melanogaster* has provided important insights into the roles of different molecular pathways involved in immunity[12,13,14]. *D. melanogaster* provides a genetically tractable system where a large numbers of replicates can be assessed while still being a whole organism system with biological complexity. Additionally, a relationship between infection, immunity, and reproductive fitness in insects has been well established. In environments where nutrients are limited, there is a negative correlation between female fecundity and resistance to bacterial infection in *D. melanogaster* [15]. Additionally, immune challenged females not only have fewer offspring, but those offspring also have shorter lifespans compared to the offspring of unchallenged female *D. melanogaster* [16]. The molecular mechanisms behind recovery from cold stress are complex and it seems that more genes/proteins are activated during the recovery phases than during the period of the cold stress itself [17,18].

Here we report on two distinct assays that are not measures of natural declines in infection, but rather measure the fly’s ability to recover from stress or react to stimuli. Such assays are commonly used in the field of aging research in *D. melanogaster*, which has long been recognized that measures of both health span and longevity are critical to understanding the biology of aging. We hypothesized that these metrics could also be used to assess health during infection because *D. melanogaster* shows an age-related up-regulation of inflammatory genes and expression patterns that characterize aging and the induction of an immune response in
these animals are related [19]. Additionally, in aged flies as well, the wasting and loss of circadian rhythms reported to occur during infection [20, 21].

Here we show infection-related deficits for the two behavioral assays, chill coma recovery and negative geotaxis. When low temperature exposures were given to insects, they enter into a reversible period of immobility referred to as chill coma. While the physiological causes behind induction and recovery from this coma are incompletely understood, the duration of time taken by D. melanogaster to be able to stand after returning to the warmer temperature is altered by the animal’s environment prior to cold exposure, its energy accumulation sites, and its age-related consequences [22, 23, 24, 25]. Negative geotaxis is the measure of how quickly a fly is able to vertically climb after being tapped to the bottom of a vessel as part of its innate escape response. Negative geotaxis is measured by either the climbed distance in a set time or the length of time it takes an animal to climb a set distance. Negative geotactic ability has been shown to be sensitive to oxidative stress, age-related parameters, and previous exposure of cold, but not to fungal infection [26, 27, 28, 29]. We finally hypothesize that the changes in physiological parameters that cause performance deficits in both of these pleiotropic assays are somehow affected by fungal infections, such that both assays can be used during fungal infection to detect decreases in different health consequences.

II. Materials and Methods

2.1 Collection and maintenance of flies

Drosophila species individuals will be collected along an altitudinal range (219m to 2202m) in a single planned collection trip to avoid seasonal variations. Before collection, climatic and geographical data of 8 - 10 sites will be analyzed and 4.5 sites, which will be showing a clinal trend of T\text{c} and RH\text{c}, will be selected. Temperature of collection site will be noted down at the time of collection and the methodology of experiment will be followed according to the temperature of collection site. Large and random samples will be made i.e. 3 or 4 places from same site. Agar-sugar food vials will be used for collection. Trap bait method and net sweeping method will be used to procure a maximum number of flies of a particular species as well as different species with minimum expenditure of time and material.

After identification of the naturally collected flies, males and females will be separated and isofemale lines will be established with the wild caught females. Ten isofemale lines in three replicates per population will be established. The experiments will be performed at 25°C, 12 h L: D in a humidified incubator.

2.2 Preparation of fungal material

B. bassiana strain will be stored as a stock spore suspension (105 spores per ml) in 25% glycerol at −80°C. Cultures will be generated from this stock by spreading 5μl of the solution onto Petri dishes with Potato dextrose agar (PDA) and culturing for 3–4 weeks at 29±0.5°C. These plates will be then used to culture the next set of Potato dextrose agar plates (inoculated by streaking with a nichrome wire loop), which when mature (after 3–4 weeks) will be used in the experiments. Before use, to make sure the fungus colonies are mature and sporulating, samples from all plates will be examined under a microscope.

2.3 Infection mode

Natural infection by entomopathogenic fungi. Anesthetized flies were shaken for a few minutes in a petri dish containing a sporulating fungal culture. Flies covered by spores were then removed to fresh Drosophila medium and incubated at 25°C. Fungi are important insect pathogens and spores of some entomopathogenic species such as B. bassiana can sporulate on the cuticle and produce chitinases and proteases to penetrate the insect.

In this study, we infected Drosophila melanogaster adults on sex basis aged 3-4 days at 25°C. Natural infection was initiated by shaking anesthetized flies in a petridish containing a sporulating culture of Beauveria bassiana was incubated at 25°C for required time period.

2.4 Chill Coma Recovery

Every day a new group of infected and control flies were transferred to clean glass vials (50 flies per vial) without anesthesia around noon and placed for three hours in melting ice in a bucket to induce chill coma. After that in a large petri plate flies were kept. A timer was started once the flies were placed in petriplates and only those flies were considered recovered only when they were able to stand on their legs. It should be noted that a single set of flies was not used throughout the course of infection experiment due to the reported effect of exposure to cold on longevity and immunity [30, 31]. Calculation is done by taking mean and standard deviation respectively.
2.5 Negative Geotaxis

Negative geotaxis is a frequently used index of locomotor behavior in flies [32,28,33,34,35,36,37,38,39,40]. Additionally, the negative geotaxis assay was performed in a manner similar to that published by [39]. 50 infected and 50 control flies were transferred once a day in the early afternoon without anesthesia into 10 cm tall clean glass vials. The glass vials were placed in a rectangular frame to keep them upright and put in front of a light box at room temperature in a chamber kept between 50% and 60% humidity. Waited for ten minutes so that all flies after transfer become fully awake and active. Individuals that did not become active were removed after waiting for an additional five minutes. After the vial was tapped three times and climbing time was recorded by a stop watch as 10 sec, 30 sec, 60 sec post startle. All work was done on sex basis. Flies were then transferred without anesthesia back into their rearing vial.

III. Results

3.1 Infection impairs chill coma recovery

Beauveria bassiana infection impairs chill coma recovery. Experiments were done on D. melanogaster that have previously been reported to undergo infection-related changes in circadian rhythms, display sickness induced anorexia, and have revealed different infection-dependent and background-dependent resistance and tolerance mechanism [41,42 unpublished observation]. To measure chill coma recovery, flies were placed on ice for three hours to induce a chill coma. Flies were then returned to room temperature and monitored for recovery as determined by their ability to stand on their legs. Both fungal infected flies showed delay in time to recover. Male flies recovered late in control as well as in infected flies.

![Figure 1](image1.png)

**Figure 1** Effect of B. bassiana infection on the mean CCRT (chill coma recovery time) of D. melanogaster collected from three different sites (Rohtak, Solan and Shimla) at 25°C. Values are expressed as mean ± SD (N=50).

3.2 Infection declines in negative geotaxis

Flies showed declines in the height climbed at 10 sec, 30 sec and 60 sec post-startle during infection as compared to control. Comparison of the data by infected versus control flies showed overall significant differences. After infection flies locomotory ability was decreased and they took more time to cover the required distance.

![Figure 2(a)](image2.png)

**Figure 2(a)** B. bassiana infection impairs climbing ability of D. melanogaster of Shimla female flies, time interval in seconds (10sec, 30 sec and 60 sec), (N=50)
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Figure 2(b)  *B. bassiana* infection impairs climbing ability of *D. melanogaster* of Shimla male flies, time interval in seconds (10 sec, 30 sec and 60 sec), (N=50)

Figure 2(c)  *B. bassiana* infection impairs climbing ability of *D. melanogaster* of Solan female flies, time interval in seconds (10 sec, 30 sec and 60 sec), (N=50)

Figure 2(d)  *B. bassiana* infection impairs climbing ability of *D. melanogaster* of Solan male flies, time interval in seconds (10 sec, 30 sec and 60 sec), (N=50)
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IV. Discussion

Chill coma temperature for cold tolerance and distribution has been used in many studies and has been shown to be a good proxy. Earlier methods generally relied on behavioural criteria, defining chill-coma temperature at which the animals became motionless, unresponsive to stimulation, or fell over because they had lost control over the righting reflex. *Drosophila melanogaster*, the fruit fly, has been used to study molecular mechanisms of a wide range of human diseases such as cancer, cardiovascular disease and various neurological diseases [43]. These behavioral assays are widely applicable for studying the role of genetic and environmental factors on fly behavior. Here we demonstrate a significant impairment in chill coma recovery during *Beauveria bassiana* infection. Chill coma recovery is a process that is known to have both genetic and environmental influences [44].

In certain insects including *D. melanogaster*, it has been shown that expression of immune response genes are increased in response to long term or repeated exposure to cold suggesting potential shared responses to cold and infection [45,46]. Similarly, the genes that are up-regulated during infection were also resulted in up-regulation of more general stress response genes after chill coma recovery assay. The heat shock proteins, particularly Hsp70, when upregulated is known to be of functional importance in response to both stresses [47,48,49], suggesting some shared protective mechanisms. The Frost gene likewise plays a significant role in

**Figure 2(e)** *B. bassiana* infection impairs climbing ability of *D. melanogaster* of Rohtak female flies, time interval in seconds (10sec, 30 sec and 60 sec), (N=50)

**Figure 2(f)** *B. bassiana* infection impairs climbing ability of *D. melanogaster* of Rohtak male flies, time interval in seconds (10sec, 30 sec and 60 sec), (N=50)
chill coma recovery [47] and is known to be up-regulated during infection [50]. These may highlight the importance of protection against cellular damage to survival of each of these stresses. The combined impact of chill and infections may overwhelm the protective mechanisms of these flies, resulting in delayed chill coma recovery or even damage resulting in death. It is also likely that relationship between the physiology of infection and delayed chill coma recovery is due to the metabolic changes that occur during infection. Mounting an immune response is known to have an energetic cost, hypothesized to result in trade offs between immunity and response to other stresses [51, 52, 53]. Increase in chill coma recovery time during infection may be contributed by insufficient energy stores.

Decline in negative geotactic ability, was also measured by how high a fly climbs in 10s post startle, in control and infected flies. In cold hardening lifespan and immunity is mainly affected [54] whereas an advantage of negative geotaxis assay is that the same set of flies were possibly used throughout the course of an infection. Hereby, both chill coma recovery and negative geotaxis could be useful in understanding the time course and degree to which different infections alter health in different genetic backgrounds of D. melanogaster. In addition to provide new measures for assessing health, these assays also suggest different pathological consequences of and metabolic up and downs that may occur over the course of an infection. Overall in brief, we can say that measures of health using these non-lethal behavioral assays may provide further insight into the mechanisms by which this model organism is able to maintain good health in facing of infections.

V. Conclusion

We demonstrated that how both chill coma recovery and negative geotaxis could be used in deep understanding the time duration and different infections duration degree that results in structural changes of health in different genetic backgrounds of D. melanogaster. Health measurements using these pleiotropic assays may further provides different measures that how an organism behaves and shows its leverage after fungal infection.

References


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