

## Study on Types, Total and Differential Haemocytes counts of Usherhopper, *Poekilocerus bufonius* Klug

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**ABSTRACT.** This study presented the identification of types, the total and differential haemocytes counts of usherhopper, *Poekilocerus bufonius*. Different histochemical tests were used to distinguish between various haemocyte types. Five types were morphologically characterized; namely, prohaemocytes (PRs), plasmotocytes (PLs), granulocytes (GRs), spherulocytes (SPs) and adipohemocytes (Ads). Total haemocytes counts in nymphal stages and both adult's male and female were 1300/mm<sup>3</sup>, 892/mm<sup>3</sup> and 838/mm<sup>3</sup>, respectively. The relative percentage of types and differential haemocytes counts of nymph, male and female were calculated. The results were discussed on the light of the existing references with emphases on the possible effect of the food consumed by usherhopper.

### Introduction

Several histochemical studies have been carried out on the identification and classification of insect haemocytes using light microscope<sup>[1-7]</sup>. There are many discrepancies in the categories of haemocyte types ranging from four to seven<sup>[6]</sup> and between three and nine<sup>[1,3,4,7-9]</sup>. In addition, there is confusion between various haemocyte types such as prohemocytes and plamatocytes as well as granulocytes and adipohemocytes<sup>[2,5]</sup>. However, many types of haemocytes were reported in different insect species; namely, prohaemocytes (PRs), plamatocytes (PLs), granulocytes (GRs), spherulocytes (SPs), oenocytoids (Oes), coagulocytes (Cos), adipohemocytes (Ads), reticular and other cells.

Total haemocyte counts (THC) and differential haemocyte counts (DHC) were determined in the haemolymph of different stages of various insects species<sup>[10-13]</sup>.

Usherhopper, *Poekilocerus bufonius* Klug (pyrgomorphidae), a black and yellow spotted locust feed on the milkweed (usher), *Calotropis procera* (Ait.), which is a perennial shrubby plant with broad, evergreen leaves and grows wildly in most arid region of Saudi Arabia. Usher is toxic to most vertebrates and invertebrates<sup>[14]</sup>. The toxicity is due to the presence of a high content of cardiac glycosides (SGs) in the latex<sup>[14, 15]</sup>. However, usherhopper ingest, sequesters and concentrates CGs in a bilobed poison gland<sup>[15, 16]</sup> and ejects them when disturbed as a means of defense against predators<sup>[16, 17]</sup>. The haemolymph CGs content of both male and female are almost similar to that of poison gland all year round<sup>[15]</sup>. There has been no study on the types, total haemocytes counts (THC) and differential haemocyte counts (DHC) of *P. bufonius*. So, the present study was undertaken.

### Materials and Methods

Various stages of male and female usherhopper, *Poekilocerus bufonius*, were collected from different locations around Jeddah. They were maintained under laboratory conditions (20-25°C) and were fed on branches of fresh usher.

To prepare blood smears, one leg of an insect (male, female and nymph of both sexes) were amputated and drops were smeared on microslides. The smears were fixed in methanol, washed in distilled water and stained with Giemsa and histochemical stains such as Periodic Acid Schiff's reagent for carbohydrates, Acid hematein for Phospholipid and Alcian blue for mucopolysaccharides.

#### *Total haemocyte counts (THC) and differential haemocyte counts (DHC)*

Total and differential haemocyte counts were carried as described by Harzarika and Gupta<sup>[18]</sup> and Chiang, *et al.*<sup>[19]</sup>. Helige true count haemocytometer was used to dilute haemolymph, and to avoid coagulation. Three drops of ethylenediaminetetraacetic acid (EDTA) buffer solution (0.1M glucose, 0.10mM EDTA, 0.45M sodium chloride, 30mM sodium citrate and 26mM citric acid) was added to the collected samples. The appropriate volume was well shaken and semiaired on Neubauer chamber slide. Degenerated and mitotic cells were included in the THC<sup>[20]</sup>.

### Results

The observation obtained using light microscope, in the present study, showed five types of haemocytes in the haemolymph of all stages and both sexes of *P. bufonius*; namely, prohaemocyte, plasmatocyte, granulocyte, spherulocyte and adipohemocyte. However, there were polymorphs of all haemocyte

types. Prohaemocytes were characterized by their small size, spherical shape and central large round nucleus, occupied most of cell cytoplasm, which form a very thin layer surround the nucleus (Fig. 1,2,3 and 4). Histochemical study (Table 1) indicated that PRs react moderately to PAS stain (Fig. 2), very strong to alcian blue (Fig. 3) and negatively to hematein test (Fig. 4). Plasmatocytes were amongst the most polymorphic and prominent types (Fig. 5). Their shapes ranged from spindle, with very pointed end, to oval and have a large centrally placed nucleus. Plasmatocytes were negatively stained with acid hematein (Fig. 6), however very strongly stained with alcian blue (Fig. 7) and PAS (Fig. 8). Granulocytes varied in shape and size, and were characterized by the presence of a small nucleus and large amount of different size granules (Fig. 9). Very few GRs have small cytoplasmic processes. Histochemical test showed that GRs were stained moderately with the three types of stains used in this study (Fig. 10 A, B, C, D, E). Spherulocytes (SPs) appeared in different shapes (polymorphic) with a small eccentric nucleus, spherules and membrane – bounded vacuoles in the cytoplasm (Fig. 11). These reacted positively with the histochemical stains (Fig. 12 A, B, C) used in this study. Adipohemocytes were very few when compared with other types of haemocytes and ranged in shape from circular to irregular (Fig. 13A, B, C and D). They contained a very large amount of lipid droplets, which occupied most of the cytoplasm. Treatment of adipohemocytes with acid hematein showed very strong reaction (Fig. 13B, C and D). All ADs from adults demonstrated negative reactions to staining with PAS (Fig. 14) and alcian blue (Fig. 15). However, ADs of nymph showed a positive reaction (Fig. 16).

The THC of hemocytes of different sex (nymph and adult) of usherhopper varied considerably (Fig.17A, B and 18A and B). The numbers of cells per  $\text{mm}^3$  were more or less similar of both sexes, male (892 cells) and female (838 cells). The nymphs of both sexes contained more THC than the adults (1300 cells/ $\text{mm}^3$ ). However, the percentage of each cell types varied (Fig. 19A, B and 20A and B). It can be seen that apart from adult males (27%), the GRs accounted for 46%, 62% and 57% of the haemocytes of female, female nymph and male nymph, respectively. In addition, the PRs form between 5-8 % of nymph THC of both sexes, and 15-24% of adult male and female. However, the PLs and GRs are the most prominent blood cell types of all stages (Fig. 17 and 18), and account for around 70% of the THC.

## Discussion

The results of the present study, using light microscope and histochemical tests, demonstrated that the haemolymph of the various stages and sexes of usherhopper contained five types of haemocytes; being prohaemocyte (PRs),

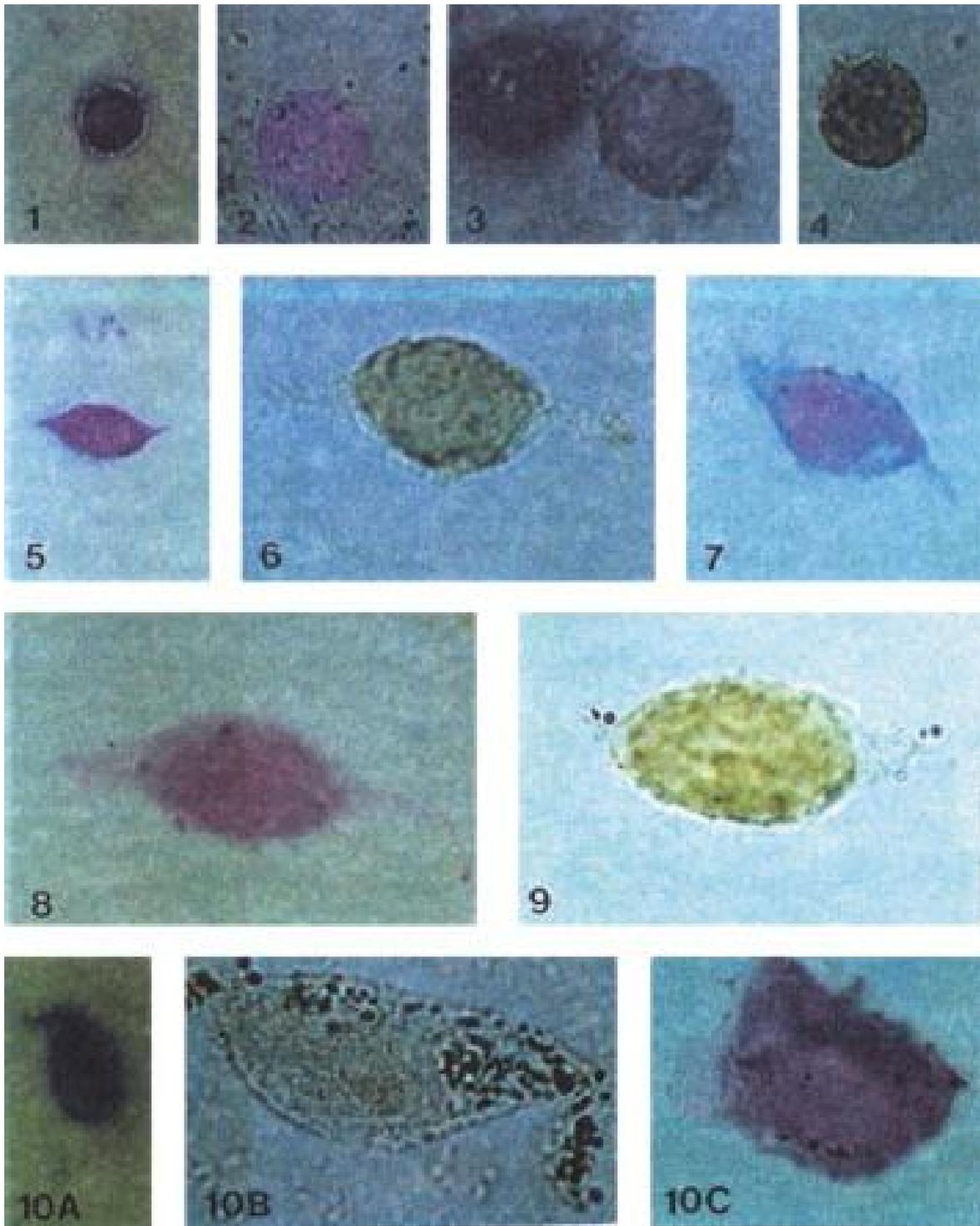


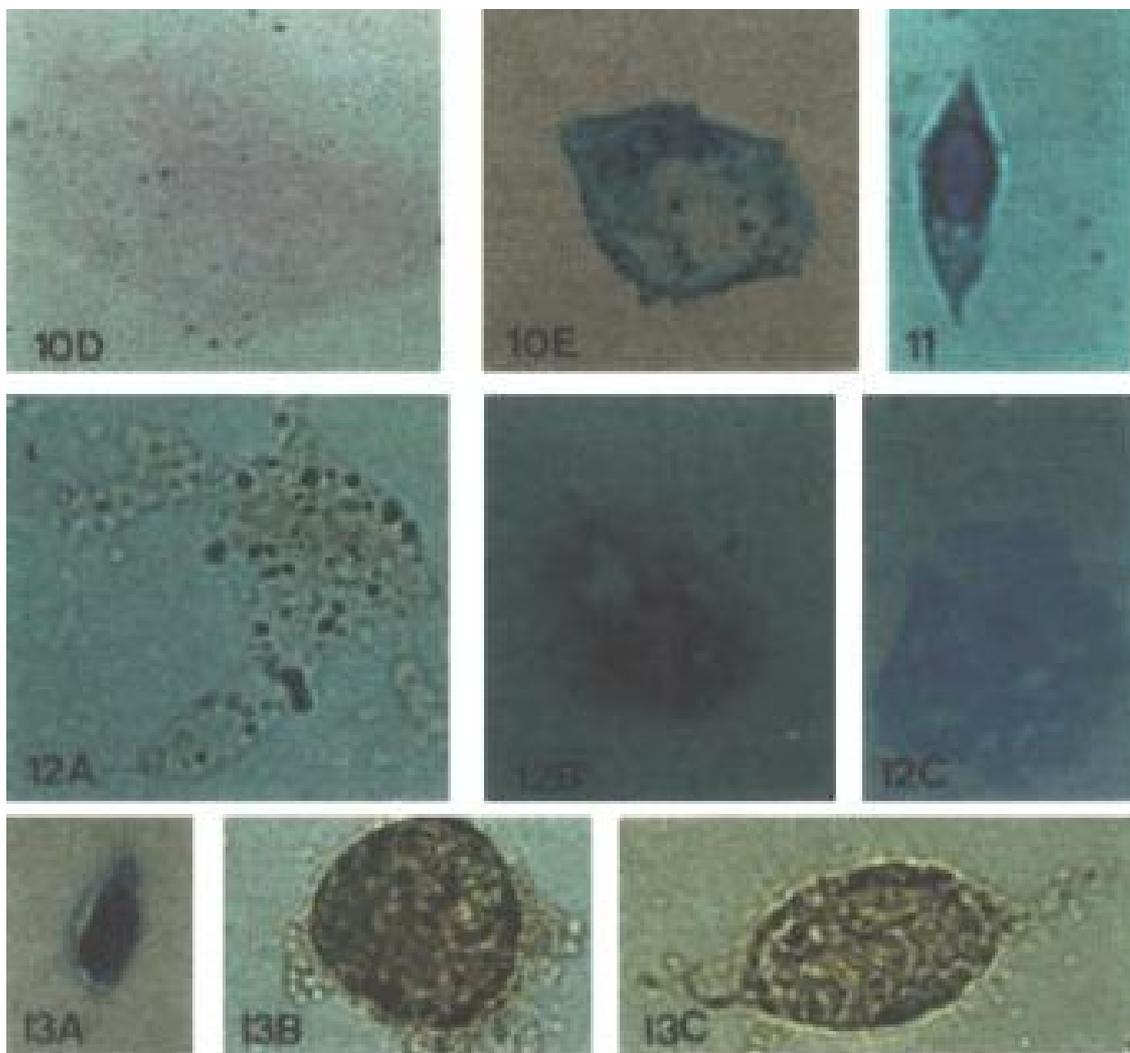
FIG. 1. Prohemocyte stained with Giemsa showing large nucleus occupying most of the cell cytoplasm ( $\times 400$ ).

FIG. 2. Prohemocyte stained with alcian blue indicating the presence of few amounts of mucoproteins ( $\times 1000$ ).

FIG. 3. Prohemocyte showing moderate reaction with PAS ( $\times 1000$ ).

FIG. 4. Prohemocyte showing negative stained with acid hematein ( $\times 1000$ ).

- FIG. 5. Plasmatocyte stained with Giemsa and characterized by spindle shape and pointed end ( $\times 400$ ).
- FIG. 6. Plasmatocyte showing negative reaction to acid hematein ( $\times 1000$ ).
- FIG. 7. Plasmatocyte stained positively with alcain blue ( $\times 1000$ ).
- FIG. 8. Plasmatocyte showing positive reaction to PAS ( $\times 1000$ ).
- FIG. 9. Plasmatocyte demonstrate trace of positive to acid hematein ( $\times 1000$ ).
- FIG. 10. Granulocyte stained with Giemsa (A) ( $\times 400$ ), react positively with acid hematein (B) ( $\times 1000$ ) and PAS from male (C) ( $\times 1000$ ) and female (D) ( $\times 1000$ ). Note the amount of granules in the cytoplasm (Fig. 10B and E) ( $\times 1000$ ).



- FIG. 11. Spherulocyte stained with Giemsa ( $\times 400$ ).
- FIG. 12. Showing various spherulocytes reacts positively with acid hematein (A) ( $\times 1000$ ) PAS (B) ( $\times 1000$ ) and alcain blue (C) ( $\times 1000$ ).
- FIG. 13. Demonstrate adipohemocytes stained with Giemsa (A) ( $\times 400$ ) and reacts positively to acid hematein whether it from nymph (B) ( $\times 1000$ ), adult male (C) ( $\times 1000$ ) and adult female (D) ( $\times 1000$ ).

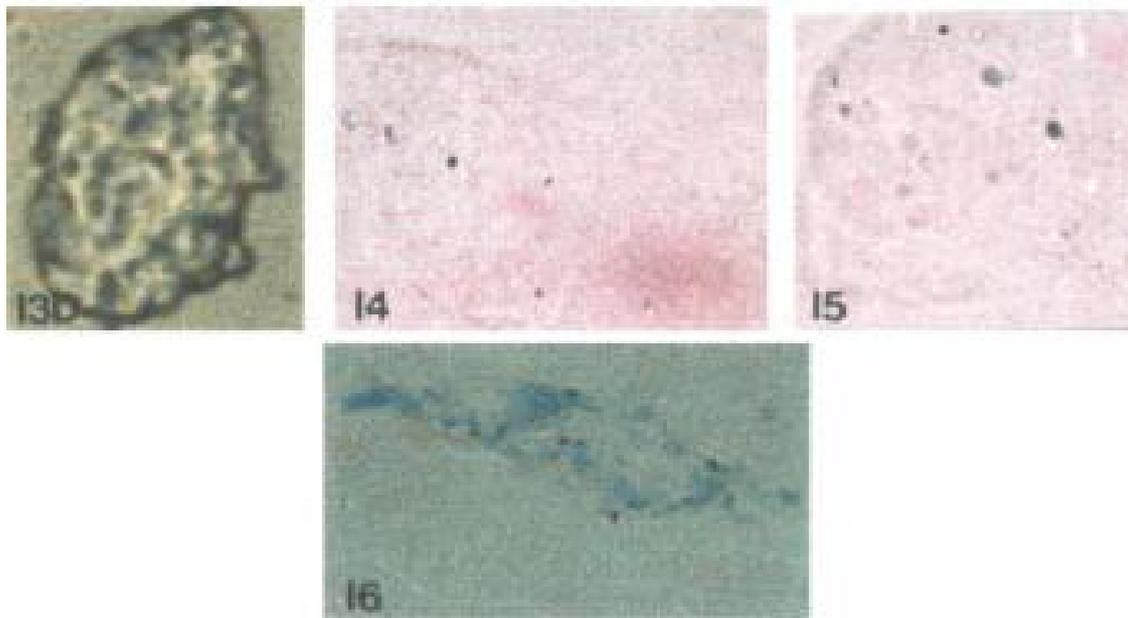


FIG. 14. Adipohemocyte react negatively to PAS ( $\times 1000$ ).

FIG. 15. Adipohemocyte react negatively to alcian blue ( $\times 1000$ ).

FIG. 16. Adipohemocyte of nymph shows positive reaction to alcian blue ( $\times 1000$ ).

plasmatocyte (PLs), granulocyte (GRs), spherulocyte (SPs) and adipohemocyte (Ads). Similar results have been reported for various insect<sup>[6]</sup>. The conflict in terminology, used to describe the same type of haemocytes<sup>[2,5,21]</sup>, between different reports may be due to the various characteristics used for classification of haemocytes. These features included cell shapes<sup>[4]</sup>, morphology, ultrastructure<sup>[22-26]</sup> and function<sup>[6,21,27-29]</sup>.

Histochemical study (Alcian blue, Acid hematein and PAS) showed that *P. bufonius* contained five types of haemocytes. However, there are different degrees of response to these stains. Similar results have been reported in haemocytes of *Garausius morosus*<sup>[6]</sup> in which three types of GRs were found. It is possible to suggest that the differences in shape and responses to various stains was due to differences in growth turnover between haemocytes, such as PRs to PLs and others. It has been suggested that the PRs are the stem cells for the other types of haemocytes<sup>[29]</sup>. This might have occurred, at least, in embryonic stages and not in post embryonic stages<sup>[6, 7]</sup>, which indicated that PRs are the origin of PLs. Some authors used its presence in all stages of insect life as strong evidence to support their role in producing other haemocytes<sup>[29, 30]</sup>. It has been proposed that many, if not all, cell types are merely stages in the development of a single type and the others are arbitrary selections from a continuous range of variation<sup>[31]</sup>. The present study indicated that some haemocytes fulfill the function of food storage (GRs, SPs and Ads) since they

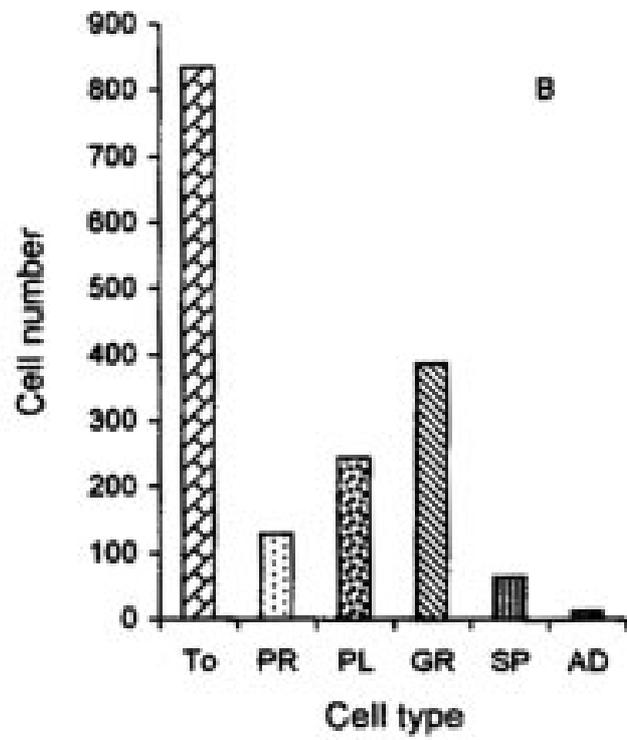
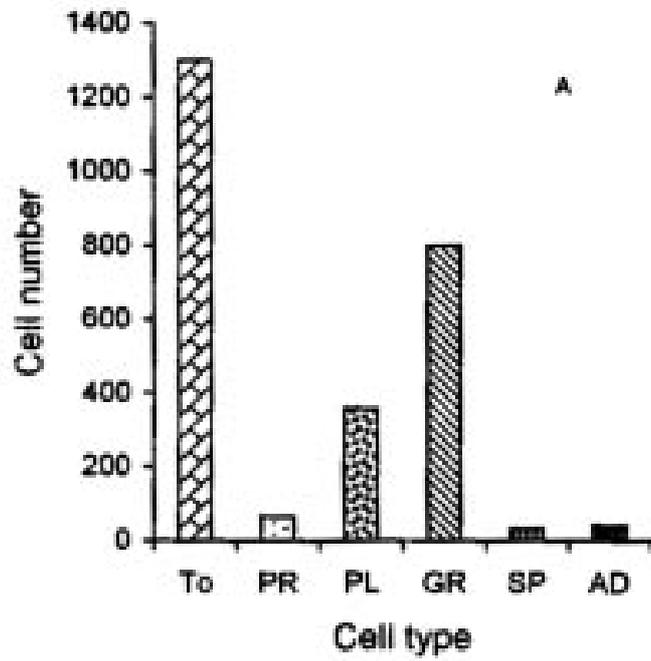


FIG. 17. Total and differential haemocytes counts of nymph (A) and adult (B) female *Poekilocerus bufonius*.

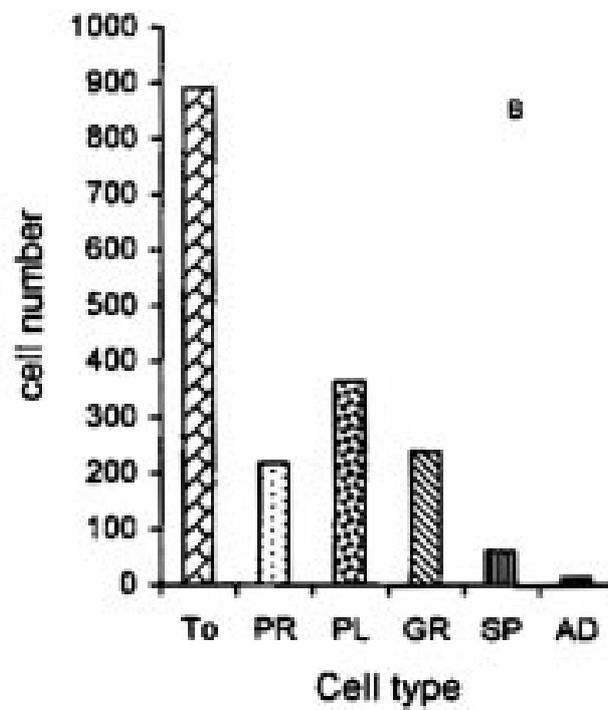
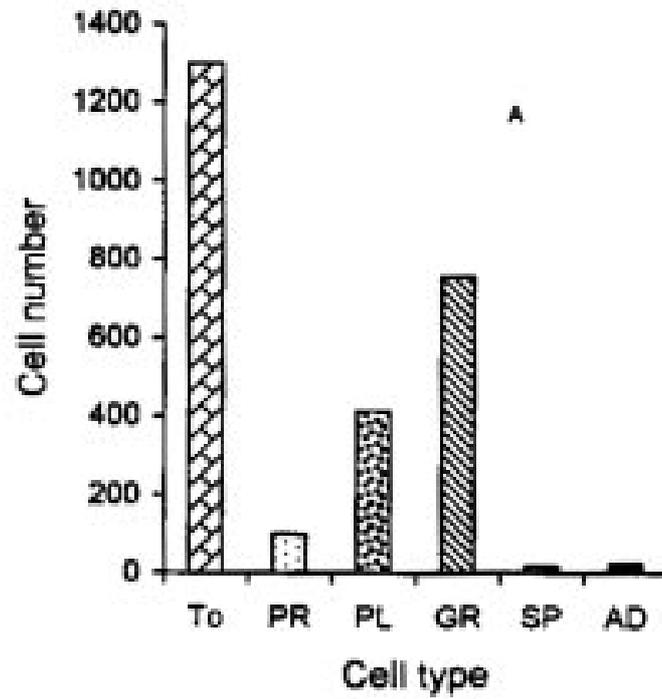


FIG. 18. Total and differential haemocytes counts of nymph (A) and adult (B) male *Poekilocerus bufonius*.

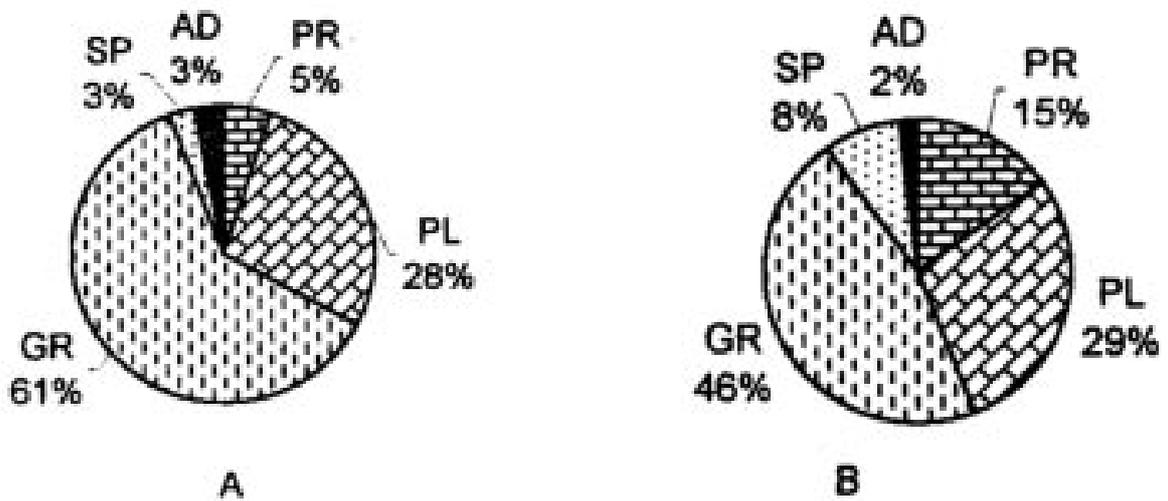


FIG. 19. Percentage of haemocytes of nymph (A) adult female (B) *Poekilocerus bufonius*.

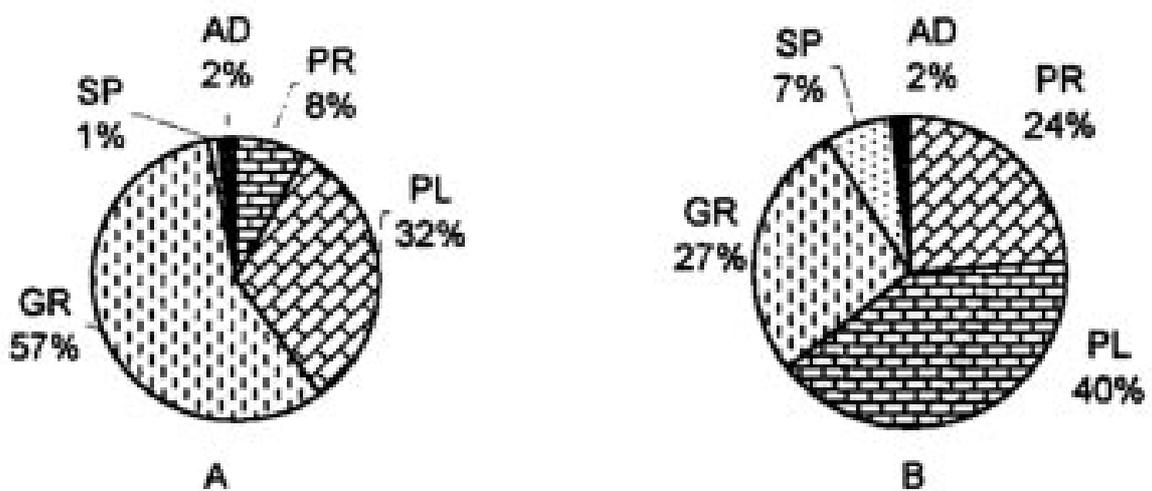


FIG. 20. Percentage of haemocytes of nymph (A) and adult male (B) *Poekilocerus bufonius*.

contained inclusions of carbohydrate and lipid, which decreased on starvation of the insect<sup>[32, 33]</sup>. The very low THC of usherhopper haemolymph, when compared with that of locusts might have been due to the type of food-consumed<sup>[15]</sup> and losing the ability to fly<sup>[13, 34]</sup>. As pointed out in the introduction usherhopper consumed toxic plants, such *Calotropis procera*, which contained cardiac glycosides. The later would have reduced the need for high number of haemocytes such that in locusts<sup>[6]</sup>.

### References

- [1] **Jones, J.C.**, Current concept concerning insect haemocytes, *Am. Zoo.*, **2**: 209-246 (1962).
- [2] **Jones, J.C.**, Normal differential counts of haemocytes in relation to ecdysis and feeding in *Rhodnius*. *J. Insect Physiol.*, **13**: 1133-1141 (1967).
- [3] **Arnold, J.W.**, A comparative study of the haemocytes (blood cells) of cockroaches (Insecta, Dictyoptera, Blattari), with a view of their significance in taxonomy, *Can. Ent.*, **104**: 309-348 (1972).
- [4] **Arnold, J.W.**, The haemocytes of insect, In: **Rockstein, M.** (eds.), *The Physiology of Insecta*, **5**: 202-254 (1974).
- [5] **Nruwirth, M.**, The structure of the haemocytes of *Galleria mellonella* (Lepidoptera), *J. Morph.*, **139**: 105-124 (1973).
- [6] **Gupta, A.P.**, *Insect haemocytes*, Cambridge University Press, Cambridge. p. 614 (1979).
- [7] **Al-Khalifa, M.** and **Siddiqui, M.**, A comparative study of haemocytes in some coleopterous species, *J. Call. Sci., King Saud Univ.*, **16**: 199-134 (1985).
- [8] **Yeager, J.F.**, The blood picture of the southern army worm (*Prodenia eridania*), *J. Agric. Res.*, **71**: 1-40 (1945).
- [9] **Wigglesworth, V.B.**, Insect blood cells, *A. Rev. Ent.*, **4**: 1-16 (1959).
- [10] **Nittono, Y.**, Study on the blood cells in the silkworm, *Bombyx mori* L., *Bull. Seric. Exp. Stn., Tokyo*, **16**: 171-266 (1960).
- [11] **Lee, R.M.**, The variation of blood volume with age in the desert locusts (*Schistocerca gregaria*), *J. Insect Physiol.*, **6**: 36-51 (1961).
- [12] **Bardoloi, S.** and **Hazarika, L.K.**, Seasonal variation of body weight, lipid reserve, blood volumes and haemocytes population of *Anthreaea assama* (Lepidoptera: Saturniidae). *J. Environ.*, **6**: 1-6 (1992).
- [13] **Chiange, S.A.**, **Gupta, A.P.** and **Han, S.S.**, Arthropod immune system: I. Comparative light and electron microscopic accounts of immunocytes and other haemocytes of *Blattella germanica* (Dictyoptera: Blattellidae), *J. Morph.*, **198**: 257-267 (1988).
- [14] **Al-Robai, A.A.**, Toxicological studies on the latex of Usher plant, *Calotropis procera* (Ait). in Saudi Arabia. IV – Effect of partly purified Usher latex and the poison gland secretion of the Ushershopper, *Poekilocerus bufonius* Klug on desert locust, *Schistocerca gregaria* Forskal (Orthoptera: Acrididae), *Arab Gulf J. Sci. Res.*, **15**: 709-716 (1997).
- [15] **Al-Robai, A.A.**, **Abo-Khatwa, A.N.** and **Jamal, Z.A.**, Toxicological studies on the latex of Usher plant, *Calotropis procera* (Ait) in Saudi Arabia. V – Seasonal variation of total cardiac glycosides in the Usher plant latex and in various tissues of the Ushershopper, *Poekilocerus bufonius* Klug. *Arab Gulf J. Sci. Res.*, **16**: 129-144 (1998).
- [16] **Euw, J.V.**, **Fishelson, L.**, **Parsons, J.A.**, **Reichstein, T.** and **Rothschils, M.**, Cardenolides (Heart poisons) in a grasshopper feeding on milkweeds, *Nature*, **214**: 35-39 (1967).

- [17] **Brewer, L.P. and Glazier, S.C.**, Localization of heart poisons in the monarch butterfly. *Science*, **188**: 19-25 (1975).
- [18] **Hazarika, L.K. and Gupta, A.P.**, Variations in haemocyte populations during various developmental stages of *Blattella germanica* L. (Dictyoptera: Blattellidae), *Zool. Sci.*, **4**: 307-313 (1987).
- [19] **Gupta, A.P.**, Studies of the blood of Meloidae (Coleoptera). 1. The haemocytes of *Epicauta cinerea* (Forster) and a synonymy of haemocyte terminologies, *Cytologia*, **34**: 300-344 (1969).
- [20] **Gupta, A.P.**, Cellular elements in the haemolymph: In **G.A. Kerkut and L.I. Gilbert** (ed.) *Comparative Insect Physiology, Biochemistry and Pharmacology*", Vol. **13**. Pergamon Press, p. 451 (1985).
- [21] **Akai, H. and Sato, S.**, Ultrastructure of the larval haemocytes of the silkworm, *Bombyx mori* L. (Lepidoptera: Bombycidae), *Int. J. Insect Morph. Embryol.*, **2**: 207-231 (1973).
- [22] **Akai, H. and Sato, S.**, An ultrastructure study of haemopoietic organs of the silkworm, *Bombyx mori* L., *J. Insect Physiol.*, **17**: 1665-1676 (1976a).
- [23] **Akai, H. and Sato, S.**, Surface ultrastructure of the larval haemocytes of the silkworm, *Bombyx mori* L. (Lepidoptera: Bombycidae), *Int. Insect Morphol. Embryol.*, **5**: 17-21 (1976b).
- [24] **Akai, H.**, Surface and internal ultrastructure of haemocytes of some insects, In **Gupta, A.P.** (ed.), *Insect Haemocytes*, Cambridge University Press, Cambridge, pp. 129-154 (1979).
- [25] **Essawy, M.A., Maleville, A. and Brehelin, M.**, The haemocytes of *Heliothis armigera*, Ultrastructure, function and evolution in the course of larval development, *J. Morphol.*, **186**: 225-264 (1985).
- [26] **Kim, C.S., Yoon, I.B. and Kim, W.K.**, Ultrastructure of the haemocytes in *Lymantria dispar* L. *Korean J. Entomol.*, **20**: 223-230 (1990).
- [27] **Gupta, A.P.**, Arthropod immunocytes: Their identification, structure, function and functional analogies with those of vertebrate B- and T-lymphocytes. In **Gupta, A.P.** (ed.), *Haemocytic and Humoral Immunity in Arthropods*, John Wiley and Sons, New York, p. 59 (1986).
- [28] **Amirante, G.A.**, Production of heteroagglutinins in haemocytes of *Lucophaea maderae* L., *Experientia*, **32**: 526-528 (1976).
- [29] **Kaya, G.P. and Ratcliffe, N.A.**, Comparative study of haemocytes and associated cells of some medically important Dipterean, *J. Morph.*, **173**: 531-365 (1982).
- [30] **Fenoligo, C. and Gervaso, M.V.**, Cytochemical characterization of the haemocytes of *Leucophaea maderae* (Dictyoptera: Balberoidae), *J. Morph.*, **218**: 115-126 (1993).
- [31] **Richards, O.W. and Davies, R.G.**, *Imam's Outline of Entomology*, (6th ed.), Chapman and Hall, London, p. 254 (1977).
- [32] **Yeager, J.F. and Munson, S.C.**, Histochemical detection of glycogen in blood cells of the south worm (*Prodenia eridania*) and in other tissues, especially midgut epithelium, *J. Agric. Res.*, **64**: 257-294 (1941).
- [33] **Munson, S.C. and Yeager, J.F.**, Fat inclusions in blood cell of the southern armyworm, *Prodenia eridania* (Cram.), *Ann. Entomol. Soc. Am.*, **37**: 396-400 (1944).
- [34] **Al-Robai, A.A. and Al-Ghamdi, H.S.**, Histological and ultrastructural study on the atrophied flight muscle of the female Usherhopper, *Poekilocerus bufonis* (Klug), *J.K.A.U. Sci.*, **6**: 49-74 (1993).

## الأنواع والعدد الكلي وأعداد الأنواع المختلفة لخلايا الدم في

### نطاط العشر *Poecilocerus bufonius* Klug

علي بن أحمد الرباعي و أحمد إبراهيم السقاف و ندى عثمان إدريس

قسم علوم الأحياء ، كلية العلوم ، جامعة الملك عبد العزيز

جدة - المملكة العربية السعودية

المستخلص . تهتم هذه الدراسة بالتعرف على أنواع وأعداد خلايا الدم في نطاط العشر *Poecilocerus bufonius* . استخدمت الصبغات الهستوكيميائية للتمييز المظهري بين الأنواع المختلفة . تم التعرف على خمسة أنواع من خلايا الدم تشمل : الأولية Prohaemocytes ، البلازمية Plasmacyte ، الحبيبية Granulocytes ، الكروية Spherulocytes والدهنية Adipohemocytes . قدر العدد الكلي في الأطوار الحورية والذكور والإناث البالغة بـ ١٣٠٠ / مم<sup>٣</sup> ، ٨٩٢ / مم<sup>٣</sup> و ٨٣٨ / مم<sup>٣</sup> ، على التوالي . تم حساب النسبة المئوية لكل نوع في الحوريات والحشرات البالغة ورسمت بيانياً . نوقشت النتائج على ضوء الدراسات السابقة مع التأكيد على تأثير نوع الغذاء الذي يستهلكه نطاط العشر . قلة عدد خلايا الدم في نطاط العشر استبدلت (تم تعويضه) بقدرتها على استهلاك النباتات السامة المحتوية على الجليكوسيدات القلبية المستخدمة كوسيلة دفاع .