

Thionate versus Oxon: Comparison of Stability, Uptake, and Cell Toxicity of (¹⁴CH₃O)₂-Labeled Methyl Parathion and Methyl Paraoxon with SH-SY5Y Cells

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The stability, hydrolysis, and uptake of the organophosphates methyl parathion and methyl paraoxon were investigated in SH-SY5Y cells. The stabilities of (¹⁴CH₃O)₂-methyl parathion (¹⁴C-MPS) and (¹⁴CH₃O)₂-methyl paraoxon (¹⁴C-MPO) at 1 μM in culture media had similar half-lives of 91.7 and 101.9 h, respectively. However, 100 μM MPO caused >95% cytotoxicity at 24 h, whereas 100 μM MPS caused 4–5% cytotoxicity at 24 h (~60% cytotoxicity at 48 h). Greater radioactivity was detected inside cells treated with MPO as compared to MPS, although >80% of the total MPO uptake was primarily dimethyl phosphate (DMP). Maximum uptake was reached after 48 h of ¹⁴C-MPS or ¹⁴C-MPO exposure with total uptakes of 1.19 and 1.76 nM/10⁶ cells for MPS and MPO, respectively. The amounts of MPS and MPO detected in the cytosol after 48 h of exposure time were 0.54 and 0.37 nM/10⁶ cells, respectively.

KEYWORDS: Uptake; radiolabel; methyl parathion; methyl paraoxon; SH-SY5Y human neuroblastoma cells; hydrolysis; MTT

INTRODUCTION

Organophosphorus (OP) insecticides **1** are a relatively safe group of agricultural chemicals used extensively in plant and crop protection and include malathion, chlorpyrifos, parathion, methyl parathion, and diazinon (**Figure 1**). Most parent OP insecticides contain the phosphorothionate group (P=S), which renders the structure relatively safe to mammals owing to its poor reactivity with target enzymes and other biomolecules. The thionate linkage also confers hydrolytic stability owing to increased electron density at phosphorus. OP insecticides can be converted from thionates **1** to oxons **2** (**Figure 1**) either in the environment or in vivo and become reactive and potentially toxic following occupational or incidental exposures by direct contact with air, food, and water (*1*). The poor reactivity of thionates is in contrast to that of organophosphorus warfare agents that contain the P=O linkage (oxons), are highly reactive, and induce rapid toxicity.

The primary mechanism of action of OP insecticides is based on their ability to inhibit acetylcholinesterase (AChE, EC 3.1.1.7) as the reactive oxon metabolite form **2** (**Figure 1**) (*2*). Methyl parathion (*O,O*-dimethyl *O*-4-nitrophenyl phosphorothioate) is representative of the thionate (P=S) class of insecticides with insect and mammal toxicity due to inhibition of AChE (*2, 3*). Parathion itself is an inherently weak cholinesterase inhibitor with an IC₅₀ in the range of 10⁻⁴–10⁻⁵ M (*4*), but its biotransformation to the reactive metabolite paraoxon (*5–7*) affords a

potent inhibitor of AChE (IC₅₀ = 10⁻⁸ M) (*8*). Inhibition of AChE by paraoxon, as for most reactive OPs, occurs mechanistically with loss of the Z group (**Figure 1**) and results in accumulation of acetylcholine in cholinergic synapses and excessive stimulation of cholinergic pathways in the central and peripheral nervous systems (*9–11*). Although AChE is the primary target and the principal mechanism underlying toxic action, highly reactive small molecules such as methyl paraoxon and ethyl paraoxon can potentially modify a number of biomolecules. Paraoxon toxicity at the cellular level has been shown in immortal cell lines in vitro (*12–17*). Subcellular targets for the initiation of cytotoxicity have not been fully elucidated, but nuclear (*18*), enzymatic (*14*), cytoskeletal (*19*), and plasma membrane (*20*) alterations have been described, and a number of alternative protein targets have been identified (*21, 22*).

One significant question that remains unanswered about OP insecticide toxicity is if there are differences in the ability or rate of the structurally similar thionate and oxon forms to penetrate cells. If this question could be answered, investigators could gain a clearer understanding of the individual or interdependent role of thionate and oxon forms of an insecticide to modify intracellular or extracellular protein targets and alter biochemical pathways that lead to toxicity. Because the conversion of thionate to oxon occurs with most thionate OP insecticides, the study of cell membrane penetration is of broad and significant impact. A differential study of cell penetration is further warranted to address a number of hypotheses regarding cell culture studies with various thionate–oxon structures. For example, toxic effects due to OP exposure likely begin with covalent modification

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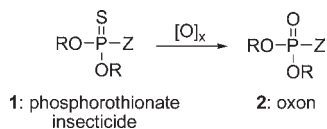


Figure 1. Structures of representative phosphorothionate insecticides and the corresponding oxons: malathion (R = Me, Z = -SCH(CO₂Et)CH₂-CO₂Et); parathion (R = Et, Z = -OPh-*p*-NO₂); methyl parathion (R = Me, Z = -OPh-*p*-NO₂); chlorpyrifos (R = Et, Z = -O-[3,5,6-trichloro-2-pyridyl]); and diazinon (R = Et, Z = -O-[2-isopropyl-6-methyl-4-pyrimidinyl]).

(adduction) of a protein by the oxon (21). Owing to the relatively high reactivity and hydrolytic instability, it is presumed that OP oxons would not be found at appreciable concentrations within cells as compared to the thionate owing to lower lipophilicity. Support for this supposition is indicated in calculations of partition coefficients in which methyl parathion (CLog *P* = 2.79; Log *P*_{o/w} = 3.0) (23) and methyl paraoxon (CLog *P* = 1.38; Log *P*_{o/w} = 1.6 for ethyl paraoxon) (24) show a 20-fold difference, suggesting that the thionate could more readily penetrate cell membranes by passive diffusion mechanisms. However, alternative mechanisms could enable oxons access to intracellular targets.

Given the reactivity, aqueous instability, and lower CLog *P* of oxons as compared to thionates, it has been presumed that cell penetration is somewhat limited. However, it is important to recognize that OP insecticide toxicity represents exposure to both thionate and oxon forms, and an understanding of cell penetration by these structures is important to validate and identify new protein targets, their localization, and distribution. Moreover, the fractional concentration of extracellular and intracellular amounts of thionate and oxon may play an important role in the mode of toxic action. In this paper, the stability, uptake, and hydrolysis of doubly labeled methyl parathion and methyl paraoxon were conducted in SH-SY5Y cells so that the key structural difference between OP insecticide (thionate) and its primary metabolite (oxon) could be better understood. The rationale for the double label was twofold: (1) placement of radiolabel at the methoxy group ensures that the isotope remains attached to proteins because the *p*-nitrophenoxy group is lost upon phosphorylation; (2) two methoxy ester radiolabels were deemed important because some proteins or biomolecules may be prone to aging or loss of a second ester group following phosphorylation. The SH-SY5Y cell line was chosen for study because it expresses AChE, a number of neuron-specific enzymes and biosynthetic pathways, dopamine hydroxylase, tyrosine hydroxylase, aromatic L-amino acid decarboxylase, enzymes unique to catecholamine neurons, and the nicotinic acetylcholine receptor (25).

MATERIALS AND METHODS

General. ¹⁴C-Methyl parathion (¹⁴C-MPS) was purchased from Perkin-Elmer, Boston, MA (NEN radiochemicals) with a radioactivity of 1 mCi (specific activity = 41.65 mCi/mmol). Liquid scintillation counting (LSC) was performed to determine the radiocarbon content of various samples using a Tricarb 2900 liquid scintillation counter. Silica gel G₂₅₄ thin layer chromatography plates (10 mm thickness; Analtech) were used for purification. Radioactive images of TLC plates were recorded using a phosphorimager (Fujifilm FLA3000). The MTT assay kit was obtained from Roche Applied Science, and the absorbance was measured using a microplate reader (Molecular Devices, Versamax).

Synthesis of ¹⁴C-Methyl Paraoxon (¹⁴C-MPO). ¹⁴C-Methyl paraoxon (¹⁴C-MPO) was prepared by oxidation of ¹⁴C-methyl parathion (¹⁴C-MPS). To a solution of ¹⁴C-MPS (1 mCi with a specific activity of 41.65 mCi/mmol) in dry methylene chloride (3 mL) was added *m*-chloroperoxybenzoic acid (1.5 mmol) at 0 °C, and the reaction mixture was stirred at room temperature for 2 h or until completion of reaction was confirmed by TLC. The reaction mixture was loaded onto a preparative

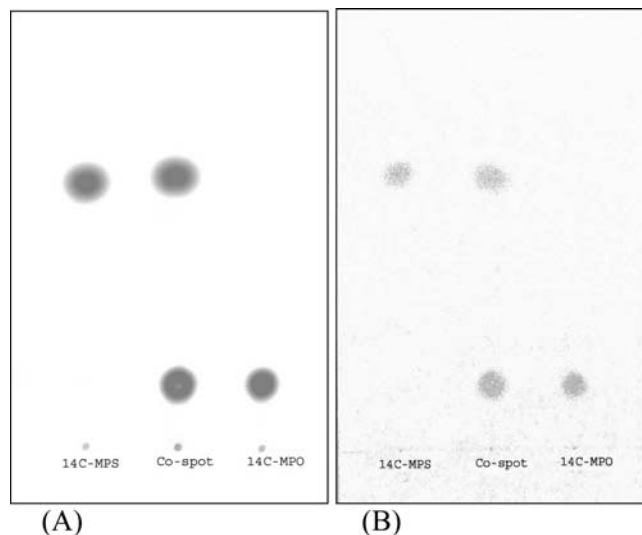


Figure 2. Thin layer chromatographic analyses of ¹⁴C-MPS and ¹⁴C-MPO (EtOAc/hex, 1:1): (A) radioactivity image; (B) UV image. *R*_f of MPS = 0.65; *R*_f of MPO = 0.18.

silica gel G₂₅₄ TLC plate and eluted with EtOAc/hex (1:1). The band of ¹⁴C-MPO at *R*_f 0.15–0.20 (*R*_f of ¹⁴C-MPS is 0.60–0.65) was correlated with the migration of cold paraoxon standard, removed, extracted with methylene chloride (5 mL × 3), and filtered, and the solvent was evaporated to afford the radiolabeled product (0.360 mCi with a specific activity of 41.65 mCi/mmol). Total counts (cpm) were converted to dpm using dpm = cpm/detector efficiency. The radiochemical yield of synthesized ¹⁴C-MPO was determined using this relationship: 1 mCi = 2.2 × 10⁹ dpm. Chemical and radiochemical purities for both ¹⁴C-methyl parathion and ¹⁴C-methyl paraoxon were >98% based on total recovery from the TLC plate (Figure 2).

SH-SY5Y Cell Culture. SH-SY5Y cells (a human neuroblastoma cell line) were obtained from American Type Culture Collection (Rockville, MD) and cultured in DMEM/F12 medium (GIBCO BRL, Grand Island, NY) supplemented with 10% fetal bovine serum (Hyclone), 100 U/mL penicillin, 100 μg/mL streptomycin, and 2 mM L-glutamine in a CO₂ incubator maintained at 5% CO₂ and 37 °C. The medium was changed every 2 days. Cells were allowed to reach 80% confluence before exposure to ¹⁴C-MPS or ¹⁴C-MPO.

3-(4,5-Dimethylthiazol-2-yl)-2,5-diphenyltetrazolium Bromide (MTT) Viability Assay. Approximately, 0.25 × 10⁵ cells/well were seeded into 96-well plates and exposed for 24, 48, or 72 h to MPS and MPO at concentrations from 10 nM to 100 μM prepared as solutions in acetonitrile (0.1% v/v). A 2% Triton X-100 solution in assay medium was used for a positive control (*n* = 8 for each MPS and MPO concentration and Triton X-100). Following exposure, cells were rinsed several times with culture medium prior to MPS and MPO exposure. Culture medium was removed, and 100 μL of fresh medium containing the graded MPS and MPO concentrations or Triton X-100 was added to each well. After incubation for appropriate time points, 10 μL of MTT labeling reagent was added to each well. Plates were incubated with MTT labeling reagent for 4 h, and then 100 μL of solubilizing solution was added to each well and incubated overnight. Absorbance of samples was measured using a microplate reader at 575 nm (formazan) using a reference wavelength at 675 nm. Viability was determined by comparing the absorbance readings of the wells containing the OP-treated cells with those of the vehicle (0.1% acetonitrile)-treated cells (Figure 3).

Stability of ¹⁴C-MPS and ¹⁴C-MPO in Culture Media. A 1 μM solution of ¹⁴C-MPS or ¹⁴C-MPO in DMEM/F12 medium 1% fetal bovine serum (Hyclone; 100 U/mL penicillin, 100 μg/mL streptomycin, and 2 mM L-glutamine) was incubated at 37 °C for 120 h. At different time points (0–120 h), samples (50 μL × 3) were loaded onto preparative silica gel G₂₅₄ TLC plates and developed with EtOAc/hex (1:1). Cold MPS and MPO were spotted as elution standards at the corner of the plate for reference. For ¹⁴C-MPS, three bands (*R*_f = 0.65, 0.18, and <0.1), and for ¹⁴C-MPO, two bands (*R*_f = 0.18 and <0.1), were isolated and the total

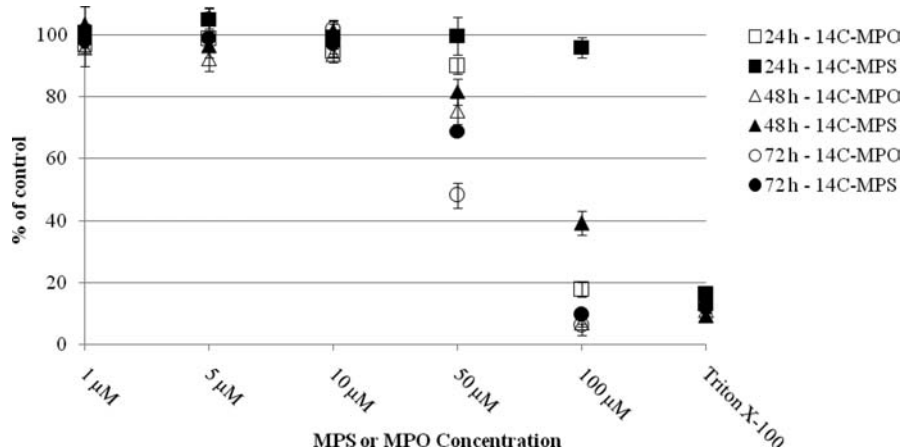


Figure 3. Cytotoxic effect of ^{14}C -MPS and ^{14}C -MPO on SH-SY5Y cells: (squares) SH-SY5Y cell viability following 24 h exposure to ^{14}C -MPS and ^{14}C -MPO; (triangles) SH-SY5Y cell viability following 48 h exposure to ^{14}C -MPS and ^{14}C -MPO; (circles) SH-SY5Y cell viability following 72 h exposure to ^{14}C -MPS and ^{14}C -MPO. Cells were treated with various concentrations of ^{14}C -MPS and ^{14}C -MPO for 24, 48, and 72 h. Cell viability was determined by 3-[4,5-dimethylthiazol-2-yl]-2,5-diphenyl tetrazolium bromide (MTT) assay. Results are presented as percent of vehicle control, determined by comparing the absorbance readings of the wells containing the OP-treated cells with those of the vehicle (0.1% acetonitrile)-treated cells. Data represent the mean \pm SEM ($n = 8$). Triton X-100 was used as a positive control (= 100% cell death). Solid symbols are used for MPS and open symbols for MPO.

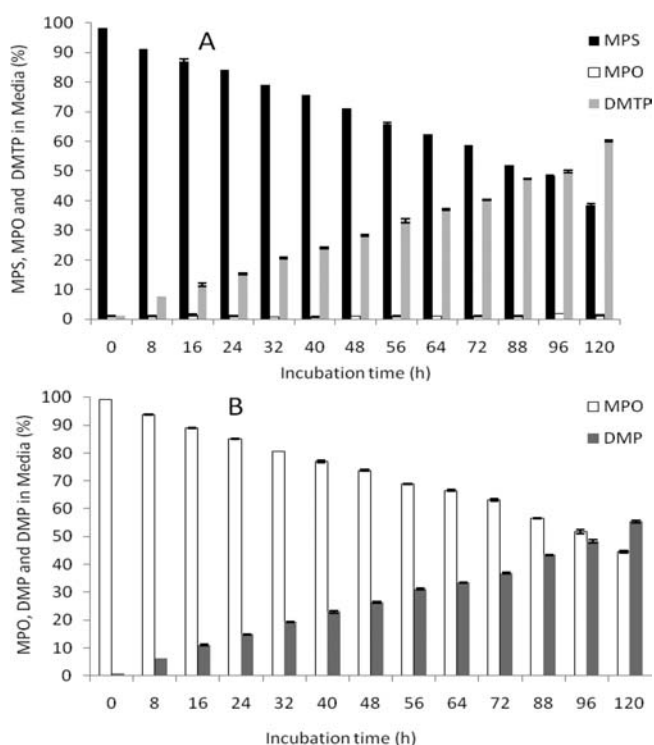


Figure 4. Stability of radiolabeled MPS or MPO in 1% FBS culture media: (A) ^{14}C -MPS; (B) ^{14}C -MPO. Data represent mean \pm SEM ($n = 3$). ^{14}C -DMTP and ^{14}C -DMP are primary metabolites of ^{14}C -MPS and ^{14}C -MPO hydrolysis.

counts (cpm) recorded. The percentage of ^{14}C -MPS or ^{14}C -MPO at each time point was calculated and plotted against incubation time (Figure 4). Rate constants for the degradation of MPS and MPO were determined by calculating the negative slope of $\ln(\%$ of MPS/MPO remaining) versus incubation time.

Uptake of ^{14}C -MPS and ^{14}C -MPO in SH-SY5Y Cells. ^{14}C -MPS and ^{14}C -MPO stock solutions were prepared in acetonitrile (500 μM) and stored at 0 $^{\circ}\text{C}$. ^{14}C -MPS and ^{14}C -MPO treatment concentrations (1 μM) were prepared by dilution of the stock solution in DMEM/F12 medium 1% fetal bovine serum (Hyclone), 100 U/mL penicillin, 100 $\mu\text{g}/\text{mL}$ streptomycin, and 2 mM L-glutamine (400 μL of stock solution into 200 mL of media to make 1 μM ^{14}C -MPS/ ^{14}C -MPO media solution).

In preliminary experiments, cells were exposed to 1 μM ^{14}C -MPS/ ^{14}C -MPO at 37 $^{\circ}\text{C}$ for 0, 8, 16, 24, 48, and 72 h in a 96-well plate (2 μL of medium in each well). At the determined time points, culture medium was removed and the cells were washed with ice-cold PBS (3 \times 1 mL) to remove residual ^{14}C -MPS or ^{14}C -MPO and placed in a scintillation vial. To the remaining washed cells was added 5 mL of ice-cold PBS, and the cells were removed using a cell scraper and transferred to centrifuge tubes. Each well was washed an additional time with PBS to remove residual cells. Cells were centrifuged for 5 min, and supernatant was transferred to a scintillation vial. Cells were treated with lysis buffer containing 20 mM Tris-HCl (pH 7.5), 150 mM NaCl, 1 mM Na_2EDTA , 1 mM EGTA, 1% Triton X-100, 2.5 mM sodium pyrophosphate, 1 mM β -glycerophosphate, and 1 mM Na_3VO_4 (Cell Signaling Technology, Beverly, MA). Additions to the lysis solution were made to give final concentrations of 0.5% sodium deoxycholate, 0.5% SDS, 1 μM okadaic acid, 1 mM phenylmethanesulfonyl fluoride (PMSF), 0.1 mg/mL benzamide, 8 $\mu\text{g}/\text{mL}$ calpain inhibitors I and II, and 1 $\mu\text{g}/\text{mL}$ each leupeptin, pepstatin A, and aprotinin. Lysis buffer (500 μL) was added to the cell pellet and vortexed. After 30 min, the cell lysate was centrifuged, and the supernatant containing the cytosolic fraction was separated. Scintillation fluid was added to all samples to obtain a standard volume and total counts (cpm) were recorded. Data were collected in triplicate for each time point. The total amount of OP inside cells was calculated using the following equation and was plotted against OP exposure time (Figure 5).

$$\begin{aligned} & \text{total amount of OP inside cells (nM)} \\ &= \left[\frac{\text{total radioactivity inside cells (cpm)}}{\text{initial OP concentration (nM)}} \right] / \\ & \left[\frac{\text{total radioactivity in medium (cpm)}}{\text{total radioactivity inside cells (cpm)}} \right] \end{aligned}$$

Fate of ^{14}C -MPS and ^{14}C -MPO in Uptake Experiments. To determine the fate of ^{14}C -MPS and ^{14}C -MPO in media during uptake experiments and the amounts of MPS/MPO and their degradation products in cell cytosol, uptake experiments were performed at 24, 48, and 72 h in a Petri dish plate (20 mL of medium in each plate; 1 μM ^{14}C -MPS or ^{14}C -MPO) using a protocol similar to that described in the previous section. After cell harvesting, total cell numbers were calculated by trypan blue counting. After cell lysis, the cytosolic fraction was collected and was loaded on preparative silica gel G_{254} TLC plate, and reference standards of cold MPS and MPO were spotted at the corner of the plate. TLC plates were eluted using EtOAc/hex (1:1). The radioactive bands were identified by correlation with standards, removed, and collected in individual liquid scintillation vials. Similarly, 100 μL of medium was also

loaded on TLC plate, and bands were collected. Scintillation fluid was added to all samples to obtain a standard volume, and total counts (cpm) were recorded. Data were collected in triplicate for each time point. Concentrations of OP in cell cytosol per million cells were calculated using the following equation and plotted against OP exposure time (Figure 6).

$$\text{nM OP in cytosol}/10^6 \text{ cells} = \frac{[\text{radioactivity for OP or its degradation product in cytosol (cpm)} \times \text{initial OP concentration (nM)}] / \{[\text{total radioactivity in media (cpm)} + \text{total radioactivity inside cells (cpm)}] \times \text{number of cells (in millions)}\}}$$

The partition coefficient values (media/cell) for the OPs and their degradation products were calculated by dividing the amount of $^{14}\text{C-MPS}/^{14}\text{C-MPO}$ in cell cytosol by the amount in the medium at 48 h

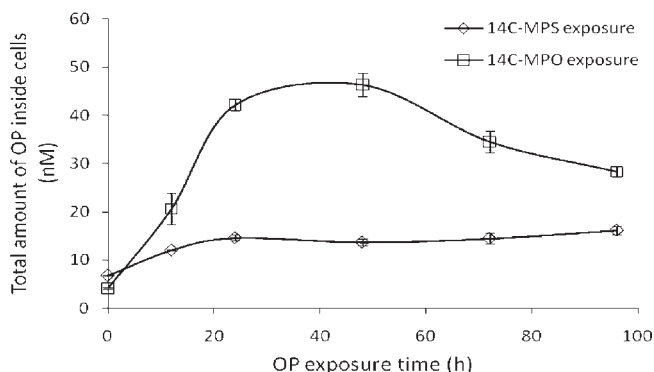


Figure 5. Uptake of $^{14}\text{C-MPS}$ or $^{14}\text{C-MPO}$ into SH-SY5Y cells at different exposure times (0, 12, 24, 48, 72, and 96 h). Data represent mean \pm SEM ($n = 3$).

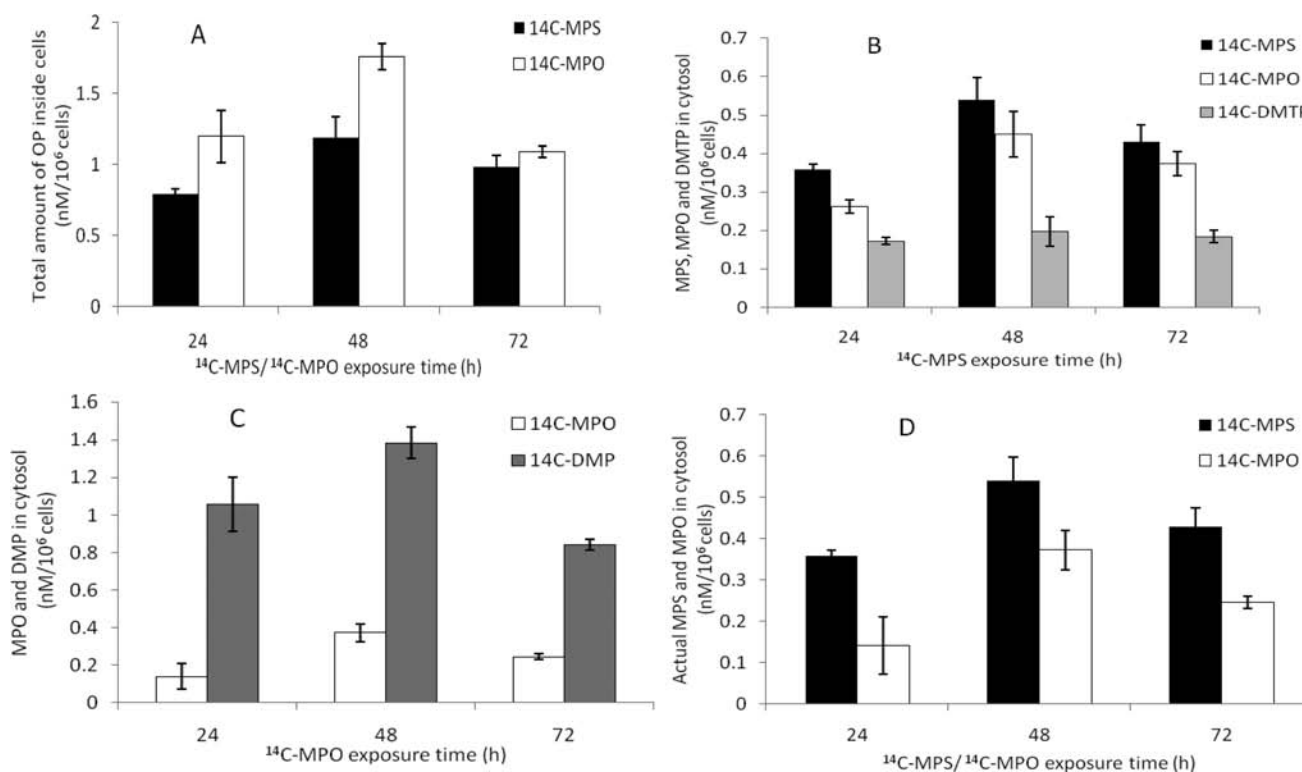


Figure 6. Uptake of $^{14}\text{C-MPS}$ or $^{14}\text{C-MPO}$ in SH-SY5Y cells at different exposure times: (A) total radioactivity (OP and their degradation products) found inside cells following exposure to $1 \mu\text{M}$ MPS or MPO; (B) uptake of $^{14}\text{C-MPS}$ ($1 \mu\text{M}$) and its degradation products; (C) uptake of $^{14}\text{C-MPO}$ ($1 \mu\text{M}$) and its degradation product; (D) Actual amount of $^{14}\text{C-MPS}/^{14}\text{C-MPO}$ found inside cells at $1 \mu\text{M}$. Cell numbers per experiment were normalized and uptake values are indicated for 10^6 cells. Data represent mean \pm SEM ($n = 3$). $^{14}\text{C-DMTP}$ and $^{14}\text{C-DMP}$ are primary metabolites of $^{14}\text{C-MPS}$ and $^{14}\text{C-MPO}$ hydrolysis.

of exposure time. The uptake rate constant was calculated from the negative slope of the plot between $\ln(\text{nM increment of OP in cytosol})$ versus OP exposure time (h).

RESULTS

Synthesis of $^{14}\text{C-Methyl Paraoxon}$ ($^{14}\text{C-MPO}$). $^{14}\text{C-Methyl}$ parathion (1 mCi) with a specific activity of 41.65 mCi/mmol was oxidized with *m*-chloroperoxybenzoic acid (*m*-CPBA) in methylene chloride to yield $^{14}\text{C-MPO}$. Purification was conducted using preparative thin layer chromatography (prep-TLC). The radioactive yield of $^{14}\text{C-MPO}$ was 0.360 mCi , and the radiochemical purities of both $^{14}\text{C-MPS}$ and $^{14}\text{C-MPO}$ were $>98\%$ (Figure 2). Radioactive MPS and MPO structures were confirmed by coelution on TLC with standard (cold) MPS and MPO, which were characterized by NMR and MS.

Viability of SH-SY5Y Cells Exposed to $^{14}\text{C-MPS}$ and $^{14}\text{C-MPO}$. SH-SY5Y cells were treated with $^{14}\text{C-MPS}$ or $^{14}\text{C-MPO}$ ranging from 10 nM to $100 \mu\text{M}$, and the cell viability was assessed by MTT assay. Some cytotoxicity was observed with both MPS and MPO at $50 \mu\text{M}$ at time points of 48 and 72 h, and at least 60% cytotoxicity occurred at all time points when the cells were exposed to $100 \mu\text{M}$ except MPS treatment for 24 h (Figure 3), which showed no significant decrease in viability. At $100 \mu\text{M}$ concentration, MPS showed 4% (24 h), 60% (48 h), and 90% (72 h) cytotoxicity, respectively, whereas MPO showed $>80\%$ cytotoxicity after just 24 h.

Stability of $^{14}\text{C-MPS}$ and $^{14}\text{C-MPO}$ in Culture Media. The stability of $^{14}\text{C-MPS}$ and $^{14}\text{C-MPO}$ in culture media was determined by incubating $1 \mu\text{M}$ OP in tissue culture media at 37°C for up to 120 h. The medium was analyzed at various time points by prep-TLC, and the amount of OP and degradation products was quantified by radioactivity counting of isolated bands. The major degradation pathways for $^{14}\text{C-MPS}$ and $^{14}\text{C-MPO}$ are oxidation/hydrolysis

and hydrolysis, respectively (**Figure 7**), forming the water-soluble DMTP and DMP as primary metabolites. The formation of DMTP and DMP from paraoxon and like compounds is well documented (26, 27). ^{14}C -MPS and ^{14}C -MPO showed similar stability profiles. After 72 h of incubation in medium, > 50% of ^{14}C -MPS (**Figure 4A**) and ^{14}C -MPO (**Figure 4B**) were recovered from the medium unchanged. Both ^{14}C -MPS and ^{14}C -MPO hydrolyzed with loss of the *p*-nitrophenol group to yield dimethyl thiophosphoric acid (thiophosphate) and dimethylphosphate (phosphoric acid) (**Figure 7**), respectively, as determined by coelution with standards. ^{14}C -MPS was found to be relatively stable to oxidation in media and formed < 1% ^{14}C -MPO after 4 days (**Figure 4A**). On the basis of the isolation and measurement of the eluted bands, 80–97 and 78–95% of the originally applied amount of ^{14}C -MPS and ^{14}C -MPO were recovered. The rate constants for degradation (hydrolysis) of ^{14}C -MPS and ^{14}C -MPO were calculated to be 7.0×10^{-3} and $6.0 \times 10^{-3} \text{ h}^{-1}$, respectively. The half-lives for ^{14}C -MPS and ^{14}C -MPO in media were 91.7 and 101.9 h, respectively.

Uptake of ^{14}C -MPS and ^{14}C -MPO in SH-SY5Y Human Neuroblastoma Cells. Using the cell viability and media stability data, time points up to 96 h and OP concentrations of $1 \mu\text{M}$ were chosen for uptake studies because little to no loss of cells was observed under these conditions. In preliminary uptake experiments, cells were exposed to $1 \mu\text{M}$ ^{14}C -MPS and ^{14}C -MPO for different exposure times, namely, 0, 8, 12, 24, 48, 72, and 96 h. A comparative plot of uptake for ^{14}C -MPS and ^{14}C -MPO is shown in **Figure 5**, indicating that MPO achieved a maximum level of 46 nM at the 48 h time point and MPS 13 nM maximum at 24 h.

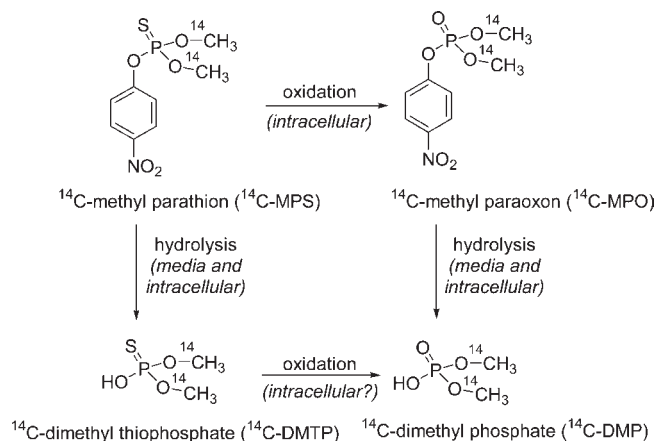


Figure 7. Degradation (hydrolysis) of ^{14}C -methyl parathion and ^{14}C -methyl paraoxon in media.

At the most, only 5% (approximately 8000 cpm; see the Supporting Information) of the total applied radioactivity (170,000 cpm) of either MPS or MPO entered the cells.

To determine the fate of ^{14}C -MPS and ^{14}C -MPO in media during the uptake experiments and the total amount of MPS, MPO, and/or the degradation products in cell cytosol, uptake experiments were performed at 24, 48, and 72 h (**Figure 6**). At each time point, a sample of the cytosolic fraction and the medium was loaded on a preparative TLC plate and the OPs separated and analyzed to determine the stability of ^{14}C -MPS or ^{14}C -MPO and formation of the hydrolysis products during the uptake experiment (**Tables 1 and 2**). ^{14}C -MPS and ^{14}C -MPO were stable in media throughout the uptake experiment. The cell counts for both experiments were normalized, and values of ^{14}C -MPS and ^{14}C -MPO uptake were reported per one million cells. As found in the preliminary uptake experiment (**Figure 5**), the total amount of MPO uptake (sum of MPO and its degradation product) was greater than the total MPS uptake (sum of MPS and its degradation products) (**Figure 6A**). It was interesting to note that in case of ^{14}C -MPO uptake, a large amount of the hydrolyzed product, dimethyl phosphate (80–90% of total radioactivity inside cells) (**Figure 6C**), was found in cells, whereas a greater amount of ^{14}C -MPS was found inside cells compared with its degradation products, MPO and DMTP (**Figure 6B**).

The maximum radioactivity inside cells was reached after 48 h of ^{14}C -MPS/ ^{14}C -MPO exposure with total uptakes of 1.19 and 1.76 nM/ 10^6 cells for MPS and MPO, respectively, whereas the actual amounts of MPS and MPO inside cells after 48 h of exposure time were 0.54 and 0.37 nM/ 10^6 cells (**Table 1**), respectively. On the basis of TLC analysis, in the case of ^{14}C -MPS

Table 2. Relative Ratios of MPS, MPO, DMTP, and/or DMP in Culture Media after Exposure of Cells to $1 \mu\text{M}$ OP (MPS or MPO)^a

incubation time (h)	relative ratios ^b		
	MPS	MPO	DMP and/or DMTP
^{14}C-MPS			
24	74.3 ± 1.0	9.8 ± 1.0	16.0 ± 0.2
48	63.9 ± 0.1	6.8 ± 0.9	29.3 ± 0.9
72	52.9 ± 0.8	10.3 ± 1.4	36.8 ± 1.6
^{14}C-MPO			
24		69.7 ± 3.3	30.3 ± 3.3
48		62.5 ± 0.6	37.5 ± 0.6
72		61.4 ± 0.2	38.6 ± 0.2

^aData represent mean ± SEM ($n = 3$). ^bRatios of OP and their degradation products were determined by preparative TLC analysis.

Table 1. Percentage of MPS, MPO, DMTP, and/or DMP in Cell Cytosol after Exposure of Cells to $1 \mu\text{M}$ OP (MPS or MPO)^a

incubation time (h)	percentage of MPS, MPO, DMTP, and/or DMP in cell cytosol ^b (per 10^6 cells)			relative ratios		
	MPS	MPO	DMP and/or DMTP	MPS	MPO	DMP and/or DMTP
^{14}C-MPS						
24	0.036 ± 0.001	0.026 ± 0.002	0.017 ± 0.001	45.3 ± 1.5	33.0 ± 0.9	21.7 ± 0.6
48	0.054 ± 0.006	0.045 ± 0.006	0.020 ± 0.004	45.8 ± 2.0	37.9 ± 1.7	16.3 ± 1.1
72	0.043 ± 0.004	0.037 ± 0.003	0.018 ± 0.002	43.5 ± 1.8	37.9 ± 2.3	18.7 ± 0.6
^{14}C-MPO						
24		0.014 ± 0.007	0.106 ± 0.014		10.9 ± 4.5	89.1 ± 4.5
48		0.037 ± 0.005	0.139 ± 0.008		21.2 ± 2.3	78.8 ± 2.3
72		0.025 ± 0.002	0.084 ± 0.003		22.5 ± 0.7	77.5 ± 0.7

^aData represent mean ± SEM ($n = 3$). ^bRatios of OP and their degradation products were determined by preparative TLC analysis.

uptake, the ratio of ^{14}C -MPS to that of degradation products inside cells was 46:38:16 (MPS/MPO/DMTP) at 48 h of exposure time, whereas the uptake of MPO showed an MPO/DMO ratio of 21:79 (MPO/DMP) (Table 1).

The amounts of ^{14}C -MPS and ^{14}C -MPO determined inside cells at various exposure times (Figure 6D) show that a larger amount of ^{14}C -MPS entered cells than of ^{14}C -MPO at each time point and only a negligible amount of ^{14}C -MPS or ^{14}C -MPO was found to be associated with the membrane fraction. The percent total recoveries of all label in the uptake experiments for ^{14}C -MPS and ^{14}C -MPO were found to be >92 and >95%, respectively. The cells–media partition coefficients (K_d) were calculated to be 6.3×10^{-4} and 5.9×10^{-4} with uptake rate constants of 0.054 and 0.025 h^{-1} for MPS and MPO, respectively.

DISCUSSION

In this study, the stability, uptake, and cytotoxicity of ^{14}C -methyl parathion and ^{14}C -methyl paraoxon were tested in a SH-SY5Y human neuroblastoma cell line. The radiolabel was placed at both methoxy groups so that attachment of the OP to biomolecules and the formation of hydrolysis products could be tracked more efficiently as compared to studies in which the radiolabel was incorporated at the *p*-nitrophenoxy group (26, 28–30). Synthesis of radiolabeled paraoxon from ($^{14}\text{CH}_3\text{O}$)₂-parathion was accomplished in >98% chemical and radiochemical purity. The specific activity was 41.65 mCi/mmol as determined by chromatography.

Stability studies in culture media indicated that >50% of either ^{14}C -MPS or ^{14}C -MPO remained in the media after 72 h (Figure 4). The degradation pathway expected for ^{14}C -MPS was oxidation/hydrolysis, and for ^{14}C -MPO, hydrolysis was anticipated (Figure 7). However, we did not observe oxidation of ^{14}C -MPS to ^{14}C -MPO in culture media. Thus, hydrolysis, or loss of the *p*-nitrophenoxy group, to yield dimethyl thiophosphate (DMTP) was the major degradation route for ^{14}C -MPS in the culture media. Direct formation of DMTP from MPS has been previously reported (26, 27, 31–33). In contrast, a significant amount of ^{14}C -MPO was found in cell lysates following ^{14}C -MPS treatment, indicative of the greater conversion of MPS to MPO in cells (Figure 6B).

To ensure that the uptake studies were conducted with continued viability during the course of the exposure, the cytotoxicity was next examined. As expected, the more reactive phosphorylating agent ^{14}C -MPO was more cytotoxic than ^{14}C -MPS at concentrations > 10 μM (Figure 3). As a result of these data, we chose a concentration of MPS and MPO (1 μM) that was below the cytotoxic threshold to examine differences in uptake of these OPs in viable cells.

Although a 20-fold difference in the lipophilicity of MPS and MPO exists, this structural difference did not result in equally dramatic change in cellular uptake; in fact, our initial uptake results showed that uptake was not related to the lipophilicity. Our experiments showed that the hydrolysis product of MPO, dimethyl phosphate (DMP), actually accounted for a higher overall amount of radioactivity in MPO-treated cells as compared with MPS-treated cells, indicating relatively equal access by MPO and MPS. However, a greater overall amount of MPS was found inside cells compared with MPO. The greater amount of MPS than MPO found in cells could be the result of the greater stability of MPS once inside the cell and/or rapid breakdown of MPO in cells to DMP, thereby reducing the net amount of MPO found. The higher amount of DMP found in cells could be due to the intracellular hydrolysis and metabolism of MPO and/or extracellular (media-mediated) metabolism followed by passive diffusion across the cell membrane (34). Although a maximum cellular uptake of OPs was not more than 5% of total OP exposure

(1 μM), it is noteworthy that uptake was concentration dependent as observed from the results of lower concentration (100 and 10 nM) uptake experiments.

The cytotoxicity induced in the ^{14}C -MPS-treated cells followed a delayed trend similar to that in the ^{14}C -MPO-treated cells after 24 h of exposure likely due to a rate-dependent oxidative conversion. The highest amount of cytotoxicity (90%) exhibited by MPS was at 100 μM concentration after 72 h of exposure time. Although very little ^{14}C -MPO (7–10%, Table 2) was found in the media, a significant amount of ^{14}C -MPO (33–37%, Table 1) was found in the cytosol in MPS-treated cells. Therefore, it is likely that conversion from MPS to MPO contributed to the cytotoxicity (16).

In conclusion, differences in the stability, uptake, and cytotoxicity of ^{14}C -MPS and ^{14}C -MPO in SH-SY5Y human neuroblastoma cells were found. The use of dual-labeled MPS and MPO allowed for precise intracellular measurements of parent and hydrolyzed product and for determining rate comparisons. Data obtained in this work suggest that the thionate MPS and the oxon MPO access cells at different rates and to different extents that could yield distinct differences in protein responses.

ABBREVIATIONS USED

MPO, methyl paraoxon; MPS, methyl parathion; DMP, dimethyl phosphate; DMTP, dimethyl thiophosphate.

SAFETY

Organophosphates are toxic reagents and should be handled in a well-ventilated hood. Methyl parathion and methyl paraoxon may be rendered safe by stirring with 1 N NaOH overnight at room temperature.

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Supporting Information Available: Radioactivity counts of OP and their degradation products in cell cytosol. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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