Research Article

Structure-Activity Relationship (SAR) and *in vitro* Predictions of Mutagenic and Carcinogenic Activities of Ixodicidal Ethyl-Carbamates

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Ethyl-4-bromophenyl-carbamate (LQM 919) and Ethyl-4-chlorophenyl-carbamate (LQM 996) are compounds that inhibit egg-laying and hatching of tick larvae that are resistant to conventional ixodicides. The structure-activity relationship (SAR) to get the endpoint predictions of mutagenicity and carcinogenicity of the LQM 919 and LQM 996 was performed and the absence of mutagenicity was confirmed by Ames test. SAR analysis show no structural alerts indicating the ability of ethyl-carbamates to bind biomolecules or estrogen receptors. Endpoint of mutagenicity with and without metabolic activation showed that the ethyl-carbamates were negative (p < 0.05) for mutagenicity induction in strains TA97, TA98, TA102, TA1535, TA1537 and TA1538 of *Salmonella typhimurium*. Pre-incubation with different ethyl-carbamate concentrations did not increase the number of spontaneously reverting colonies; moreover, the compounds did not induce a concentration-dependent increase in the number of reverting colonies in any of the strains used. This confirmed the absence of mutagenic activity in this test system. Exogenous metabolic activation did not modify these observations; suggesting that no metabolites with mutagenic activity were present. The endpoint of carcinogenicity in rats were negative for LQM 919 (p < 0.05,) and LQM 996 (p < 0.001). The results of the present study strongly suggest that ethyl-carbamates do not represent a risk for cancer in mammals.

1. Introduction

Currently, regulation of chemical substances that are used daily as pesticides (i.e. ixodicides) is of great relevance globally. The European Chemicals Agency (ECHA), the Organization for Economic Cooperation and Development (OECD), and the Environmental Protection Agency (EPA), are the most important organizations responsible for setting the guidelines for regulation of chemical substances.

These organizations generally accept *in silico* methods based on computational modeling of chemical structure and biological activity of a compound to predict its toxicity without the use of biological tests. This allows reducing costs, the number of experimental animals, risks in handling hazardous chemicals, and the time used to perform *in vitro* or *in vivo* tests [1].

*Rhipicephalus microplus*, considered the most important parasitic tick on cattle in tropical and subtropical areas
worldwide, represents a great threat to the livestock industry
due to the diseases it transmits as a vector and the direct eco-
nomic losses it causes [2]. Excessive use of chemical sub-
stances for the control of ticks in cattle (ixodicides) has
resulted in development of tick strains that are resistant to
the majority of commercial ixodicides in several regions of
the world [3]. This has generated the need to develop new
molecules to which no tick strains are currently resistant.
We have shown that ethyl-4-bromophenyl-carbamate
(LQM 919) and ethyl-4-chlorophenyl-carbamate (LQM 996)
inhibit egg-laying and hatching of Rhipicephalus microplus
larvae and this inhibition was observed both in strains that
are resistant to other ixodicides and in those that are suscep-
tible to them [4, 5]. These carbamates alter the reproductive
organs, vitellogenesis, and viability of ovarian cells in R.
microplus and cause harm that is suggestive of apoptosis
[6]. These effects were found to be independent of acetylcho-
linesterase inhibition [7].

The toxicity of LQM 919 and LQM 996 must be evalu-
ated before these compounds are proposed for the safe con-
trol of ticks. Following the criteria of the OECD, these
carbamates are found to have low acute and subchronic tox-
icity in rats [8, 9]. Nevertheless, these compounds produce an
increase in micronucleated reticulocytes in the peripheral
blood of rats as well as alterations in the cell cycle of human
lymphocytes suggesting that these compounds cause DNA
damage [10]. Other carbamates such as vinyl-carbamate
and ethyl-carbamate (urethane) are carcinogenic and pro-
duce tumors in several organs of rodents [11, 12]. This sug-
gests a potential risk for the application of carbamates in
animals used for human food and to the persons who apply
these products. Therefore, this study aimed to evaluate the
mutagenic and carcinogenic potential of the ethyl-
carbamates LQM 919 and LQM 996 in silico, as well as to cor-
robate their mutagenic capacity using the Ames test.

2. Materials and Methods

2.1. Carbamates. The ethyl-carbamates used in this study
were designed, synthesized, and patented at the National
Autonomous University of Mexico using a benzimidazole
molecule as the structural base, as described by Angeles
et al. [13]. The chemical structures, nomenclature, molecular
weights, simplified molecular input line entry system
(SMILES), octanol/water partition coefficient (Kow) and
identification codes of ethyl-carbamates are shown in
Table 1.

2.2. In silico Studies. Structure-activity relationship (SAR) for
endpoint predictions of mutagenicity and carcinogenicity of
the ethyl-carbamates LQM 919 and LQM 996 was performed
by Read-across-analysis using the QSAR Toolbox software
according to workflow of the hazard assessment process:
Input; Profiling; Endpoint data collection; category defini-
tion; data gap filling, and finally, reporting [14]. The partition
coefficients (Log Kow) of each carbamate were used as
descriptors. A conformational study of LQM 919 and LQM
996 was conducted; the molecules were modeled using the
Toolbox modeler tool and were double checked by compar-
ing each SMILES code. Profiling was carried out using the
OECD HPV chemical categories, substance type, and US-
EPA New Chemical Categories profiling methods [15, 16].

The carbamate structures were optimized by PM3
(Molecular Orbital Model) and were categorized using the
DICE (Dynamic Integrated model of Climate and the Eco-
nomy) model to obtain the DICE coefficient [17, 18]. Molecu-
lar feature parameters (atom centered, fragments and cycles)
and atomic characteristics (count H attach, hybridization
and cyclic) were used to obtain a set of chemical analogues
with greater than 50% similarity to the carbamates LQM
919 and LQM 996. The endpoint predictions of mutagenicity
on strains TA97, TA98, TA102, TA1535, TA1537, and
TA1538 of Salmonella typhimurium with and without simu-
lation of S9 metabolic activation, and endpoint predictions
of carcinogenicity on rats by the oral route of ethyl-carbamates
were determined. The endpoint value predictions were based
on five neighbors’ values. Prediction confidence was estab-
lished and categorized as moderate (p < 0.05), strong (p <
0.01), or very strong (p < 0.001).

2.3. Ames’ in vitro Assay. S. typhimurium strains TA98 (his
D3052, rfa National Autonomous University of Mexico,
uvrB, bio-, pKM101[ApR]), TA100 (his G46, rfa, uvrB, bio-
, pKM101[ApR]), TA1535 (his G46, rfa, uvrB, bio-),
TA1537 (his C3076, rfa, uvrB, bio-), and TA102 (His G428,
rfra, bio-, pKM101[ApR], pAQ1[Tr]) were used; these were
maintained at the Mutagens and Carcinogens Laboratory of
the Biomedical Research Institute, National Autonomous
University of Mexico. Before the assays were carried out, sin-
gle colonies were isolated from each strain, and their geno-
typic characteristics were confirmed using the procedure
described by Maron and Ames [19]. The toxicity of varying
concentrations of the studied compounds was assessed by
microscopic observation of bacterial growth in the selected
dishes. Homogeneous distribution of bacteria was consid-
ered indicative of no toxicity, whereas loss of homogeneity and
the dose-dependent decrease in reverting colonies was consid-
ered indicative of toxicity.

Extract S9 from the livers of Sprague-Dawley male rats,
induced with phenobarbital and B-naphthoflavone, was pre-
pared and stored (-80°C) according to the modified method
described by Escobar-García et al. [20].

One colony from each strain grown in NB solid medium
was transferred to 10 mL of NB-Oxoid broth and kept under
constant stirring (150 rpm) for 12 h. Subsequently, 0.1 mL of
cultures from each strain (approximately \(10^8\) cells) was pre-
icubated for 30 min with a mixture of 0.5 mL of S9 mix or
0.5 mL of phosphate buffer at pH 7.4 and varying final con-
centrations of the studied carbamates (0, 0.1, 0.2, 0.4, 0.5
and 1 mg/mL) that had been previously diluted with
dimethyl sulfoxide (DMSO). The lowest concentration
(0.1 mg/mL) was selected considering the results of previous
assays that demonstrated no toxicity, and the concentration
was increased for the remaining assays.

The pre-incubated mix was homogenized into 2 mL of
molten top agar (0.6% agar, 0.5% NaCl, 0.5 mM biotin, and
0.05 mM L-histidine) and subsequently distributed homoge-
neously on the surface of minimal agar plates. The plates
were incubated at 37°C for 48 h, and the number of reverting colonies in each plate was counted. The criteria for considering carbamates as mutagenic were (1) a dose-dependent increase in the number of reverting colonies or (2) the number of induced reverting colonies greater than or equal to double the amount of reverting colonies found in the negative (0 mg/mL) control plates [21, 22].

Positive controls (known specific mutagens) and negative controls were included in each assay. The specific positive controls were 2-nitrofluorene (10 μg/plate for TA98 without S9), sodium azide (1 μg/plate for TA100 and TA1535 without S9), picrolonic acid (100 μg/plate for TA1537 without S9), 4-Nitroquinoline-N-oxide (1 μg/plate for TA102 without S9), and 2-aminoanthracene (2 μg/plate for all strains with S9). All the mutagens were previously diluted in DMSO except for sodium azide, which was dissolved in water. The negative control for all strains was phosphate buffer at pH 7.4 with 2% DMSO [23].

3. Results

3.1. Profiling. Table 2 shows the profiling results of LQM 919 and LQM 996. The qualitative analysis did not show structural alerts that indicated the ability of these ethyl-carbamates to either produce damage or bind to biomolecules.

3.2. Mutagenicity Predictions. Table 3 shows the endpoint predictions of mutagenicity in six strains of S. typhimurium. The ethyl-carbamates studied were negative (p < 0.05) for the mutagenicity endpoint in all strains studied with and without simulation of metabolic activation using S9.

3.3. Carcinogenicity Predictions. The read across predictions of carcinogenicity in rats were negative for LQM 919 (p < 0.05, moderate confidence) as well as for LQM 996 (p < 0.001, strong confidence). Figure 1 shows the summary carcinogenicity predictions from LQM 919 and neighbours (analogues) used for read-across analysis (a similar result was observed for the LQM 996).

3.4. In vitro Ames’ Assay. Tables 4 and 5 show the average number of reverting colonies in strains exposed to different LQM 919 and LQM 996 concentrations with and without S9 metabolic activation, respectively. The number of spontaneous reverting colonies at all the concentrations used for both carbamates did not show a concentration-dependent increase; moreover, this number was not greater than or equal to double the number of spontaneous reverting colonies of each strain in the respective negative control. In general, greater concentrations of these ethyl-carbamates resulted in thinning out of background growth.

4. Discussion

The genotoxicity induced by pesticides is considered one of the most severe undesirable effects in mammals. When a pesticide reacts with nuclear DNA, it is considered potentially mutagenic and carcinogenic to the exposed organisms; therefore, its use in therapeutics may be limited. For the aforementioned reasons, when the use of new pesticides is proposed, their potential as a possible mutagen and carcinogen should be evaluated before being widely used in the environment. In the present study, in silico predictive analyses and studies on in vitro effects showed that the evaluated ethyl-carbamates do not present a mutagenic or carcinogenic risk.

Most mutagens and carcinogens or their metabolites covalently bind to cellular macromolecules such as DNA and proteins before producing changes in their structure, sequence, or function [24]. Some ethyl-carbamates such as urethane reportedly bind to liver cell DNA before producing tumors in mouse liver or to lung cell DNA before producing tumors in rat lungs [25, 26]. In this study, read-across-analysis revealed no DNA-binding or protein-binding alerts in the molecular structure of LQM 919 and LQM 996. These results suggest that these ethyl-carbamates have a low probability of affecting DNA replicative processes that could alter its sequence.

High levels of estrogen have been associated with tumor production both in experimental models and in epidemiological studies in humans [27–29]. One of the mechanisms of estrogen carcinogenicity is its binding to the specific nuclear receptor ER-α, exerting a potent stimulus on cell proliferation through increased production of growth factors [30]. Some pesticides such as Acetochlor, Dicofol, Endosulfan, Propoxur, and Lindane mimic estrogen activity in terms of structure and function (xenoestrogens) as these can bind estrogen receptors and stimulate tumor production [31, 32]. In silico analysis did not show structural alerts that indicate
probable estrogen receptor binding in the ethyl-carbamates studied; therefore, the risk that they can induce cancer by this mechanism is low. In addition to the above, the read across prediction of carcinogenicity in rats was negative for both ethyl-carbamates; therefore, their chemical structure does not seem to present a risk of cancer induction either. However, other alerts indicated possible toxicity. According to the Cramer classification (estimation of the threshold of toxicological concern) these compounds were categorized within class III, which includes compounds with a chemical structure that does not allow an initial impression of safety, and suggests significant toxicity. The US-EPA New Chemical Categories prediction indicated that these are neutral organic compounds related to probable environmental toxicity for some aquatic species [24].

The Ames’ assay is used worldwide to determine the mutagenic capability of pesticides [33, 34]. This test is highly sensitive, and depending on the strain of S. typhimurium
### Table 4: Mutagenicity of carbamate LQM 919 towards *Salmonella typhimurium* TA98, TA100, TA1535, TA1537 and TA102 strain with or without S9 metabolic activation.

| Concentration (mg/mL) | TA98  | TA100 | TA1535 | TA1537 | TA102 |
|-----------------------|-------|-------|--------|--------|-------|
|                       | -S9   | +S9   | -S9    | +S9    | -S9   |
| 0 (-)                 | 18.5 ± 3.2 | 27.7 ± 2.5 | 104.1 ± 10.8 | 120 ± 9 | 9.1 ± 3 |
| 0.1                   | 16.3 ± 4.9 | 31 ± 6.2 | 112 ± 20.7 | 133.3 ± 10.5 | 10 ± 1.7 |
| 0.2                   | 20.3 ± 4.1 | 28.7 ± 7.1 | 107.6 ± 15.8 | 118.6 ± 4.5 | 6 ± 3.4 |
| 0.4                   | 16.3 ± 2 | 30 ± 7.5 | 125.6 ± 19.4 | 89.6 ± 19.8 | 11 ± 2.6 |
| 0.5                   | 20.3 ± 4.5 | 32.3 ± 8.6 | 78 ± 2 | 106.3 ± 16.8 | 4 ± 3.4 |
| 1                     | 10 ± 5.2 | 25.6 ± 5.1 | 35 ± 22.9 | 28.3 ± 19.4 | 5 ± 1 |
| Limit value:          | 37 | 55.4 | 208.2 | 240 | 18.2 |
| Positive controls:    |       |       |       |       |       |
| 2-NF                  | 2017 ± 460 |       |       |       |       |
| Sodium azide          |       |       | 1666 ± 99 |       | 731 ± 77.2 |
| Picrolonic acid       |       |       |        | 48.33 ± 25.1 |       |
| 4-NQO                 |       |       |        |        | 1573 ± 61 |
| 2-AA                  | 1831 ± 163 | 2277 ± 146 | 765 ± 45 | 441 ± 99 | 2050 ± 93 |

*S9 = Microsomal fraction of liver of rats treated with phenobarbital and beta-naphthoflavone enriched with NADP and glucose-6-P; 2-NF = 2-nitrofluorene; 4-NQO = 4-nitroquinilino N-oxide; 2-AA = 2-aminoanthracene.*
**Table 5:** Mutagenicity of carbamate LQM 996 towards *Salmonella typhimurium* TA98, TA100, TA1535, TA1537 and TA102 strain with or without S9 metabolic activation.

| Concentration (mg/mL) | -S9  | +S9  | -S9  | +S9  | -S9  | +S9  | -S9  | +S9  | -S9  | +S9  | -S9  | +S9  | -S9  | +S9  |
|-----------------------|------|------|------|------|------|------|------|------|------|------|------|------|------|------|
| 0 (-)                 | 18.5 ± 3.2 | 27.7 ± 2.5 | 104.1 ± 10.8 | 120 ± 9 | 9.1 ± 3.9 | 16.5 ± 1.9 | 9.1 ± 3 | 14.6 ± 3 | 319.5 ± 75.7 | 330 ± 66 |
| 0.1                   | 20.3 ± 4.1 | 28.3 ± 6.1 | 94.6 ± 27.7 | 92.6 ± 5.5 | 8.3 ± 3.2 | 7.6 ± 4.6 | 7.3 ± 3.2 | 8.3 ± 1.1 | 395.3 ± 6.1 | 473 ± 17.5 |
| 0.2                   | 20.0 ± 5  | 30.3 ± 4.0 | 96 ± 3   | 101.3 ± 12.3 | 4 ± 1.7 | 10.6 ± 3.5 | 4.6 ± 2  | 8.3 ± 2.5 | 251.6 ± 49.8 | 419.6 ± 31.6 |
| 0.4                   | 17 ± 4    | 22.6 ± 8  | 83.6 ± 17.6 | 95 ± 4.3 | 2.6 ± 1.5 | 11.6 ± 1.5 | 1.6 ± 2.8 | 5.3 ± 3  | 241.3 ± 49.1 | 367.3 ± 58.2 |
| 0.5                   | 18.3 ± 4  | 30.3 ± 1.1 | 96.6 ± 12.5 | 81.6 ± 16.6 | 2.6 ± 2.3 | 4.6 ± 3.2 | 0      | 0      | 194.3 ± 43  | 335.6 ± 18 |
| 1                     | 13 ± 1.7  | 21.3 ± 3  | 34.6 ± 26.2 | 0    | 0      | 6.6 ± 2.8 | 0      | 0      | 150 ± 16.7  | 185.3 ± 29.2 |
| Limit value:          | 37 ± 5.4  | 208.2 ± 240 | 18.2 ± 33 | 18.2 ± 29.2 | 639 ± 66 |
| Positive controls:    | 2-NF 2017 ± 460 | Sodium azide 1666 ± 99 | TA100 731 ± 77.2 | TA1535 48.33 ± 25.1 | 4-NQO 1573 ± 61 |
| 4-AA                  | 1831 ± 163 | 2277 ± 146 | 765 ± 45 | 441 ± 99 | 2050 ± 93 |

*S9 = Microsomal fraction of liver of rats treated with phenobarbital and beta-naphthoavone enriched with NADP and glucose-6-P; 2-NF = 2-nitrofluorene; 4-NQO = 4-nitroquinolino N-oxide; 2-AA = 2-aminoanthracene.*
used, it can reveal different types of mutations. Strains TA98, TA1535, and TA1538 are sensitive to frameshift mutations, strains TA97, TA100, and TA1535 are sensitive to C/G base pair exchanges, and strain TA102 is sensitive to A/T base pair exchanges and mutations caused by oxidative damage [22, 35]. In this study, the results of read-across analysis predicted that the evaluated ethyl-carbamates have no mutagenic potential in any of the six strains of *S. typhimurium* studied. These predictions were consistent with the results obtained in the Ames *in vitro* assay, and in both cases, metabolic activation of ethyl-carbamates was considered. Some carbamates such as Propoxur, Carbaryl, and Methomyl acquire mutagenic activity when subjected to exogenous metabolic activation [36, 37]. We did not observe differences in spontaneous reversion induced by the ethyl-carbamates under study with or without exogenous metabolic activation. The *in silico* and *in vitro* results obtained in this study indicated the absence of mutagenic activity in the ethyl-carbamates or their metabolites.

In previous studies, we observed that the ethyl-carbamates used in this study induce an increase in micronucleated reticulocytes in rats, which could be the result of DNA damage [10]. Furthermore, we observed an increase in thiobarbituric acid reactive species (oxidative stress) that can cause damage to macromolecules including DNA [8]. However, the results of the present study suggest that these ethyl-carbamates did not represent a risk for cancer in mammals.

### 5. Conclusions

*In silico* studies predicted that the ethyl-carbamates LQM 919 and LQM 996 are negative for mutagenicity and carcinogenicity. The *in vitro* Ames test confirmed the absence of mutagenic capacity in these compounds.

### Data Availability

The data used to support the findings of this study are available from the corresponding author upon request.

### Conflicts of Interest

The authors declare that there is no conflict of interest regarding the publication of this paper.

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