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## Evaluation of residual and fumigant effects of Deltamethrin WG250 on different surfaces against *Anopheles stephensi* for malaria control

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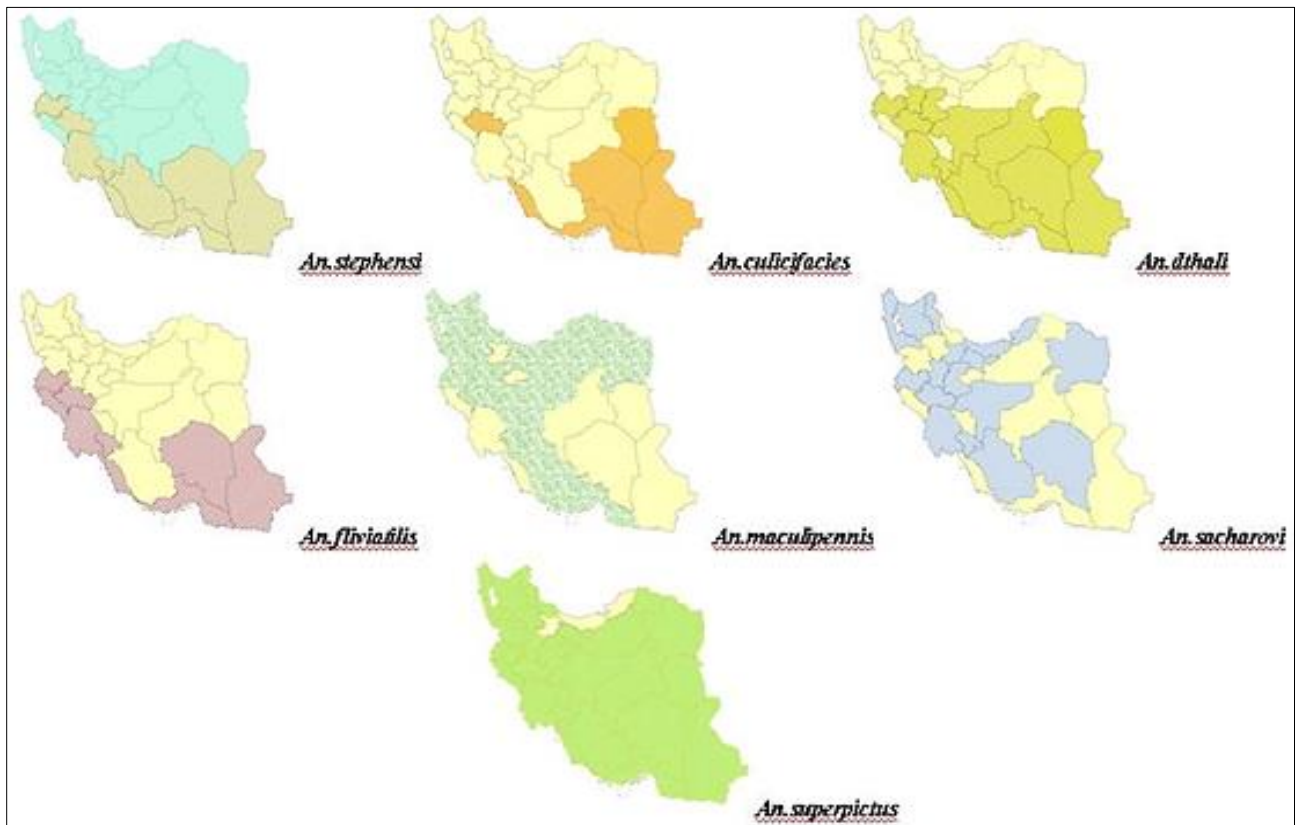
**Abstract**

Malaria is one of the public health problem in southern Iran. The main activity of vector control is indoor residual spraying using pyrethroids. The aims of study was to evaluate the biological assays of deltamethrin WG250 at different surfaces of wall. The persistency of deltamethrin WG250 at 25 mg/m<sup>2</sup> was studied on different local surfaces of rooms such as plaster and mud surfaces (sorberent) as well as wooden and thatch roofs (as non-sorberent). Contact bioassays were carried out using WHO standard cones and lab-bred sugar-fed, 48-72 h old females of *Anopheles stephensi* (Iranshahr strain). In contact bioassays carried out on sprayed surfaces for 150 days. Deltamethrin WG250 (25 mg/m<sup>2</sup>) caused 43.43-100.0% on sorberent surfaces and 61.11-100.0% on non-sorberent surfaces and persist about 3.5 months in the environmental condition of Saravan district. The fumigant bioassay of deltamethrin WG250 revealed 50-100% mortalities after one month.

**Keywords:** Deltamethrin, residual effects, fumigant efficacy, anopheles stephensi, malaria, Iran

**Introduction**

Malaria and other mosquito-borne disease are the major problems worldwide. Malaria presents a major health problem globally. It is estimated that globally 243 million cases of malaria led to 863,000 deaths in 2008. In most countries where malaria is endemic, the disease disproportionately affects poor and disadvantaged people, who have limited access to health facilities and can barely afford the recommended treatment. (WHO, 2020) [1]. WHO recommended insecticides for indoor residual spraying against malaria vectors are: DDT, Malathion, Fenitrothion, Pirimiphos-methyl, Bendiocarb, Propoxur, Alpha-cypermethrin, Bifenthrin, Cyfluthrin, Deltamethrin, Etofenprox, Lambda-cyhalothrin, Clothianidin [2]. Similarly main malaria cases in the country has been reported from southern and southeastern areas. The most routes of malaria cases are immigration from neighboring countries to southern and southeastern areas of the country (WHO, 2017) [3]. The main important vector control in the country are using insecticides as indoor residual spraying, impregnate bednets and larviciding by applying *Bacillus thuringiensis* in the breeding places. Currently there are proven and effective tools to fight against malaria including vector control measures [4]. A total of 228 million cases of malaria occurred worldwide in 2018. Most malaria cases (93%) were in African Region. *Plasmodium falciparum* is the most prevalent malaria parasite [5]. Iran is one of the malaria-endemic countries in the world, especially in southern provinces. The total number recorded cases have dropped to less than 89 locally-transmitted cases in 2017. Iran started a malaria elimination programme with a goal to achieve this target by 2025. There has been excellent progress since, but the continued risk of importation of malaria cases from Pakistan and Afghanistan. Main malaria vectors are *Anopheles stephensi*, *An. culicifacies*, *An. dthali*, *An. fluviatilis*, *An. superpictus* *An. maculipennis* and *An. sacharovi* [6]. Distribution of malaria vectors is shown in Fig.1.



**Fig 1:** Spatial distribution of malaria vectors in Iran

Campaign against malaria vectors was started from 1952 by DDT spraying and then replaced by dieldrin, Malathion, propoxur, lambdacy halothrin and deltamethrin, respectively. The chemical control of vectors now is restricted to endemic malarious areas of south-eastern part of the country with Deltamethrin and residual spraying and long lasting permethrin impregnated nets (Olyset) for personal protection, while biological control is conducting by *Bacillus thuringiensis* as larvicide. There are different aspects of malaria including insecticide resistance monitoring [7, 18], novel methods for vector control [19, 31], using bednets and

long lasting impregnated nets [32-38], vector control [39], and repellent evaluation [40]. The aim of current study was to evaluate the residual and fumigant effects of deltamethrin wg250 on different surfaces against *Anophels stephensi* for malaria control

**Materials and methods**

**Study area**

The experiments was carried out in a malarious areas in Saravan, Sistan and Baluchistan province, southern Iran (Fig.1)



Insecticide sued: deltamethrin wg250 were used (Fig.2)

**Fig 2:** Map of study area



Fig 3: Deltamethrin wg250 were used for evaluation

**Composition of mosquitoes in the region:** Fig. 3 shows the composition of Anopheles mosquito in the study area:

**Strain used for bioassay test:** the adult females of *Anophels stephensi* (Iranshahr strains were used for bioassay tests)

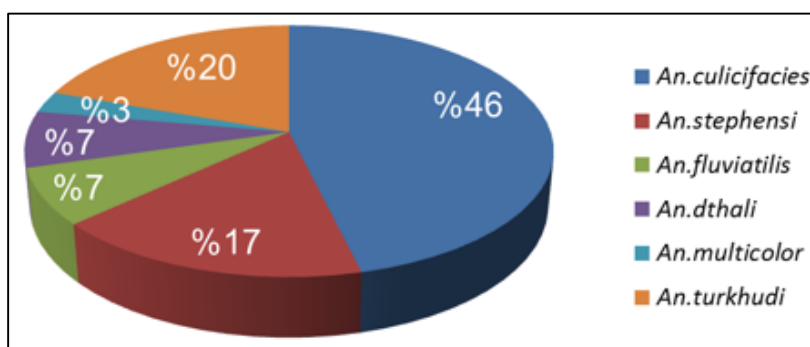


Fig 4: Composition of Anopheles mosquito in the study area

**Method of bioassay tests**

Bioassay aspirator tube especially narrow with glass arm were used. The conical exposure chamber made of transparent polished plastic and then adhesive sponge plastic for lining rim of exposure chamber.

**Method of fumigant effects**

3 cylindrical cages (14 x 20 cm) were employed. The number of mosquitoes per cage was 30-35, female, unfed, 2-3 days old. They were exposed 60 min in conical chamber and then the mortality was calculated after 24-hours.

**Results**

**Result of contact bioassay on Deltamethrin WG250 (25 mg/m<sup>2</sup>).** The results of bioassay test on different surfaces are illustrated in Figures 4-7. Residual effect at different surfaces is about four months. Fumigant effect is about 5-30 days.

**Residual effect of deltamethrin in plaster surfaces:** In plaster surface the standard mortality which is 80% could be obtained 105 days after application. After 90 days 100% mortality were observed.

**Residual effect of deltamethrin in Mud surfaces:** In mud surface a total of 643 mosquitoes were used. The standard mortality which is 80% could be obtained 135 days after application. After 90 days of application 100% mortality were observed

**Residual effect of deltamethrin in thatch surfaces**  
A total of 603 mosquitoes were used. 100% mortality was observed after 120 days. The residual effect in this surface is 4.5 months.

**Residual effect of Deltamethrin in Wood surfaces:** A total of 488 mosquitoes were used. 100% mortality was observed after 60 days. The residual effect in this surface is 5 months.

**Fumigant effect deltamethrin**  
In fumigant test a total of 1012 mosquitoes were used and 50% mortality were obtained after 30 days.

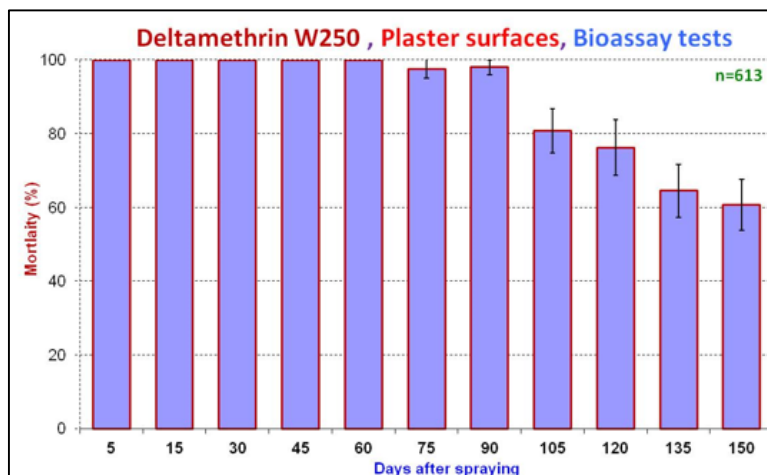


Fig 5: Results of contact bioassay test of deltamethrin on plaster surface

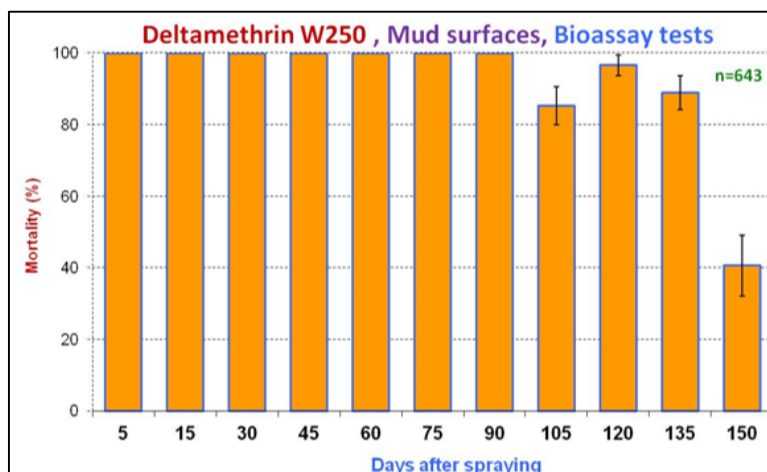


Fig 6: Results of contact bioassay test of deltamethrin on mud surface

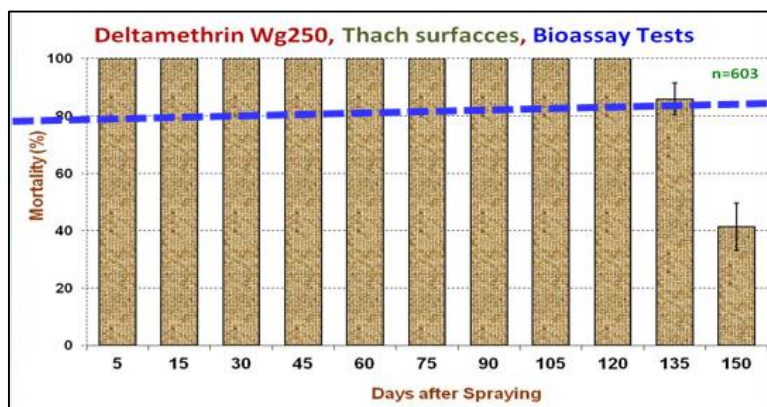
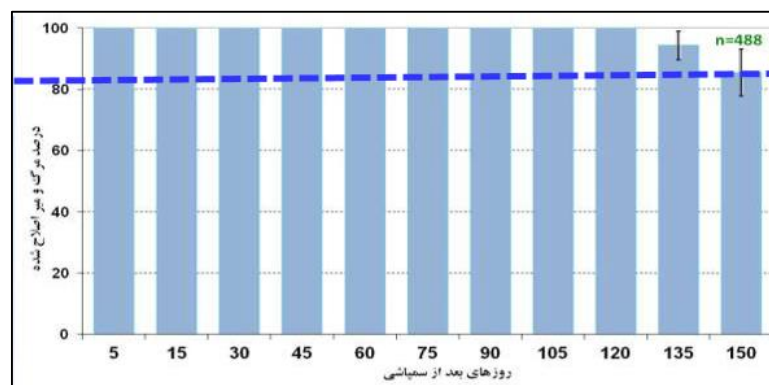
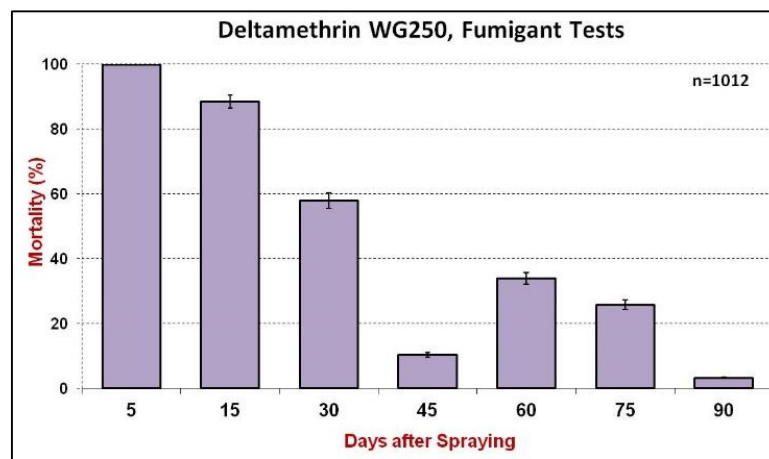


Fig 7: Results of contact bioassay test of deltamethrin on wood surface



**Fig 8:** Results of contact bioassay test of deltamethrin on Thatch surface

The fumigant effect of deltamethrin is shown in Fig 8.



**Fig 9:** The fumigant effect of deltamethrin

## Discussion

In the present study the residual effect of deltamethrin WG250 according the WHO method at dosage of 25 mg/m<sup>2</sup> using local species of *An. stephensi* at different surfaces of plaster, mud, thatch, wood is 4 months and fumigant effect= 5-30 days (MR=50-100%). There are several reports on bioefficacy of deltamethrin against different malaria vectors worldwide. The residual effects of deltamethrin WG 25% on different surfaces was assessed in southern part of Iran using *An. stephensi*. based on results, from 100% to about 70%. At 25, 40 and 50 mg a.i./m<sup>2</sup> the WG formulation of deltamethrin had a bioefficacy for about 2, 3 and 4 months respectively [41]. Deltamethrin at dosage of 50 mg/m<sup>2</sup> resulted 100% mortality against *An. gambiae* for 5.5 months [42]. The bioefficacy of indoor residual sprayed deltamethrin wettable granule (WG) formulation for the control of malaria was compared with the current dose of deltamethrin wettable powder (WP) in malaria endemic areas in Balai Ringin, Sarawak. Doses of 20 mg/m<sup>2</sup> WP (control), 20 mg/m<sup>2</sup> WG, 30 mg/m<sup>2</sup> WG and 40 mg/m<sup>2</sup> WG were sprayed separately on different surfaces namely, wooden, rough-bamboo, smooth-bamboo and brick surfaces. Residual activity of WP and WG formulations was tested against lab-bred *An. maculatus* using WHO standard procedure. Deltamethrin at 30 mg/m<sup>2</sup> WG exhibited the highest sustainable level of effectiveness against *An. maculatus*. Mortality was between 95%-100% up to week 60 post -spraying when sprayed on smooth-bamboo surface [43].

## Conclusion

The results of study revealed that deltamethrin as

recommended dosage as residual spraying could control the malaria vector during the transmission season. Monitoring and mapping of insecticide resistance to WHO recommended adulticide is recommended periodically.

**Conflict of interest (mandatory for all article types, also for Editorials, Commentaries and Letters):** The authors declare that there is no conflict of Interest.

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**Ethical approval (including committee and record number):** Ethical committee of Tehran University of Medical Sciences approved the research under code number 23550.

**Informed consent:** All the authors agree for submission of paper.

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