Distribution Behaviour of Dimethoate in Tea Leaf

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ABSTRACT

A study was undertaken to assess the distribution behaviour of Dimethoate in tea leaf. Tea bushes were subjected to Dimethoate spray at recommended dose and double the recommended doses. The extraction of pesticide was done using chloroform and the analysis was done using a Hewlett-Packard 5890 series II gas chromatograph with Nitrogen Phosphorus Detector (NPD). The penetration behaviour of Dimethoate was studied in dry and wet seasons. Variations in penetration were observed in dry and wet seasons which was attributed to climatic factors like temperature, humidity, rainfall, sunlight and physicochemical properties of the residue like water solubility, partition coefficient and formulation type. Residues observed in the cell wall and tissues confirm its good penetrating ability inspite of its hydrophilic nature. Higher penetration in wet season as indicated can be attributed to the route through the stomatal pores.

Keywords: Tea, Cuticle, Dimethoate, Dissipation

1. Introduction

Tea, Camellia sinensis (L) O. Kurtze, is the most important plantation crop in India occupying 420,000 ha of land. All over the world it is a popular invigorating and refreshing drink having excellent medicinal properties. Considering that an estimated amount of 18 - 20 billion cups are consumed daily in the world its economic and social interest is clear [1]. Like any other crop tea plantations are also subjected to ravages of insects, mites, plant pathogens, nematodes etc. The perennial nature of the crop and the more equitable weather pattern prevailing in tea areas are favorable for insect pests resulting in 250-500 million kg of annual loss of crop. In terms of monetary loss it could be approximately 500 million-1 billion US $ [2]. The insects target almost all plant parts such as roots, stem, leaves and buds. Thus every part of the tea plant is a potential target for a wide spectrum of pests and disease causing organisms.

Tea is an unusual crop where leaves are sprayed directly with pesticides, harvested and processed even without washing. The shoots of the tea plant are thin and tender (233 - 291 μm) with cuticle thickness 4 - 10 μm and surface area per unit weight of leaf (tender leaf 41.7 cm²/gm and mature leaf 15.9 cm²/gm respectively). Tea leaf, with thin cuticle, relatively larger surface area and short interval between pesticide application and harvest, is expected to be permeable to various groups of chemicals. Pesticide thus could have negative impact both on ecology and quality of tea when applied [3]. Consequently tea represents a significant potential of human exposure to pesticide residues by virtue of high application of pesticides to tea crop coupled with the average intake of 6 gms of dried (made) tea per day per individual [4,5]. Some basic studies on plant cuticles as barriers against the diffusion of chemicals have been reported [6,7]. Pesticide penetration into the leaf surface plays important practical implications as it allows the residues to stick/persist thus maintaining its efficacy even if it rains after the treatment [8]. Moreover once the pesticide has penetrated the epicuticular wax solar radiations acting on the pesticide molecule have to cross the cuticle thus reducing the photodegradative activity [9]. Furthermore, cuticular wax can hold on the pesticide residues resulting in low volatility from the leaf surface. The information available on the fate of pesticides on tea leaf and role leaf cuticle, plays in the dissipation under different environmental conditions is scanty in literature. Hence a study was initiated to understand the penetrating and dissipation behavior of dimethoate on tea leaf surface. Fate of dimethoate in tea and its brew have been reported but still its exact behaviour on the tea leaf surface is not known thoroughly [10,11].
2. Experimental Details

2.1. Field Trials

Field trials were carried out in dry and wet seasons at Institute of Himalayan Bioresource Technology (IHBT) tea experimental farm at Banurri, Palampur (1300 msl, 32°6′20″N × 76°33′29″E), India. The maximum and minimum temperature during dry and wet seasons as recorded from the experimental farm was 32°C, 23°C and 30°C, 18°C respectively. Relative humidity was 64% and 82% respectively and the total rainfall was 97 mm during wet season where as dry season was without any rainfall. The commercial formulation of dimethoate was sprayed (spray volume 400 l/ha) on tea bushes considering each plot of 100 bushes per replicate at recommended dose (200 gm a.i./ha) and double of the recommended dose (400 gm a.i./ha). In control treatment (T0), water was sprayed. Treatments were carried out in dry and wet seasons. Spray was done with a calibrated Knapsack sprayer. The weather parameters were continuously recorded in the experimental period to monitor the effect of environmental conditions. Green tea leaves (two leaves and a bud) were plucked from each replicate of both the treatment and control plots and brought to the laboratory each time at 0 (immediately and 4 hours after spraying), 1, 3, 5, 7, 10, 14 and 21 days after the treatment. To see the effect of leaky cuticle damaged leaves were also collected from the same plots. Further to analyze the effect of cuticle on the photodegradation of Dimethoate a laboratory experiment was carried out in parallel by spraying dimethoate on glass plate (i.e. without leaf) and other with leaf under same environmental conditions to provide estimated behavioral information. The weather parameters during the experimental time are graphically represented in figure below (see Figure 1).

2.2. Analytical Standards and Working Solutions

An analytical grade of dimethoate was obtained from Dr.Ehrenstorfer Laboratories, Augsburg, Germany (reported purity > 98%). For the field studies, formulation grade of dimethoate (Rogor® 30 Emulcifiable Concentrate EC, Isagro Agrochemicals Pvt. Ltd., Mumbai) was procured from the local market. Standard solution (1000 mg/l) was prepared in acetone and the spiking solution (50 mg/l) was diluted from the stock solution and the solutions required for preparing a standard curve (0.2, 0.5, 1.0, 1.5, 5.0 and 10.0 mg/l) were prepared from the stock solution by serial dilutions. All chemicals used for extraction and analysis of the residues and activated carbon were products of Merck India Limited, Mumbai India. Anhydrous sodium sulphate AR (Analytical Reagent) used was supplied by S.d. fine-chemicals, Mumbai.

Figure 1. Weather parameters.

2.3. Apparatus

Gas Chromatograph

A Hewlett-Packard 5890 series II gas chromatograph (Avondale, PA-USA) supported by a nitrogen phosphorus detector (NPD), a HP-7673 autosampler and integrator (Hewlett-packard) using split-splitless injector, connected to HP 3365 Chemstation system software (Hewlett-packard) was used. The detection was done using Nitrogen Phosphorus Detector (NPD). Peak resolution was done on a HP-17 medium polar capillary column (25 m length × 0.2 mm id) containing 50% phenyl and 50% methyl polysiloxane coated fused silica (0.25 µm film thickness) (Hewlett-Packard, Co., Wilmington, DE). Detailed analytical conditions were as follows. The injection was made using a split mode (50:1), injector temperature held at 280°C. The temperature of the NPD was held at 260°C. The carrier gas was nitrogen (purity 99.99%) at column flow rate of 1 ml/min. The samples were filtered through millipore
membrane teflon filters (0.45 µm particle size) before injection into the chromatographic column.

3. Standard Calibration Curve

Standard curve was prepared by diluting the stock solution to five different concentrations in acetone. The column was conditioned by repeated injections (3 times) of each matrix (green leaves, dried leaves and soil). These samples were prepared by adding known amount of standard in matrix before extraction. The extraction was carried out as described below in section. The duplicate injections of each extract were made in Gas Chromatograph (GC-NPD).

4. Extraction of Pesticide from Leaf Surface

Extraction of dimethoate was done with Chloroform (recovery > 90%) The extract was agitated mechanically with acetonitrile for 3 hours on a horizontal shaker. The mixture was filtered through Whatman no.1 filter paper and the cake was washed twice with 20 ml solvent each time. The combined water extract was partitioned with 150ml of acetonitrile twice in a 500 ml separating funnel. Discarding the aqueous layer, the organic layer was concentrated to near dryness on a water bath and reconstituted with 1 ml of acetone for final analysis. The effect of washing was thus confirmed by laboratory washing and in case of samples collected from the treated fields in dry and wet seasons, the residue was extracted in dichloromethane. The final eluate was evaporated to dryness and reconstituted with 1 ml of acetone for quantification.

4.1. Extraction of Pesticide from Epicuticular Wax

Epicuticular wax extraction was done using the method described by Mc. Donald et al. [12]. After thorough washing to remove the surface pesticides, the tea leaves were soaked in 100 ml of chloroform and shaken on an automatic horizontal shaker for 1 minute. Extract was filtered through Whatman No.1 filter paper and concentrated to 5 ml followed by passing through an adsorbent column containing florisil topped with 1 cm of anhydrous sodium sulphate prewashed with chloroform. The extract was eluted with 200 ml of chloroform, concentrated on a vacuum rotary evaporator using a water bath at 35°C - 40°C. The extract was finally eluted with dichloromethane (200 ml) from the florisil column. The eluate was evaporated to dryness and reconstituted to 1 ml in acetone and 2 ml of it was analysed by GC (NPD).

4.2. Extraction of Pesticide from Cell Wall

Extraction of dimethoate was done by soaking the tea leaves in 100 ml of chloroform and shaken on an automatic horizontal shaker for 1 minute. Extract was filtered through Whatman No.1 filter paper and concentrated to 5 ml followed by passing through an adsorbent column containing florisil topped with 1 cm of anhydrous sodium sulphate prewashed with chloroform. The extract was eluted with 200 ml of chloroform, concentrated on a vacuum rotary evaporator using a water bath at 35°C - 40°C. The extract was finally eluted with dichloromethane (200 ml) from the florisil column. The eluate was evaporated to dryness and reconstituted to 1 ml in acetone and 2 ml of it was analysed by GC (NPD).

5. Detection and Quantification

Detection limit test

To determine the limit of detection made tea samples were spiked with different concentration levels of dimethoate standard and analysed by GC (NPD). The detection limit was evaluated by the peak signal/noise (S/N) ratio. An S/N ratio greater than 3 was considered as a detectable peak.

6. Results and Discussion

6.1. Quantification

The GC analytical conditions were optimized in terms of temperature program that allowed an improvement of the time and the chromatographic run resolution. Moreover, the adopted oven programming allowed a good chromatographic separation of dimethoate. The total run time was 14 minutes and the retention time of the dimethoate in the given chromatographic conditions was 8.68 minutes and constant for each series of samples. Chromatographic separation by HP-17 column provided good results for the quantification of the samples.

6.2. Linearity

The calibration curve of the analysed dimethoate gave a good regression line (R² = 0.8827) in the range of explored concentrations, 0.1 - 10.0 mg/l. The detection limit of dimethoate was taken to be 0.01 mg/kg, which were much lower than the maximum residue limits fixed by European Commision for dimethoate in tea (0.2 mg/kg). Residues below 0.01 mg/kg were detected but not quantified. This low detection limit was achieved...
because of the efficient cleanup step that allowed the elimination of all the possible interfering peaks, giving a low noise value.

6.3. Method Validation

The recovery of the fortified (50 mg/kg) samples of dimethoate in made tea (in triplicate) ranged from 92.9% - 94.6%. The results showed good recovery and reproducibility. The compounds of interest were well resolved from other co-extractives. These results indicated that the method used in this study provided a good cleanup.

6.4. Field Studies

The shoots of the tea plant are thin and tender (233 - 291 \( \mu m \)) with cuticle thickness 4 - 10 \( \mu m \) and relatively larger surface area per unit weight of leaf (tender leaf 41.7 and mature leaf 15.9 cm\(^2\)/gm respectively). It is an unusual crop where leaves are sprayed directly with pesticides, harvested and processed even without washing. In the preliminary experiments with tea leaf cuticle penetration of dimethoate was measured immediately after application as evaporation of water also affected the rate of penetration [13].

As indicated by data presented in Tables 2-5, the residues of dimethoate on leaf surface at 0 day (immediately after spray) was found to be 9.07 ± 0.14 mg/kg and 18.31 ± 0.34 mg/kg whereas its concentration was 9.01 ± 0.10 mg/kg and 18.19 ± 0.34 mg/kg respectively at two different treatments when the samples were collected 4 hours after spray. No penetration effect observed in the epicuticular wax and cell wall when the leaves were collected immediately after spray. This indicated that during the evaporation of water the penetration rate was negligible. In dry season the residue concentration in the epicuticular wax was 0.14 ± 0.05 and 0.11 ± 0.10 mg/kg and in the cell wall was 2.97 ± 0.02 and 4.41 ± 0.08 mg/kg in normal leaf when the samples were collected 4 hours after spray at 200 and 400 gm a.i/ha respectively. While the concentration in injured leaf was 0.17 ± 0.02 and 0.68 ± 0.06 mg/kg in epicuticular wax and 3.04 ± 0.06 and 6.12 ± 0.12 mg/kg in cell wall.

Further, the concentration in leaf surface was found to be 0.02 ± 0.00 mg/kg and 0.04 ± 0.00 mg/kg at two different treatments on 7th day after spray. No residues were detected on leaf surface after 10th day of treatment.

In wet season, the residue concentration in the cell wall was 3.24 ± 0.06 mg/kg and 6.25 ± 0.19 mg/kg respectively at two different treatments in normal leaf and 3.32 ± 0.11 mg/kg and 7.20 ± 0.17 mg/kg in injured leaves whereas in wax the residue concentration was 0.03 ± 0.00 mg/kg, 0.08 ± 0.01 mg/kg and 0.08 ± 0.00 mg/kg, 0.08 ± 0.00 mg/kg in normal and injured leaves respectively. Further, this concentration in wax was negligible in normal leaf and injured leaf on 3rd day after spray, which declined to no detectable limit in 7th day after spray. As observed from the data, the residues found on the epicuticular wax and cell wall was comparatively more in the damaged cuticle.

Residues observed in the cell wall and tissues confirm its good penetrating ability inspite of its hydrophilic nature. Higher penetration in wet season as observed from the tables indicated the route could be through the stomatal pores also. Little variation in the results as observed from Tables 2-5 for normal and damaged cuticle confirmed that cuticle was also acting as a barrier during penetration of dimethoate residue. The tea leaf having thin cuticle is expected to be permeable to various groups of pesticides and dimethoate residue in the wax and cell wall confirmed it but cuticle thickness alone may not be responsible for the dimethoate penetration in the leaf. Data in Table 1 proved that physical properties of the

<table>
<thead>
<tr>
<th>COMMON NAME</th>
<th>DIMETHOATE</th>
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</thead>
<tbody>
<tr>
<td>CHEMICAL NAME</td>
<td>O,O-DIMETHYLS-METHYL-CARBAMOYL-METHYL PHOSPHORODITHIOATE</td>
</tr>
<tr>
<td>EMPIRICAL FORMULA</td>
<td>C(<em>{5})H(</em>{12})NO(<em>{3})PS(</em>{2})</td>
</tr>
<tr>
<td>MOLECULAR WEIGHT</td>
<td>229.3</td>
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<tr>
<td>VAPOUR PRESSURE</td>
<td>2.5 × 10(^{-4}) Pa at 25°C</td>
</tr>
<tr>
<td>PHYSICAL STATE</td>
<td>COLOURLESS CRYSTALLINE SOLID.</td>
</tr>
<tr>
<td>MELTING POINT</td>
<td>45 - 52.5°C</td>
</tr>
<tr>
<td>Boiling point</td>
<td>107°C at 0.05 mmHg</td>
</tr>
<tr>
<td>SOLUBILITY IN WATER</td>
<td>3g/l</td>
</tr>
<tr>
<td>SOLUBILITY IN ORGANIC SOLVENTS</td>
<td>HIGHLY SOLUBLE IN CHLOROFORM, METHYLENE CHLORIDE, BENZENE, TOULENE, ALCOHOLS, ESTERS AND KETONES.</td>
</tr>
<tr>
<td>SPECIFIC GRAVITY AT 25°C</td>
<td>1.281</td>
</tr>
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</table>

Figure 2. Standard curve.
### Table 2. Dimethoate distribution in tea leaf in dry season (200 gm a.i/ha).

<table>
<thead>
<tr>
<th>DAS</th>
<th>Leaf surface</th>
<th>Wax</th>
<th>Cell wall</th>
<th>Leaf surface</th>
<th>Wax</th>
<th>Cell wall</th>
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<td>0(1)</td>
<td>9.07 ± 0.14</td>
<td>ND</td>
<td>ND</td>
<td>8.96 ± 0.17</td>
<td>0.02 ± 0.00</td>
<td>ND</td>
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<td>0(2)</td>
<td>9.01 ± 0.10</td>
<td>0.14 ± 0.05</td>
<td>2.97 ± 0.02</td>
<td>8.65 ± 0.13</td>
<td>0.17 ± 0.02</td>
<td>3.04 ± 0.06</td>
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<td>6.17 ± 0.15</td>
<td>0.11 ± 0.09</td>
<td>2.03 ± 0.00</td>
<td>5.92 ± 0.12</td>
<td>0.21 ± 0.02</td>
<td>2.79 ± 0.06</td>
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<tr>
<td>3</td>
<td>1.46 ± 0.07</td>
<td>0.02 ± 0.00</td>
<td>0.03 ± 0.00</td>
<td>1.30 ± 0.01</td>
<td>0.02 ± 0.00</td>
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</tr>
<tr>
<td>5</td>
<td>0.05 ± 0.00</td>
<td>ND</td>
<td>ND</td>
<td>0.04 ± 0.00</td>
<td>ND</td>
<td>ND</td>
</tr>
<tr>
<td>7</td>
<td>0.02 ± 0.00</td>
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<tr>
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</table>

### Table 3. Dimethoate distribution in tea leaf in dry season (400 gm a.i/ha).

<table>
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<th>Leaf surface</th>
<th>Wax</th>
<th>Cell wall</th>
<th>Leaf surface</th>
<th>Wax</th>
<th>Cell wall</th>
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<tr>
<td>0(1)</td>
<td>18.31 ± 0.34</td>
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<td>ND</td>
<td>17.60 ± 0.30</td>
<td>0.03 ± 0.00</td>
<td>0.02 ± 0.00</td>
</tr>
<tr>
<td>0(2)</td>
<td>18.19 ± 0.31</td>
<td>0.11 ± 0.10</td>
<td>4.41 ± 0.08</td>
<td>17.36 ± 0.25</td>
<td>0.68 ± 0.06</td>
<td>6.12 ± 0.12</td>
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<tr>
<td>1</td>
<td>11.39 ± 0.23</td>
<td>0.90 ± 0.07</td>
<td>4.10 ± 0.10</td>
<td>11.18 ± 0.16</td>
<td>0.33 ± 0.07</td>
<td>5.93 ± 0.06</td>
</tr>
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<td>3</td>
<td>3.13 ± 0.13</td>
<td>0.06 ± 0.00</td>
<td>0.04 ± 0.00</td>
<td>2.95 ± 0.07</td>
<td>0.03 ± 0.00</td>
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<td>0.10 ± 0.01</td>
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<td>0.06 ± 0.00</td>
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<td>0.04 ± 0.00</td>
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### Table 4. Dimethoate distribution in tea leaf in wet season (200 gm a.i/ha).

<table>
<thead>
<tr>
<th>DAS</th>
<th>Leaf surface</th>
<th>Wax</th>
<th>Cell wall</th>
<th>Leaf surface</th>
<th>Wax</th>
<th>Cell wall</th>
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<tr>
<td>0 (1)</td>
<td>4.15 ± 0.13</td>
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<td>ND</td>
<td>3.86 ± 0.10</td>
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<td>ND</td>
</tr>
<tr>
<td>0 (2)</td>
<td>4.00 ± 0.01</td>
<td>0.03 ± 0.00</td>
<td>3.24 ± 0.06</td>
<td>3.65 ± 0.08</td>
<td>0.02 ± 0.10</td>
<td>3.32 ± 0.11</td>
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<td>1</td>
<td>0.08 ± 0.00</td>
<td>0.05 ± 0.05</td>
<td>4.14 ± 0.09</td>
<td>0.06 ± 0.00</td>
<td>0.08 ± 0.13</td>
<td>4.25 ± 0.03</td>
</tr>
<tr>
<td>3</td>
<td>0.02 ± 0.00</td>
<td>0.02 ± 0.00</td>
<td>0.82 ± 0.03</td>
<td>0.03 ± 0.00</td>
<td>0.01 ± 0.00</td>
<td>0.94 ± 0.02</td>
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<td>5</td>
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residue and environmental conditions might be equally involved. Low partition coefficient $K_{ow}$ (0.7) and high water solubility of 39 g/l might have enabled it to penetrate inside the inner region of the cuticle. They are rapidly absorbed under high humid conditions, supporting the view that there was an aqueous route traversing the cuticle and maximum penetration might be due to stomatal pores. Thus the permeation of active ingredients was influenced by their solubility characteristics as evident by their partition coefficients. The $K_{ow}$ reflected the lipophilicity of the compound and was related to the degree and rate at which it would be absorbed by leaf [14]. Furthermore, penetration as observed in the leaf surface could be attributed to the emulsifiable concentrate formulation used as they allow better deposits and different adjuvants keep the deposit in binded form and minimize the early loss of pesticide by increasing the rate of penetration. The adhesion, retention and distribution of agrochemicals sprayed on plant surfaces also depends on target wettability [15]. However it could be envisaged that temperature and humidity could have roles to play.

8. Conclusions

Above studies confirmed that the dimethoate distribution on the tea leaf surface and decrease in surface residue is due to the collective behaviour of cuticle, its solubility, penetration, partition coefficient, vapour pressure and the formulation along with the environmental conditions at the time of experiments.

9. Acknowledgements

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