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Safety Assessment of Etofenprox, S-Methoprene, and Piperonyl Butoxide in Dogs Topically Exposed to Bio Spot Defense

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Abstract

Use of ectoparasiticides on pets seems inevitable since pets are commonly infested with fleas, ticks, and many other external parasites. Currently, pyrethroids are more commonly used than any other class of ectoparasiticides because they are considered relatively safe. Due to paucity of data, serious concerns have been raised not only about their safety in dogs but their transferable residues to owners, veterinarians, veterinary technologists, and dog handlers who come in contact with treated dogs on a daily basis. The present investigation was therefore undertaken to determine the toxicity and safety of “Bio Spot Defense” applied to six adult dogs, and to determine the residue of active ingredients (etofenprox, s-methoprene, and piperonyl butoxide) in dogs and cotton gloves worn for five minutes to pet dogs at 24, 48, 72 hours, and 1, 2, 3, 4, and 5 weeks post-application. At these time intervals, dogs were evaluated for physical examination. Residues of active ingredients were confirmed and quantitated using GC/MS. In the blood, etofenprox was detected as early as 48 hours (18.42 ± 5.05 µg/g) and the residue persisted until 1 week (0.80 ± 0.35 µg/g). S-methoprene and piperonyl butoxide were not detected in the blood. In the gloves, the highest concentrations of etofenprox, s-methoprene, and piperonyl butoxide were determined at 24 hours (9,552.00 ± 1551.83; 3,307.86 ± 456.70; and 1286.13 ± 0.49 µg/g, respectively). Residues of all three compounds were detected in appreciable concentrations in the gloves until 1 week (294.86 ± 27.22; 80.62 ± 10.06; and 40.49 ± 5.78 µg/g, correspondingly). Their residues persisted in insignificant amounts in gloves until 5 weeks. In conclusion, findings of this investigation suggest that Bio Spot Defense product appears to be safe for dogs and their owners, but the veterinary personnel can be exposed to significant levels of etofenprox, s-methoprene, and piperonyl butoxide following chronic exposure, if not properly protected.

Keywords: Etofenprox; S-methoprene; Piperonyl butoxide; Pyrethroids; Ectoparasiticide toxicity; Ectoparasiticide safety; Flea and tick killer; Bio spot defense

Introduction

Blood sucking ticks, fleas, mosquitoes, and other ectoparasites commonly infest dogs and pose a serious global health concern as they can transmit infectious diseases between humans and animals. The use of ectoparasiticides on pets therefore is not only imminent but seems inevitable. Currently, pyrethroids are more widely used than any other class of ectoparasiticides (organophosphates, carbamates, neonicotinoids, etc) because they are considered relatively safe due to their species selectivity [1,2]. Due to lack of safety data, serious concerns have been raised for their safe use in dogs and about transferable residues to the owners, veterinarians, veterinary technologists, and dog handlers who come in contact with these ectoparasiticides on a daily basis. Bio Spot Defense Flea and Tick Spot On’ (hereafter referred to as Bio Spot Defense) is a relatively new over the counter topical ectoparasiticide for dogs, which kills and prevents ticks, fleas, and mosquitoes. The product consists of three active ingredients: 1. Etofenprox (2-(4-ethoxyphenyl)-2-methylpropyl 3-phenoxybenzyl ether), 30.0%; 2. S-methoprene (2(E)-4E)-11-methoxy-3,7,11-trimethyl-2,4-dodecadienoic acid 1-methylthyl ester), 3.6%; and 3. Piperonyl butoxide (5-[2-(2-butoxyethylthio)ethoxyethyl]-6-propyl-1,3-benzodioxole), 5.0% [3]. Chemical structures of etofenprox, s-methoprene, and piperonyl butoxide are shown in Figure 1.

According to the label, the product is effective against ticks (Ixodes spp.), mosquitoes (Anopheles quadrimaculatus), fleas (Ctenocephalides spp.) eggs, flea larvae, and adult fleas for up to two and a half months, usually during the summer. Etofenprox is a rapidly acting pyrethroid insecticide, which can kill adult fleas and ticks on dogs as early as

Figure 1: Chemical structures of etofenprox, s-methoprene, and piperonylbutoxide.

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within 15 minutes of application [4]. Etofenprox is an insecticide with contact and systemic action, with applications in agriculture, public health and animal health. S-methoprene is an insect growth regulator due to its ovicidal and larvicidal actions; and piperonyl butoxide is a synergist, which enhances the persistence and effects of etofenprox and s-methoprene [5-7].

According to the Bio Spot Defense website, the possible side effects of this product include drowsiness, itching, redness or rashes, hair discoloration, and hair loss at the site of application [4]. There is a high confidence level about safety of Bio Spot Defense to the persons applying and to their pets due to a low acute mammalian toxicity of etofenprox, (s)-methoprene, and piperonyl butoxide; and minimal potential of residue transfer to humans once the product is dried, usually between 12-24 hours. Although the product label states that it is safe once dried, there have been concerns regarding transferable residue to humans coming into contact with Bio Spot Defense treated dogs on a daily basis. The Environmental Protection Agency has deemed both etofenprox and (s)-methoprene “safe”. According to section 408(b)(2)(A)(ii) of the Federal Food, Drug, and Cosmetic Act, “safe” is defined as “there is a reasonable certainty that no harm will result from aggregate exposure to the pesticide residue, including all anticipated dietary exposures and all other exposures for which there is reliable information.” This does not include occupational hazard [8]. Currently, there is no reported all other exposures for which there is reliable information. “This does not include occupational hazard [8]. Currently, there is no reported safety of etofenprox, (s)-methoprene and piperonyl butoxide that humans may be exposed to from coming in contact with dogs treated with Bio Spot Defense. In previous studies, we reported transferable residues of fipronil, imidacloprid and selamectin to humans from dogs treated topically with Frontline™, Advantage®, and dogs treated with Bio Spot Defense. In previous studies, we reported data regarding the levels of etofenprox, (s)-methoprene, and piperonyl butoxide that humans may be exposed to from coming in contact with dogs treated with Bio Spot Defense. In previous studies, we reported transferable residues of fipronil, imidacloprid and selamectin to humans from dogs treated topically with Frontline™, Advantage®, and Revolution™ respectively, using the glove sampling method for petting dogs and assessing transferable residue [9-12].

The acute oral LD₅₀ values for etofenprox, s-methoprene, and piperonyl butoxide in rats are reported to be 2880 mg/kg, 34500 mg/kg, and 7500 mg/kg respectively. In dogs, the acute oral LD₅₀ for racemic and s-methoprene is between 5000 to 10000 mg/kg. Dermal LD₅₀ values for etofenprox and piperonyl butoxide in rats are 2140 mg/kg and 7950 mg/kg respectively. The acute dermal LD₅₀ for both racemic and s-methoprene in rabbits is >2000 mg/kg. These toxicity data suggest that insecticides, such as etofenprox and s-methoprene, and a synergist piperonyl butoxide, are of low mammalian toxicity [4,5,13-16].

The present investigation was undertaken to assess the toxicity and safety of etofenprox, s-methoprene and piperonyl butoxide in dogs, and possible exposure to owners and veterinary personnel after topical application on dogs of Bio Spot Defense.

Materials and Methods

Animals

Six mixed breed adult dogs (medium length hair), weighing between 30-40 pounds, were used in this investigation. The dogs were not treated with any ectoparasiticide for at least two months prior to this study.

Chemicals

Bio Spot Defense Flea & Tick Spot On®, having etofenprox, s-methoprene, and piperonyl butoxide, was purchased from Farnam Pet Products (Phoenix, AZ). Technical grade Etofenprox (99.0%), s-methoprene (99.2%), and piperonyl butoxide (97.5%) were purchased from Chem Service (West Chester, PA). All other chemicals with highest purity were obtained from Fisher Scientific (Fair Lawn, NJ).

Experimental design

Ectoparasiticide application: Bio Spot Defense Flea & Tick Spot On® (Farnam Pet Products, Phoenix, AZ) in the amount of 4.44 mL (etofenprox, 30%; s-methoprene, 3.6%; and piperonyl butoxide, 5%) was topically applied on each dog. The product was applied directly to the skin on the back (at three evenly spaced spots) starting at the shoulder blades and ending at the base of the tail. The Smart Shield™ applicator was used to apply the product without humans coming in direct contact with Bio Spot Defense [3]. Following application of the product, the active ingredients redistribute into the skin, with a high concentration going into the sebaceous glands. The sebaceous glands release the drug continuously in the sebum along with other natural oils and waxes. This allows the drug to coat the hair shaft and skin surface providing long-lasting protection against fleas, ticks, and mosquitoes [4].

Physical Examination: Dogs topically treated with Bio Spot Defense were evaluated for physical parameters. At specified time intervals, dogs were examined for body weight, behavior, and skin reaction at the Bio Spot Defense application site.

Sample collection for pesticide residue analysis: Cotton gloves used for petting dogs were collected at 0, 24, 48, 72 hour, and 1, 2, 3, 4, and 5 week intervals; and blood samples were collected at 0, 24, 48, 72 hours, and 1 and 2 week intervals, for pesticides (etofenprox, s-methoprene, and piperonyl butoxide) residue analysis.

Glove sampling included the wipe sampling method, which consisted of petting the dog forward and back along its back and sides for five minutes with a 100% cotton glove. After sampling, the glove was immediately placed in a labeled 800 mL glass jar and kept at room temperature until analyzed (<72-96 hours). Blood samples were collected from the cephalic vein using a 6 mL syringe with a 22-gauge needle. At each time interval, approximately 5 mL blood was collected in an EDTA anticoagulant tube and kept in the refrigerator until analyzed.

Sample extraction: Dog blood and cotton glove samples were extracted in methylene chloride: petroleum ether (50:50, vol/vol). Extracts were passed through the sodium sulfate, evaporated to dryness overnight, and reconstituted in methylene chloride: petroleum ether just prior to GC/MS analysis.

GC/MS analysis: All three active ingredients present in Bio Spot Defense (etofenprox, s-methoprene, and piperonyl butoxide) were confirmed using an Agilent Gas Chromatograph (GC model 7990A)/Mass Spectrometer (MS model 5975C) coupled with a computer, and their concentrations were expressed in terms of µg/g. The evaporated extract was reconstituted in an appropriate volume of extraction solvents (methylene chloride: petroleum ether) and passed through a Sep-Pak® Cartridge (Waters Corp, Milford, MA). One µL of sample extract was injected into the GC. The column used was Ultra II Cross-linked with 5% phenyl methyl siloxane coating and the following dimensions (capillary 25 mm x0.52 µm), which was directly connected to the Mass Selective Detector via a heated transfer line. The carrier gas was ultrapure (99.9999%) helium at a flow of 1 mL/min, and the injector temperature was 200°C. The injector was operated in the splitless mode. A temperature program for the GC-oven was used comprising a starting temperature of 150°C for 1 min, and then increased to a final temperature of 300°C in 20°C/min increments. The final temperature was maintained for 5 min. The total duration of the chromatography for each injection was 14 min. The transfer line temperature was 280°C, the source temperature was 230°C. The instrument was operated in electron ionization mode, and the ion energy was 70eV. Peaks of s-methoprene, piperonyl butoxide, and etofenprox were eluted at 6.524
min, 7.967 min, and 10.105 min, respectively. Sensitivity of GC/MS for these compounds was in the range of ng, and limit of detection was in the range of low µg/g.

Results

A single application of Bio Spot Defense, having etofenprox, s-methoprene and piperonyl butoxide, caused no adverse effects in dogs at any time during the study period of five weeks. There was no significant change in body weight and no skin reaction at the site of insecticide application.

The present study simulated a model for ectoparasiticides’ exposure to veterinary personnel from handling canine patients treated with Bio Spot Defense. Residues of etofenprox, s-methoprene, and piperonyl butoxide transferred from the canine coat were measured on cotton gloves at various intervals using GC/MS. Total Ion Chromatogram revealed that the peaks of s-methoprene, piperonyl butoxide, and etofenprox eluted at 6.524, 7.967, and 10.105 min, respectively (Figure 2).

The concentrations of etofenprox, s-methoprene, and piperonyl butoxide present in the cotton gloves transferred from petting dogs at various time intervals are shown in Figures 3-5. The highest concentrations of all three pesticides were found at 24 hr post Bio Spot Defense application (9,552.00 ± 1551.83; 2,307.86 ± 456.70; and 1286.13 ± 0.49 µg/g, respectively). Significant concentrations of etofenprox, s-methoprene, and piperonyl butoxide were found in the gloves until one week (294.86 ± 27.22; 80.62 ± 10.06; and 40.49 ± 5.78 µg/g, correspondingly).

Figure 2: GC/MS-Total Ion Chromatogram (TIC) of s-methoprene (Mol. Wt., 310.47), piperonyl butoxide (Mol. Wt., 338.44), and etofenprox (Mol. Wt., 376.49). Peaks of s-methoprene, piperonyl butoxide, and etofenprox were eluted at a retention time of 6.524 min, 7.967 min, and 10.105 min, respectively. For detailed GC/MS conditions, see the text.

Figure 3: Concentrations of etofenprox (ppm) in gloves (Mean ± SEM; n=6).
Thereafter, with a steep decline trend, residues of these pesticides were present on the canine coat until five weeks post-application of Bio Spot Defense.

Dog blood analysis for pesticide residues revealed the presence of only etofenprox (Figure 6). Its highest concentration was detected after 48 hours (18.42 ± 5.05 µg/g). By 72 hours, etofenprox concentration reduced to 1.22 ± 0.42 µg/g, and after one week, the level was 0.80 ± 0.35 µg/g. Thereafter, etofenprox residue was undetectable in blood. At no time was residue of either s-methoprene or piperonyl butoxide detected in the dog blood.

Discussion

The present investigation was carried out in response to common queries of veterinarians about the safety of Bio Spot Defense to dogs and veterinary personnel. Although literature abounds showing the basic toxicity data of etofenprox, s-methoprene, and piperonyl butoxide (major ingredients in Bio Spot Defense) in rats and mice, no reports are there to support the safety of this product with regards to dogs, dog owners, or veterinary personnel. This study was therefore done in an attempt to simulate exposure of veterinary personnel to transferrable residue of etofenprox, s-methoprene, and piperonyl butoxide from the coat of dogs treated with Bio Spot Defense.

Based on types of symptoms and syndromes produced, pyrethroids are categorized into two types: (a) type I that lack cyano group termed as T (tremors), e.g., permethrin; and type II that contain cyano group termed as CS (choreoathetosis/salivation), e.g. deltamethrin [2,14,17-19]. Pyrethroids are neurotoxicants [20,21] and primarily affect the sodium channel of cells, but chloride and calcium channels are also...
Pyrethrins and pyrethroids slow the opening and closing of the sodium channels, resulting in excitation of the cells [25,26]. The duration of the sodium action potential is much longer for type II pyrethroids than for type I. Paresthesia results from the direct action of pyrethroids on sensory nerve endings, causing repetitive firing of these fibers. There is marked stereospecificity of the action of pyrethroids on the sodium channel, i.e. the cis isomers are usually more toxic than trans isomers [27,28]. It is noteworthy that unlike the other pyrethrins and pyrethroids, etofenprox is a non-ester pyrethroid, which does not contain a cyano group and centers of molecular asymmetry and therefore does not show stereoisomerism.

In dogs, toxic reactions to pyrethroids include itching and allergic reactions (paw flicking, ear twitching, and respiratory distress), and hypersalivation, that can eventually progress to diarrhea, in coordination, depression and muscle tremors. Clinical signs associated with pyrethroid exposure are usually resolved within 24 to 72 hours. Death occurs in dogs in rare cases [2]. The American Association of Poison Control Centers reported more than 20,000 pyrethroid exposures of canine patients every day. Therefore, it is important to assess the levels of transferrable residue of pyrethroids on dog’s coat to veterinary personnel. Currently, there are no published studies regarding the effects of these pesticides on human health from Bio Spot Defense use on dogs. In previous studies, we reported the transferrable residue of fipronil from Frontline™ [9], ingestion of products having pyrethrins and pyrethroids meant for dermal application.

There are some reports on pharmacokinetic and PBPK values of some pyrethroids [31-33], but very little is known about etofenprox. A wide species and age-dependent variation appears to exist in pharmacokinetics and susceptibility to pyrethroid toxicity [31,34-37]. The absorption of pyrethroids in mammals through the skin is very low and it is lesser in humans than in animals [37-39], but following oral administration it appears to be consistently 40-60% [14]. Hawkins et al. [40] reported that following oral administration of etofenprox (30 mg/kg or 180 mg/kg) in rats, its bioavailability was about 14-51%, and the residue was detected in blood (5-16 ppm), liver (0.34 ppm), fat (16.6 ppm), and brain (0.002-0.004 ppm). Phase I metabolites of etofenprox include desethyl-etofenprox, 4'-hydroxy-etofenprox, α-CO (2-(4-ethoxyphenyl)-2-methyl propyl-3-phenoxybenzoate), ethoxyphenyl-2-methyl propanol, 3-phenoxybenzoic acid, and 4'-OH-3-phenoxybenzoic acid; and phase II metabolites include glucuronide and sulfate conjugates of oxidation products [14,41,42]. Hawkins et al. [40] also indicated that etofenprox was eliminated 8-10.8% in the urine and 86.4-88% in the feces.

Peak blood/plasma and brain concentrations of pyrethroids tend to correlate with clinical signs of neurotoxicity after acute oral dosing of a pyrethroid [14,43,44]. Such signs usually dissipate within hours after a single gavage dosing, correlating with the reduction in blood/plasma pyrethroid concentration. In the present investigation, after 48 hours of Bio Spot Defense application, the concentration of etofenprox in the blood was 18.42 µg/g, which was 180 times less than that found on the coat (333.34 ± 43 µg/g). Lack of s-methoprene or piperonyl butoxide in the dog blood appears to be due to their inability to dermal penetration.

In the present study, residues of etofenprox, s-methoprene, and piperonyl butoxide on dog’s coat appeared to be within safe limits following dermal application of Bio Spot Defense. Currently, dogs and cats with pyrethrins and pyrethroids poisoning often seen in veterinary clinics and confirmed at the diagnostic centers might be due to oral death occurs in dogs in rare cases [2]. The American Association of Poison Control Centers reported more than 20,000 pyrethroid exposures of canine patients every day. Therefore, it is important to assess the levels of transferrable residue of pyrethroids on dog’s coat to veterinary personnel. Currently, there are no published studies regarding the effects of these pesticides on human health from Bio Spot Defense use on dogs. In previous studies, we reported the transferrable residue of fipronil from Frontline™ [9], ingestion of products having pyrethrins and pyrethroids meant for dermal application.

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Veterinarians and veterinary technicians handle a large number of canine patients every day. Therefore, it is important to assess the levels of transferrable residue of etofenprox, (s)-methoprene, and piperonyl butoxide from treated dogs to veterinary personnel. Currently, there are no published studies regarding the effects of these pesticides on human health from Bio Spot Defense use on dogs. In previous studies, we reported the transferrable residue of fipronil from Frontline™ [9],
imidacloprid from Advantage® [10], and selamectin from Revolution™ [11,12] treated dogs. Dogs treated with Advantage and Revolution also had in their blood significant residue of imidacloprid and selamectin, respectively, for several days post-treatment. The present study demonstrated that levels of etofenprox, s-methoprene, and piperonyl butoxide on the canine coat could be transferred to human skin surfaces directly or through glove contamination.

Conclusions

This investigation was undertaken to assess the safety and toxicity of Bio Spot Defense in dogs, and related health concerns to humans. Findings revealed the persistence of etofenprox, s-methoprene, and piperonyl butoxide residues on the canine's coat in significant amounts that can be transferred to humans through their contact with the dogs. The levels of these pesticides on the dogs' coat were highest at 24 hours post-application, which can be the time of greatest risk to human exposure. After 1 week, the concentrations rapidly declined, yet insignificant levels were present until week five. Etofenprox level in the dogs' blood was highest at 48 hours, and persisted until 1 week. At no time was s-methoprene or piperonyl butoxide detected in the dog blood. Bio Spot Defense was found to be safe for topical application, as none of the dogs showed any adverse effects, and no skin reactions occurred at the application site. Veterinarians, veterinary technologists, and dog handlers can be exposed to significant levels of ectoparasiticides following chronic exposure, being in close contact with treated dogs, if not properly protected.

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