



HAL
open science

2,4-Di-tert-butylphenol Induces Adipogenesis in Human Mesenchymal Stem Cells by Activating Retinoid X Receptors

Xiao-Min Ren, Richard C Chang, Yikai Huang, Angélica Amorim Amato, Coralie Carivenc, Marina Grimaldi, Yun Kuo, Patrick Balaguer, William Bourguet, Bruce Blumberg

► **To cite this version:**

Xiao-Min Ren, Richard C Chang, Yikai Huang, Angélica Amorim Amato, Coralie Carivenc, et al.. 2,4-Di-tert-butylphenol Induces Adipogenesis in Human Mesenchymal Stem Cells by Activating Retinoid X Receptors. *Endocrinology*, 2023, 164 (4), pp.bqad021. 10.1210/endocr/bqad021 . hal-04600470

HAL Id: hal-04600470

<https://hal.science/hal-04600470>

Submitted on 4 Jun 2024

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



Distributed under a Creative Commons Attribution - NonCommercial - NoDerivatives 4.0 International License

2,4-Di-tert-butylphenol Induces Adipogenesis in Human Mesenchymal Stem Cells by Activating Retinoid X Receptors

Xiao-Min Ren,^{1,2} Richard C. Chang,¹ Yikai Huang,¹ Angélica Amorim Amato,¹ Coralie Carivenc,³ Marina Grimaldi,⁴ Yun Kuo,¹ Patrick Balaguer,⁴ William Bourguet,³ and Bruce Blumberg^{1,5,6}

¹Department of Developmental and Cell Biology, University of California, Irvine, CA 92697-2300, USA

²Faculty of Environmental Science and Engineering, Kunming University of Science and Technology, Kunming 650500, China

³Centre de Biologie Structurale, Université de Montpellier, CNRS, Inserm, Montpellier, France

⁴Institut de Recherche en Cancérologie de Montpellier (IRCM), Inserm U1194, Université Montpellier, Institut régional du Cancer de Montpellier (ICM), Montpellier, France

⁵Department of Pharmaceutical Sciences, University of California, Irvine, CA 92697-2300, USA

⁶Department of Biomedical Engineering, University of California, Irvine, CA 92697-2300, USA

Correspondence: Bruce Blumberg, PhD, University of California Irvine, 2011 BioSci 3, Irvine, CA 92697-2300, USA. Email: blumberg@uci.edu.

Abstract

2,4-Di-tert-butylphenol (2,4-DTBP) is an important commercial antioxidant and a toxic natural secondary metabolite that has been detected in humans. However, there is scant information regarding its toxicological effects. We asked whether 2,4-DTBP is a potential obesogen. Using a human mesenchymal stem cell adipogenesis assay, we found that exposure to 2,4-DTBP led to increased lipid accumulation and expression of adipogenic marker genes. Antagonist assays revealed that 2,4-DTBP increased lipid accumulation by activating the peroxisome proliferator-activated receptor (PPAR) γ -retinoid X receptor (RXR) heterodimer. 2,4-DTBP likely activated the PPAR γ /RXR α heterodimer by activating RXR α but not directly binding to PPAR γ . We confirmed that 2,4-DTBP directly bound to RXR α by solving the crystal structure of this complex, then predicted and demonstrated that related compounds could also activate RXR α . Our study demonstrated that 2,4-DTBP and related chemicals could act as obesogens and endocrine disruptors via RXRs. These data showed that 2,4-DTBP belongs to a family of compounds whose endocrine-disrupting and obesogenic effects can be strongly modulated by their chemical composition. Structure–activity studies such as the present one could help guide the rational development of safer antioxidants that do not interact with important nuclear receptors having broad effects on human development and physiology.

Key Words: endocrine disruptor, RXR, PPAR, MSC, obesogen, adipogenesis

Abbreviations: DBD, DNA-binding domain; DMSO, dimethylsulfoxide; DTBB, di-tert-butylbenzene; DTBP, di-tert-butyl phenol; EDC, endocrine disrupting chemical; FABP4, fatty acid binding protein 4; FBS, fetal bovine serum; FSP27, fat-specific protein of 27 kDa; LBD, ligand-binding domain; LBP, ligand-binding pocket; LOEC, lowest effective concentration; LPL, lipoprotein lipase; LXR, liver “X” receptor; MSC, multipotent mesenchymal stromal stem cell; PPAR, peroxisome proliferator activated receptor; PPAR γ 2, peroxisome proliferator activated receptor gamma 2; RAR, retinoic acid receptor; ROSI, rosiglitazone; RXR, retinoid “X” receptor; SPA, synthetic phenolic antioxidant; TIF2, transcriptional intermediary factor 2; TR, thyroid hormone receptor; TTBB, tri-tert-butylbenzene; TTBP, tri-tert-butyl-phenol.

Obesity has become a serious health problem worldwide in infants, children, adolescents, and adults (1). Approximately 13% of the world population is clinically obese (body mass index ≥ 30 kg/m²) and 39% of adults are overweight (body mass index ≥ 25 kg/m²) (2). Obesity increases the risk of many chronic diseases such as type 2 diabetes, heart disease, hypertension, stroke, and certain cancers (3) and elevates risk of serious illness from COVID-19. According to the World Health Organization, at least 2.8 million deaths annually are related to overweight or obesity worldwide (4). Obesity is an even greater problem in the United States where 42% of adults are currently obese (5).

Obesity is considered to be a result of the combined effects of both genetic and environmental factors (6). Traditional risk

factors include lack of physical activity, excessive consumption of food and genetic susceptibility (7). However, these factors cannot fully explain the dramatic increase in the obesity rate worldwide. An increasing number of epidemiological studies together with experimental evidence in animal models indicated that some environmental chemicals could be important contributors to obesity (6). Such “obesogens” include natural chemicals, pharmaceutical chemicals, and xenobiotic chemicals that can promote obesity (8). Known obesogens include organotins, bisphenols, many agrochemicals, perfluoroalkyl substances, phthalates, and polybrominated diphenyl ethers (reviewed in 9). Obesogens can contribute to obesity by increasing white adipocyte cell number or increasing lipid storage in existing adipocytes (10). These chemicals can also

lead to obesity by affecting appetite and satiety, changing basal metabolic rate, altering energy balance, or influencing gut microbiota (6). Identifying which environmental pollutants are obesogens and revealing the underlying mechanisms will contribute to ameliorating the obesity pandemic and related diseases.

2,4-Di-tert-butyl phenol (2,4-DTBP) is an important commercial antioxidant that is mainly used as an intermediate to produce other high molecular weight antioxidants and UV stabilizers for hydrocarbon-based products and plastics (11). 2,4-DTBP has been detected in indoor dust (12), river sediment (13), wastewater, and sludge (14). 2,4-DTBP is also a natural toxic secondary metabolite produced by bacteria, fungi, plants, and animals (15). Human exposure can result from ingestion of dust (12), packaged food intake because tris(2,4-di-tert-butylphenol) phosphite in food packaging materials can break down to 2,4-DTBP (16), and intake from foods containing 2,4-DTBP (17). A recent study analyzing synthetic phenolic antioxidants (SPAs) in humans detected 2,4-DTBP in urine samples from the United States (18). 2,4-DTBP was detected in 100% of investigated human urine samples and was the predominant congener, contributing 88.2% to total target concentrations of SPAs in the urine samples. The concentration of total 2,4-DTBP (7.3-130 ng/mL [0.03 μ M-0.63 μ M]; geometric mean, 25.8 ng/mL; 0.12 μ M) was much higher than 2,6-di-tert-butyl-4-methylphenol (0.85 ng/mL, 3.8 nM) and 3-tert-butyl-4-hydroxyanisole (0.39 ng/mL, 2 nM), which were 2 other SPAs of concern. The relatively high levels of 2,4-DTBP found in human urine led us to investigate its possible adverse impacts.

In the present study, we assessed the capability of 2,4-DTBP to induce adipogenesis in human multipotent mesenchymal stromal stem cells (a.k.a. mesenchymal stem cells or MSCs) and explored its mechanism of action. We found that 2,4-DTBP, but not its structural analog 2,6-di-tert-butylphenol (2,6-DTBP), which is also used industrially as a UV stabilizer and antioxidant for hydrocarbon-based products (11), induced adipogenesis and expression of white adipocyte marker genes in MSCs. We used receptor selective agonist and antagonist studies to link 2,4-DTBP exposure with activation of the peroxisome proliferator activated receptor gamma (PPAR γ) and its heterodimeric partner, the retinoid X receptor (RXR) to promote adipogenesis in MSCs. Reporter gene assays revealed that 2,4-DTBP, but not 2,6-DTBP, activated RXR α , the RXR α subunit of the PPAR γ /RXR α heterodimer and RXR α in the context of 2 other nuclear receptor complexes: the liver X receptor alpha (LXR α)/RXR α heterodimer and the thyroid hormone receptor β (TR β)/RXR α heterodimer. We solved the crystal structure of 2,4-DTBP complexed with RXR α and identified the mechanistic basis underlying binding of 2,4-DTBP to RXR α and activation of RXR α by 2,4-DTBP. This analysis suggested that other related chemicals such as 2,4,6-tri-tert-butyl phenol and 1,3,5-tri-tert-butylbenzene would activate RXR α whereas 2,6-DTBP and 1,3-di-tert-butylbenzene would not. Reporter gene assays confirmed these predictions. Overall, our studies identified 2,4-DTBP as an RXR α activator and therefore a potential endocrine disrupting chemical (EDC) and obesogen.

Materials and Methods

Chemicals and Reagents

Dexamethasone, isobutylmethylxanthine, Nile red, Hoechst 33342, HX531, LG100268 (LG), 2,4-di-tert-butylphenol

(2,4-DTBP), 2,6-di-tert-butylphenol (2,6-DTBP), 2,4,6-tri-tert-butylphenol (2,4,6-TTBP), 1,3-di-tert-butylbenzene (1,3-DTBB), 1,3,5-tri-tert-butylbenzene (1,3,5-TTBB), LG100268, GW3065, T3, and TTNPB were purchased from Sigma-Aldrich. CD3254 was purchased from Tocris Bioscience. Rosiglitazone (ROSI) was purchased from Cayman Chemical Company. T0070907 was from Enzo Life Sciences. Dimethylsulfoxide (DMSO) was purchased from Thermo Fisher Scientific.

Human MSC Adipogenesis Assay

Human bone marrow MSCs were purchased from PromoCell at passage 2 (lot 429Z001-Female). Human MSCs were cultured in alpha Modification of Eagle's minimal essential medium (α MEM) (Thermo Fisher Scientific) supplemented with 15% fetal bovine serum (FBS) (Gemini Bio-Products), 2% penicillin, 5000 IU/mL-streptomycin 5000 μ g/mL (Corning), and 10 μ M HEPES (Fisher Chemical). The cells were split when the confluency reached 90% and used at passage 6 for adipogenesis assays. A total of 40 000 cells per well were seeded in a 24-well plate or 80 000 cells per well were seeded in a 12-well plate. When cells reached 100% confluency, media were replaced with differentiation media (α MEM, 15% FBS supplemented with adipogenic induction cocktail [500 μ M isobutylmethylxanthine, 1 μ M dexamethasone, 5 μ g/mL human recombinant insulin]) and ligands. Specific ligands (dissolved in DMSO) were administered twice weekly for 2 weeks. Antagonist assays were performed similarly, except that T0070907 was added twice daily (to 10 μ M). DMSO concentrations in the medium were identical between vehicle controls and test chemicals and never exceeded 0.1%. At the end of each assay, cells in the 24-well plates were fixed in 3.7% formaldehyde in phosphate-buffered saline for 30 minutes at 25 $^{\circ}$ C for lipid staining; cells grown in 12-well plates were homogenized and total RNA extracted for gene expression analysis.

Gene Expression Analysis

Total RNA was isolated from cells using Trizol reagent as recommended by the manufacturer (Invitrogen Life Technologies). Reverse transcription and quantitative real-time reverse transcriptase polymerase chain reaction were performed using Transcriptor reverse transcriptase and Sybr Green Master Mix (Roche Diagnostics Corp., Indianapolis, IN). The following genes were examined: *FABP4* (fatty acid binding protein 4), *FSP27* (fat-specific protein of 27 kDa), *PPAR γ 2*, and *LPL* (lipoprotein lipase). The details of the experiment and sequences of primers used for quantitative real-time reverse transcriptase polymerase chain reaction were described in our previous studies (19, 20).

Transient Transfection and Luciferase Reporter Gene Assays

The CMX-GAL4 effector plasmids using pCMX-GAL4 DNA-binding domain (DBD) fused to nuclear receptor ligand-binding domains (LBDs) for human PPAR γ (pCMX-GAL4-PPAR γ), human RXR α (pCMX-GAL4-RXR α), human LXR α (pCMX-GAL4-LXR α), and human TR β (pCMX-GAL4-TR β) were previously described (21-23). Transient transfections were performed in COS-7 cells (ATCC CRL-1,651TM) as described (20). Briefly, COS-7 cells were

seeded at 15 000 cells/well in 96-well tissue culture plates in Dulbecco's modified Eagle's medium (HyClone) supplemented with 10% calf bovine serum (Atlanta Biologicals). Twenty-four hours later, cells were transfected in Opti-MEM reduced-serum medium (Invitrogen Life Technologies, Grand Island, NY) when they reached ~90% confluency (~24 hours after seeding). Cells were transfected with 1 μ g of CMX-GAL4 effector plasmids (0.3 μ g for pCMX-GAL4-PPAR γ plasmid) with or without 1 μ g of pCMX-LBD of RXR α (L-RXR α) (0.3 μ g of pCMX-L-RXR α for pCMX-GAL4-PPAR γ plasmid) together with 5 μ g of tk-(MH100)4-luciferase reporter and 5 μ g of CMX- β -galactosidase transfection control plasmids in Opti-MEM using Lipofectamine 2000 reagent (Invitrogen Life Technologies, Grand Island, NY), following the manufacturer's recommended protocol. After 24 hours of incubation, the medium was replaced with phenol red-free Dulbecco's modified Eagle's medium (Invitrogen Life Technologies)/10% resin charcoal-stripped FBS plus ligands at concentrations indicated in the figure legends for an additional 24 hours before luciferase and β -galactosidase assays. 2,4-DTBP and 2,6-DTBP were tested from 1.2 μ M to 40 μ M with 2-fold dilutions. No noticeable cytotoxicity was produced at 40 μ M, as judged by reduction of β -galactosidase activity (data not shown). The ligands were dissolved in DMSO. The DMSO concentrations in the medium were identical between vehicle controls and test chemicals and never exceeded 0.1%. Twenty-four hours after adding the ligands to the media, cells were lysed in 165 μ L of lysis buffer and allowed to shake for 30 minutes at 25 $^{\circ}$ C. For luciferase assay, 35 μ L of the lysate from each well was transferred to a well in a clean, nontreated, white, flat-bottom, polystyrene, 96-well plate (Costar). Additionally, 100 μ L of freshly made luciferase solution was added to each well with the cell lysate, and that plate was shaken for 30 seconds at room temperature. Plates were placed in a Dynatech ML3000 luminometer, and data were acquired with luminometer ML3000 (version 3.07) software. For β -galactosidase assays, 35 μ L of the cell lysate were transferred to a clear, flat-bottom, 96-well plate (Thermo Fisher Scientific) and 100 μ L of β -galactosidase solution added to each well. Plates were shaken for 30 seconds, then incubated at 25 $^{\circ}$ C for 15 minutes. Absorbance was measured at 405 nm on a Versamax microplate reader (Molecular Devices) using SOFTmaxPRO 4.0 software. Each luciferase read was normalized with the corresponding β -galactosidase read coming from the same transfection well and multiplied by the number of minutes the β -galactosidase plate was incubated (usually 15 minutes). Data are reported as relative light units (Luc/Gal). The median effective concentration and maximum activations were calculated using GraphPad Prism 9.0 (GraphPad Software, Inc.). Transfections were performed in triplicate and reproduced in multiple experiments.

Statistical Analysis

The data are expressed as the mean value \pm standard error of the mean (SEM) ($n = 3$). Unpaired t -test was used to determine the significance of effects elicited by the chemicals relative to the control group in human MSC adipogenesis assays and to determine the significance of differences comparing +T0070907 vs -T0070907 or +HX531 vs -HX531 with each other. In the transfection assays, the significance of the experimental data was analyzed using 1-way analysis of variance, followed by a least significant difference multiple

comparisons test. A P value of $<.05$ was considered to be statistically significant. Statistical analysis used GraphPad Prism 9.0 (GraphPad Software Inc., San Diego, CA).

Retinoic Acid Receptor β Reporter Cell Line

The HELN retinoic acid receptor (RAR) β cell line was already described (24). Briefly, we firstly generated the HELN cell line by transfecting HeLa cells with the p-ERE- β Glob-Luc-SVNeo plasmid. Secondly, we transfected HELN cells with pRAR β (ER α DBD)-puro plasmid. pERE- β Glob-Luc-SVNeo contains a luciferase gene driven by an estrogen receptor binding site in front of the β -globin promoter and a neomycin phosphotransferase gene under the control of the SV40 promoter. In pRAR β (ER α DBD), the encoded chimeric RAR β protein contains the DBD of the estrogen receptor α . Puromycin N-acetyl transferase selection marker expression confers resistance to puromycin.

To measure the activity of the chemicals, HELN RAR β cells were seeded at a density of 40 000 cells per well in 96-well white opaque tissue culture plates (Dutscher, Brumath, France). Compounds were added 8 hours later alone in presence of the pan RAR-agonist TTNPB 100 nM or the pan RXR-agonist CD3254 100 nM, and cells were incubated for 16 hours. At the end of incubation, the culture medium was replaced by a medium containing 0.3 mM luciferin. Luciferase activity was measured for 2 seconds in intact living cells using a microBeta Wallac luminometer (Perkin Elmer, Villebon sur Yvette, France). Tests were performed in quadruplicate in at least 3 independent experiments and data were expressed as mean \pm SEM. Results are expressed in percentage of the maximal activity obtained in presence of TTNP 100 nM. Dose-response curves were fitted using the sigmoid dose-response function of a graphics and statistics software package (Graph-Pad Prism, version 4, 2003, Graph-Pad Software Inc., San Diego, CA, USA).

Crystallographic Analysis

The histidine-tagged LBD of human RXR α (residues 223-462 in a pET15b vector) was expressed in *Escherichia coli* BL21(DE3). Cells were grown in ZYM-5052 autoinduction medium supplemented with 100 μ g/mL ampicillin. After 24 hours of incubation at 25 $^{\circ}$ C, cell cultures were harvested by centrifugation at 8000g for 20 minutes. The cell pellet from 2 L of culture was resuspended in 250 mL of buffer A (20 mM Tris-HCl pH 8.0, 500 mM NaCl, 5 mM imidazole) supplemented with 100 μ g/mL of lysozyme and a protease inhibitor cocktail (Complete, mini, EDTA-free tablet, Roche Applied Science) then incubated with shaking for 30 minutes at 4 $^{\circ}$ C. The suspension was lysed by sonication and centrifuged at 35 000g for 30 minutes at 4 $^{\circ}$ C. The supernatant was loaded onto a 5-mL Ni²⁺ affinity column (HiTrap chelating column, GE healthcare), pre-equilibrated with buffer A, using the Äkta pure system (GE Healthcare). The column was washed with 10 volumes of buffer A. Protein was eluted in 0 to 500 mM imidazole gradient using 25 column volumes of the buffer. The fractions containing RXR α LBD were pooled and incubated overnight at 4 $^{\circ}$ C with thrombin to remove the histidine tag. The protein was further purified using a Superdex 75 26/60 gel filtration column (GE healthcare) pre-equilibrated with buffer C (20 mM Tris-HCl pH 8.0, 150 mM NaCl, 5% glycerol, 5 mM dithiothreitol).

5wFractions containing the purified receptor were pooled, mixed with 1:3 molar ratio of 2,4-DTBP and 1:2 molar ratio of the TIF2 coactivator peptide and concentrated to 8 mg/mL. Crystals were obtained by vapor diffusion at 293 °K by mixing 1 μ L of protein with 2 μ L of precipitant solution. The well buffer contained 20% polyethylene glycol 3350 and 0.2 M sodium formate. A single crystal was mounted from the mother liquor onto a cryoloop (Hampton Research, Aliso Viejo, CA, USA), soaked in the reservoir solution supplemented with 20% glycerol and 40 nM 2,4-DTBP and then frozen in liquid nitrogen. Diffraction data were collected on the ID30A-1 at the European Synchrotron Radiation Facility (Grenoble, France). Data were processed and scaled with XDS (25) and Scala (26). Crystals belong to space group P212121. The structure was refined using REFMAC5 (27) and COOT (28). Figures were generated with PyMol (<https://pymol.org/2/>). Structural data were deposited in the Worldwide Protein Data Bank with accession number 7NKE (29). Data collection and refinement statistics can be found elsewhere (Table S1 (30)).

Results

2,4-DTBP Increased Adipogenesis in Human MSCs

MSCs have been widely used to study the obesogenic properties of environmental chemicals (19, 20, 31, 32). We tested the potential obesogenic properties of 2,4-DTBP and 2,6-DTBP using a human MSC adipogenesis assay. The selective PPAR γ agonist, ROSI was used as a positive control, DMSO vehicle (0.1% v/v) was the negative control. Treatment of MSCs with adipogenesis cocktail supplemented with 500 nM ROSI for 14 days elicited a significant and substantial increase in lipid accumulation, as measured by Nile Red fluorescence normalized to cellular DNA content measured by Hoechst 33342 fluorescence. After treatment with 10 μ M 2,4-DTBP for 14 days, adipogenesis was increased significantly whereas lower doses of 2,4-DTBP or all doses of 2,6-DTBP did not (Fig. 1A). Induction of adipogenesis by LG100268 was similar to that of 2,4-DTBP (Fig. 1A).

We confirmed the adipogenic effects by evaluating the expression of white adipocyte marker genes in response to chemical exposure. Levels of mRNAs encoding FABP4, FSP27, LPL, and PPAR γ -2 were significantly increased in the 500 nM ROSI-treated cells compared with vehicle (DMSO) (Fig. 1B and 1E). Levels of FSP27, LPL, FABP4, and PPAR γ 2 mRNAs were increased in cells treated with 10 μ M 2,4-DTBP compared with 2,6-DTBP and vehicle control, both of which were inactive.

2,4-DTBP-Induced Adipogenesis Was Blocked by the PPAR γ Antagonist T0070907 or the RXR α Antagonist HX531

PPAR γ plays a key role in regulating the transcription of genes involved in adipogenesis in MSCs (33). Therefore, we explored whether antagonizing PPAR γ activation with the specific antagonist, T0070907 affected 2,4-DTBP-mediated increases in lipid accumulation in human MSCs. We found that cotreatment with 10 μ M T0070907 (34) significantly ($P < .05$) inhibited lipid accumulation induced by treatment with 100 nM ROSI or by 10 μ M 2,4-DTBP (Fig. 2A).

PPAR γ regulates the transcription of adipogenic genes as a heterodimer with RXRs (33). To assess the extent of RXR involvement in 2,4-DTBP-induced lipid accumulation, we

conducted an RXR antagonist assay by comparing the effects of 2,4-DTBP in the presence or absence of the specific antagonist HX531 (35). Cotreatment with 1 μ M HX531 significantly ($P < .05$) inhibited lipid accumulation induced by 100 nM of the specific RXR agonist, LG100268 (22), and by 10 μ M 2,4-DTBP (Fig. 2B). Taken together, these results implicate the PPAR γ -RXR heterodimer in 2,4-DTBP-induced adipogenesis, but do not identify which partner is responsible.

2,4-DTBP Showed Agonistic Activity Toward GAL4-PPAR γ and GAL4-RXR α

We next assessed whether 2,4-DTBP could activate human PPAR γ or RXR α using transient transfection in COS-7 cells with GAL4-receptor chimeras. ROSI enhanced the activity of a GAL4-PPAR γ in a dose-dependent manner with a lowest effective concentration (LOEC) value of \sim 10 nM (Fig. 3A). Similarly, 2,4-DTBP enhanced GAL4-PPAR γ activation in a dose-dependent manner with a LOEC value of \sim 10 μ M (Fig. 3B). 2,6-DTBP did not activate GAL4-PPAR γ between 1.2 and 40 μ M (Fig. 3C).

Interestingly, 2,4-DTBP also showed agonistic activity toward human RXR α . Treatment with LG100268 strongly induced activity by GAL4-RXR α in transient transfection assays also with a LOEC of \sim 10 nM (Fig. 3D). 2,4-DTBP also enhanced GAL4-RXR α activity with a LOEC value of \sim 10 μ M (Fig. 3E). 2,6-DTBP was inactive (Fig. 3F). Taken together, these results confirm that 2,4-DTBP appeared to have agonistic activity toward both PPAR γ and RXR α .

RXR α Mediated the Agonistic Activity of 2,4-DTBP Toward PPAR γ

We next asked whether 2,4-DTBP could interfere with the ability of ROSI to activate GAL4-PPAR γ as would be expected by a bona fide PPAR γ ligand. We tested the ability of different concentrations of 2,4-DTBP to affect activation of GAL4-PPAR γ by 500 nM ROSI and found no significant increase or decrease in activity (Fig. 4A), suggesting that 2,4-DTBP and ROSI did not compete for the same binding site in PPAR γ . Since 2,4-DTBP did not further activate GAL4-PPAR γ , we inferred that it does not directly interact with PPAR γ .

Since PPAR γ /RXR is a permissive heterodimer, activation of RXR in the heterodimer could lead to activation of the complex or further enhance the activity of the agonist-bound PPAR γ (23). Because COS-7 cells express endogenous RXR α (36), it is possible that there is some interaction between endogenous full-length RXR α and the GAL4-PPAR γ heterodimer in the transient transfection assays. Since 2,4-DTBP had agonistic activity toward RXR α (Fig. 3E), we hypothesized that 2,4-DTBP might activate GAL4-PPAR γ via a heterodimer with endogenous RXRs. In support of this hypothesis, we found that LG100268 could also elicit modest (\sim 2-fold) activation of GAL4-PPAR γ (Fig. 4D).

To further explore the involvement of RXR α in activation of the PPAR γ /RXR α heterodimer by 2,4-DTBP, we cotransfected COS-7 cells with 2 plasmids: 1 expressing GAL4-PPAR γ and the other expressing only the LBD of RXR α (L-RXR α). L-RXR α lacks the ability to bind DNA and must exert any observed actions via its heterodimeric partner (23). Cotransfection with L-RXR α (Fig. 4E) strongly potentiated the ability of LG100268 to activate GAL4-PPAR γ (compare Fig. 4E with Fig. 4D). Similarly, 2,4-DTBP was now able

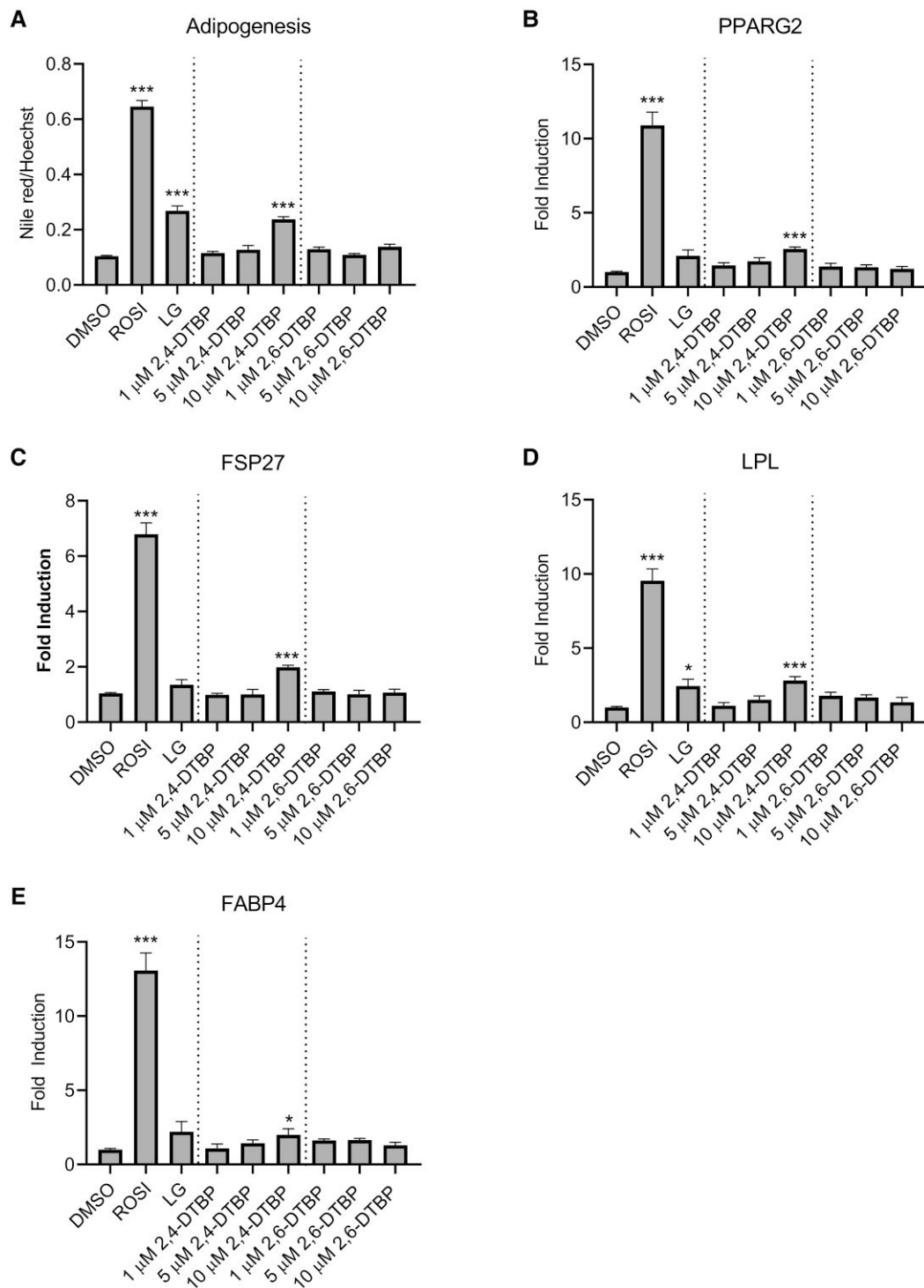


Figure 1. Effect of ROSI, LG100268, 2,4-DTBP, and 2,6-DTBP on lipid accumulation and adipogenic marker gene expression in human MSCs. Cells were treated with 500 nM ROSI, 100 nM LG100268, or different concentrations of 2,4-DTBP or 2,6-DTBP as indicated. Three replicate wells were included for each group in a 24-well plate for lipid accumulation assays or in 12-well plates for adipogenic marker gene expression assay. (A) effects of chemical exposure on adipogenesis in hMSCs. Data are expressed as lipid accumulation normalized to cellular DNA content. (B-E) effects of chemical exposure on expression of adipogenic markers in hMSCs. Data are expressed as fold induction over vehicle control. Error bars represent the standard error of the mean for 3 replicates. * $P \leq .05$, ** $P \leq .01$, *** $P \leq .001$ compared with cells treated with 0.1% DMSO vehicle control.

to induce robust activation of the GAL4-PPAR γ -L-RXR α heterodimer at 1 μ M (Fig. 4F). Both LG100268 (Fig 4B) and 2,4-DTBP (Fig 4C) potentiated reporter gene activity induced by 25 nM ROSI in cells cotransfected with GAL4-PPAR γ and L-RXR α , although LG100268 potentiation was only

significant at 10 nM and higher. We inferred that 2,4-DTBP action on the PPAR γ -RXR heterodimer is mediated through RXR since expression of exogenous RXR α LBD potentiated action of 2,4-DTBP and the RXR-selective agonist LG100268 on GAL4-PPAR γ .

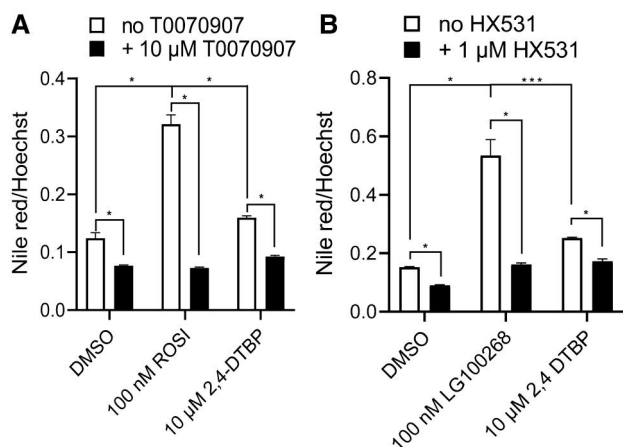


Figure 2. Study of the inhibitory effects of PPAR γ antagonist T0070907 and RXR α antagonist HX531 on lipid accumulation induced by 2,4-DTBP in human MSCs. (A) Cells were treated with 100 nM ROSI or 10 μ M 2,4-DTBP in the absence or presence of 10 μ M T0070907. (B) Cells were treated with 100 nM LG100268 or 10 μ M 2,4-DTBP in the absence or presence of 1 μ M HX531. Data are expressed as lipid accumulation normalized to cellular DNA content. Error bars represent the standard error of the mean for 3 replicates. * $P \leq .05$, ** $P \leq .01$, *** $P \leq .001$ compared with cell samples treated with 0.1% DMSO vehicle. 10 μ M T0070907 or 1 μ M HX531.

2,4-DTBP Activated GAL4-LXR α and GAL4-TR β Through RXR α in the Heterodimers

In addition to PPAR γ , RXR α also heterodimerizes with many other nuclear receptors, including LXR, TR, RAR, constitutive androstane receptor, farnesoid X receptors, pregnane X receptor, and vitamin D receptor (37, 38). Since we found that 2,4-DTBP activation of RXR α in the heterodimers could affect the function of PPAR γ , we next asked whether 2,4-DTBP could also modulate the function of other nuclear receptors by activating RXR α in the heterodimers.

To test this hypothesis, we evaluated the effects of 2,4-DTBP on human GAL4-LXR α and GAL4-TR β . The results obtained for GAL4-LXR α were similar to those shown for GAL4-PPAR γ in Fig. 4. In COS-7 cells transfected with GAL4-LXR α alone, LG100268 (Fig. S1A) and 2,4-DTBP (Fig. S1B) increased reporter gene activity significantly (30). This increase was strongly potentiated by cotransfection with L-RXR α (Fig. S1E and S1F (30)). Neither LG100268 (Fig. S1C) nor 2,4-DTBP (Fig. S1D) (30) showed any interference with activation of GAL4-LXR α by its specific agonist GW3965 (39). When cells were additionally cotransfected with L-RXR α , activation by GW3965 was potentiated by LG100268 (Fig. S1G) and 2,4-DTBP (Fig. S1H) (30). We inferred that LG100268 and 2,4-DTBP activated GAL4-LXR α heterodimer via its RXR α component, rather than LXR α directly.

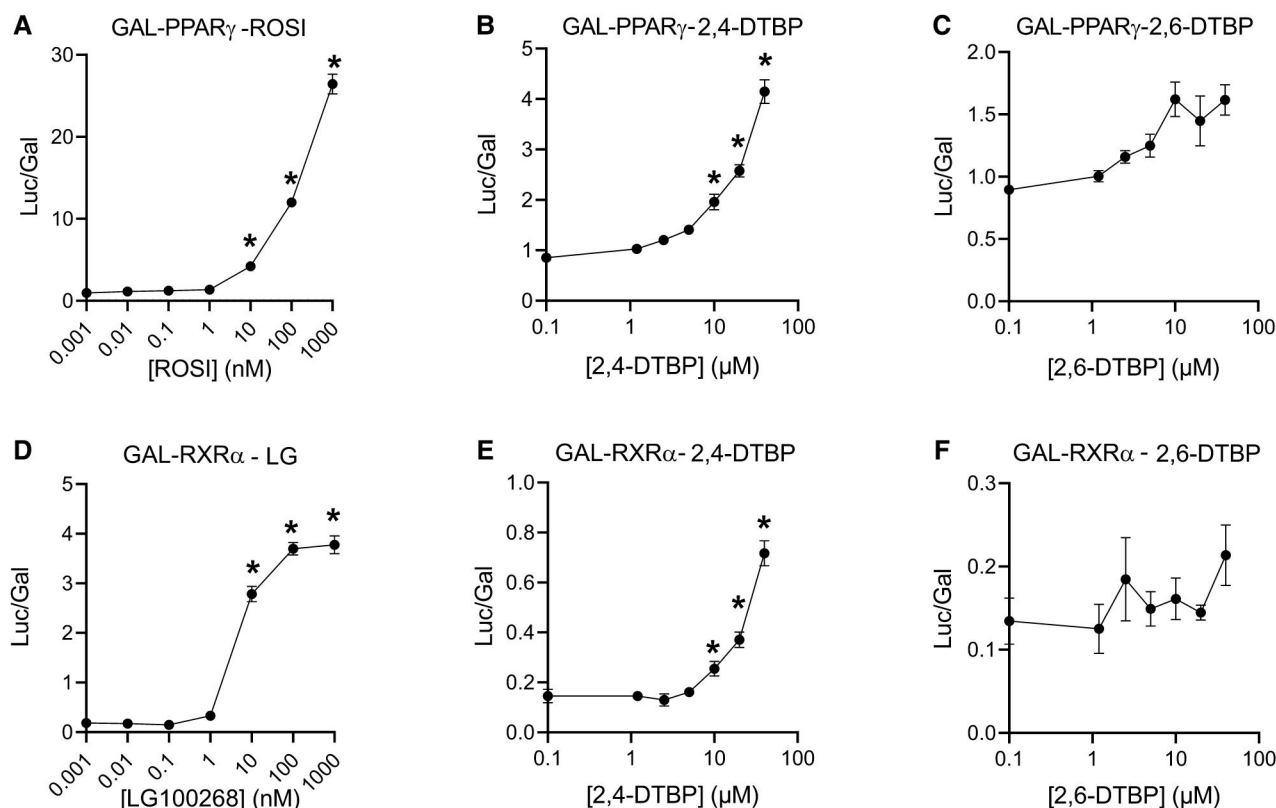


Figure 3. The effect of ROSI, LG100268, 2,4-DTBP, and 2,6-DTBP on GAL4-PPAR γ - and GAL4-RXR α -regulated luciferase activity in COS-7 cells. (A-C) GAL4-PPAR γ or (D-F) GAL4-RXR α were transfected into COS-7 cells. Plasmid transfected cells were treated with different concentrations of chemicals for 24 hours. Three replicated wells were included for each group in a 96-well plate. Data are reported as relative light units (Luc/Gal). Each luciferase read was normalized with the corresponding β -galactosidase read coming from the same transfection well and multiplied by the number of minutes the β -galactosidase plate was incubated (15 minutes). Error bars represent the standard error of the mean for 3 replicates. * $P \leq .05$, compared with cell samples treated with 0.1% DMSO vehicle.

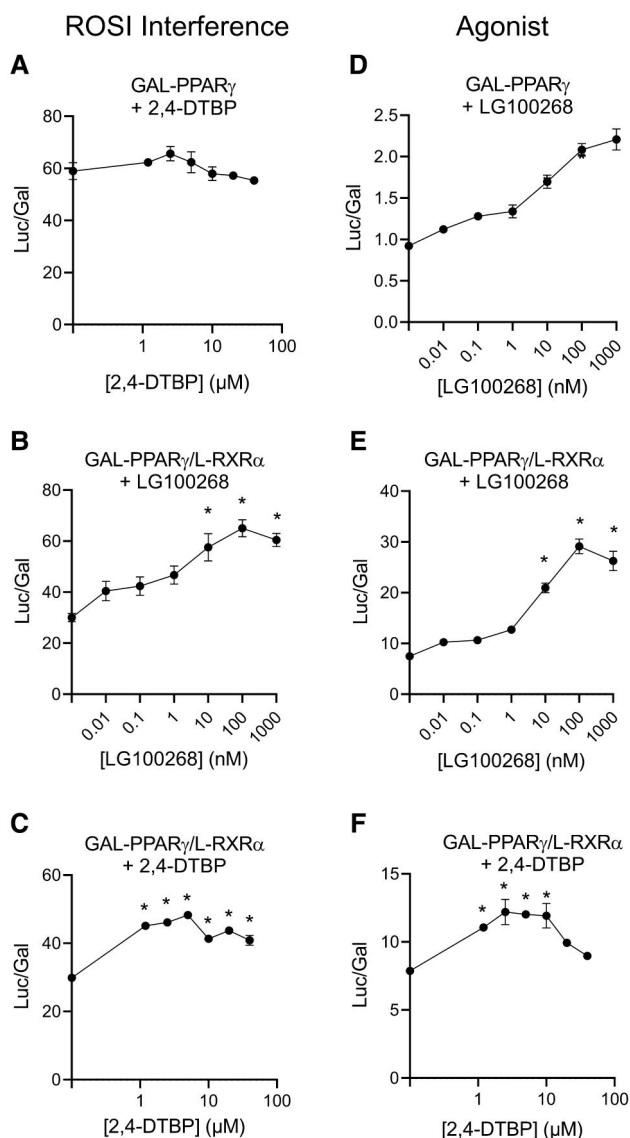


Figure 4. The effect of LG100268 and 2,4-DTBP on GAL4-PPAR γ - or GAL4-PPAR γ /L-RXR α -regulated luciferase activity in COS-7 cells. (A-C) ROSI interference assays evaluating the ability chemicals to interfere with ROSI-induced reporter activity on GAL4-PPAR γ (A), or GAL4-PPAR γ /L-RXR α -induced reporter activity (B, C). (D-F) Agonist assays evaluating the effects of chemicals on ability of GAL4-PPAR γ (D), or GAL4-PPAR γ /L-RXR α (E, F) to elicit luciferase reporter activity. The concentration of ROSI were 500 and 25 nM in the GAL4-PPAR γ - and GAL4-PPAR γ /L-RXR α -regulated luciferase reporter gene assay, respectively. Transfected cells were treated with indicated concentrations of chemicals for 24 hours in triplicate in 96-well plates. Data are reported as relative light units (Luc/Gal). Error bars represent the standard error of the mean for 3 replicates. * $P < .05$, compared with cell samples treated with 0.1% DMSO vehicle. (D-F) or with 500 nM ROSI (A) or 25 nM ROSI (B, C).

We next tested the effects of these chemicals on TR β and observed that LG100268 (Fig. S2A, C) and 2,4-DTBP (Fig. S2B, D) neither activated nor interfered with activation of GAL4-TR β by T3 (30). However, cotransfection with L-RXR α allowed LG100268 (Fig. S2E, G) and 2,4-DTBP (Fig. S2F, H) to activate GAL4-TR β in the absence (Fig. S2E, F) or presence (Fig. S2G, H) of its specific ligand, T3 (30). These results further supported a model in which

2,4-DTBP alters the function of nuclear receptor heterodimers via activation of the RXR α partner.

Crystallographic Analysis of the Binding of 2,4-DTBP to RXR α

Prototypical synthetic or natural retinoids (the subclass of retinoids targeting RXR α such as 9-cis-retinoic acid) contain a carboxylic head group involved in a salt bridge with an arginine residue and a long aliphatic/aromatic chain making numerous van der Waals contacts with essentially non-polar residues distributed all over the ligand-binding pocket (LBP) of RXR α . 2,4-DTBP neither structurally nor chemically resembles classical retinoids. To gain structure-based insight into its mode of action, we solved the crystal structure of the RXR α LBD (hereafter RXR) bound to 2,4-DTBP and a short LxxLL-containing peptide derived from the transcriptional intermediary factor 2 (TIF2), at 2.35 Å resolution (Table S1(30)). The structure reveals a RXR homodimer with 2,4-DTBP and the coactivator peptide bound to both protomers (Fig. 5A). Each subunit displayed the canonical active conformation with the C-terminal helix H12 (also termed AF-2 or activation helix) capping the LBP, and the 2,4-DTBP could be precisely placed in its electron density. Interestingly, the density map allowed the unambiguous identification of 2 alternative orientations of 2,4-DTBP that can be reached through rotation by 180° around the hydroxyl axis (Fig. 5A, inset).

2,4-DTBP bound between helices H3 and H11, and in close proximity to C432. While not directly in contact with the AF-2 helix, the tert-butyl group at position 2 (and the alternative 2' site) plays an important role in stabilizing key residues of helices H3, H5 and H11 to which helix H12 docks in the transcriptionally active conformation (green residues in Fig. 5B). On the other hand, the tert-butyl at position 4 makes extensive van der Waals contacts with hydrophobic residues deeply buried in the LBP (Fig. 5B, residues are blue). These interactions are critical for the binding affinity of 2,4-DTBP as well as for its activity, by maintaining the tert-butyl group at position 2 in close proximity to residues involved in receptor activation. The hydroxyl moiety of 2,4-DTBP appears to be involved in a hydrogen bond with the main chain carbonyl group of C432.

Comparison of the present structure with that of RXR α bound to the reference synthetic agonist CD3254 (40) shows that 2,4-DTBP occupies only a small fraction of the LBP, but that interactions with residues crucial for stabilization of the receptor-active form are preserved in both complexes (Fig. 5C). Interestingly, similar observations were previously made with the potent RXR environmental agonist tributyltin (41). Indeed, superposition of the 2,4-DTBP- and TBT-bound RXR α structures revealed that the binding sites of the 2 compounds largely overlap (Fig. 5D). Overall, it appears that, as for TBT, although 2,4-DTBP interacts with only a subset of binding pocket residues in the H11 region, it is engaged in enough essential contacts to stabilize RXR α in its active conformation. However, the high-affinity binding of TBT is ensured by a covalent bond linking the metal atom of the organotin and cysteine residue 432 in helix H11 (masked by the ligands in Fig. 5D). This covalent bond does not exist for 2,4-DTBP, which, as a consequence, binds much less avidly than TBT to RXR α .

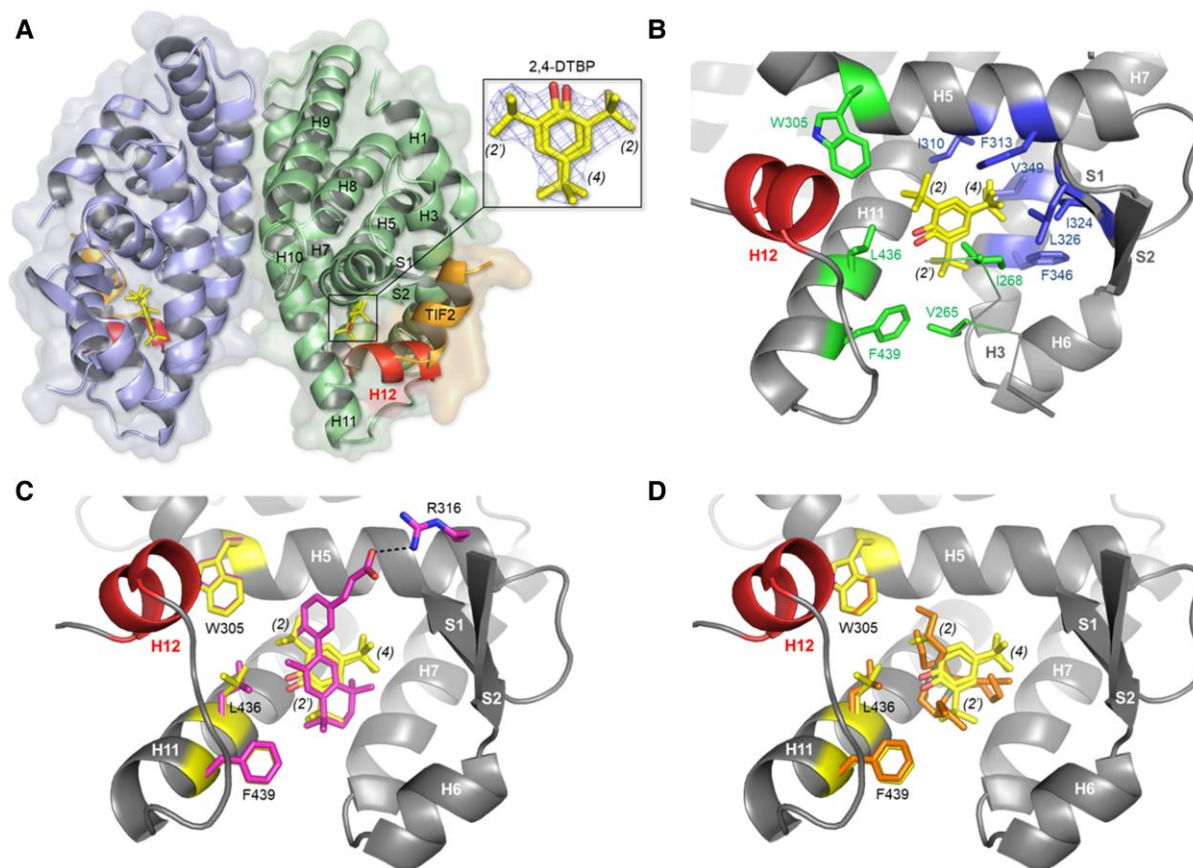


Figure 5. Structural basis for RXR interaction with and activation by 2,4-DTBP. (A) Overall structure of RXR LBD homodimer in complex with 2,4-DTBP (yellow sticks) and TIF2-NR2 peptide (orange). The activation helix H12 is highlighted in red. The inset shows the 2Fo-Fc electron density map of 2,4-DTBP. (B) Close-up view of the ligand-binding pocket of RXR bound to 2,4-DTBP. Key RXR residues in contact with the tert-butyl moieties at positions 2, 2' and 4 are depicted as green and blue sticks, respectively. (C) Superposition of the 2,4-DTBP- and CD3254-bound RXR structures (PDB code 3E94). As a prototypical rexinoid, CD3254 (magenta sticks) contains an acidic head group and a long aliphatic/aromatic chain bridging arginine 316 (H5) on 1 side and H11 on the other. (D) Superposition of the 2,4-DTBP- and TBT-bound RXR structures (PDB code 3FUG) showing that the binding sites of the 2 compounds overlap. TBT is displayed as orange sticks.

Structure–Activity Analysis of 2,4-DTBP and Congeners on RXR α Based on Stably Transfected HELN RAR β Reporter Cells

The crystallographic data revealed the structural basis for the agonistic activity of 2,4-DTBP toward RXR α and could allow a better understanding of the binding and activation properties of its congeners. We next investigated the activity of 2,4-DTBP and closely related compounds toward RXR α /RAR β heterodimer by using a stably transfected HELN RAR β reporter cell line in which both RXR and RAR agonist are able to induce luciferase expression. We monitored the agonist potential of 2,4-DTBP and 4 related compounds (Fig. S3A (30)), namely the 2,4,6-tri-tert-butylphenol bearing an additional tert-butyl group at position 6 (2,4,6-TTBP), the 2,6-DTBP containing 2 tert-butyl groups in ortho positions, and the 2 corresponding benzene derivatives lacking the hydroxyl moiety, the 1,3-DTBB and 1,3,5-tri-tert-butylbenzene (1,3,5-TTBB). We found that 2,4,6-TTBP is, by far, the most potent RXR α agonist, followed by 1,3,5-TTBB and 2,4-DTBP, while 1,3-DTBB and 2,6-DTBP are inactive (Fig. S3A (30)). To assess the specific effects of 2,4-DTBP, 2,4,6-TTBP, and 1,3,5-TTBB, HELN RAR β cells were coincubated with saturating concentrations of the RAR and RXR agonists CD3254 or TTNPB. 2,4-DTBP, 2,4,6-TTBP, and 1,3,5-TTBB further activated the TTNPB-saturated

RXR-RAR β heterodimer (Fig. S3B (30)). However, 2,4-DTBP, 2,4,6-TTBP, and 1,3,5-TTBB appear unable to act in conjunction with CD3254 to enhance the activity of RXR-RAR β (Fig. S3C (30)). These transactivation experiments confirmed the ability of 2,4-DTBP, 2,4,6-TTBP, and 1,3,5-TTBB to activate the RXR α -RAR β heterodimer through RXR α . These activity profiles are in full agreement with our structural analysis and confirm that the tert-butyl groups at positions 2, 4 and 6 (2' site in the 2,4-DTBP complex structure) are important for compound affinity and activity. With 3 tert-butyl groups, 2,4,6-TTBP displays the best potency and efficacy values followed by 1,3,5-TTBB differing from the latter by the lack of the hydroxyl moiety. The specific role of the tert-butyl at position 4 suggested by the crystal structure was also confirmed by cellular assays since the 2,6-DTBP containing only 2 tert-butyl groups on both sides of the hydroxyl moiety shows no detectable activity whereas with 2 tert-butyl groups in ortho and para positions, 2,4-DTBP acts an effective RXR α activator.

Discussion

2,4-DTBP has been detected in urine samples from the general US population at high concentrations (18); however, information about its toxicity is very limited. Since 2,4-DTBP is widely

used as an antioxidant in plastics, is known to interact with nuclear hormone receptors (42), and has been detected in humans, we asked whether 2,4-DTBP is a potential obesogen and, if so, what is its underlying mechanism of action.

We used an established MSC assay (19, 20, 22, 34, 43) to evaluate the potential of 2,4- and 2,6-DTBP for obesogenic activity. We found that 2,4-DTBP exposure induced lipid accumulation in human MSCs with a corresponding increase in the induction of marker genes for white adipocyte differentiation (Fig. 1). In contrast, 2,6-DTBP was inactive in these assays, indicating that 2,4-DTBP is a potential obesogen. We further found that 2,4-DTBP activated the PPAR γ -RXR α heterodimer—the so-called “master regulator” of adipogenesis (33). Activation of PPAR γ is a well-established predictor of adipogenic activity *in vivo*, and many chemicals have been shown to promote adipogenesis by activating PPAR γ (reviewed in (31, 32)). We used specific PPAR γ and RXR α antagonist assays to confirm the requirement for this heterodimer in promoting adipogenesis (Fig. 2). Transient transfection assays suggested that 2,4-DTBP specifically modulated activation of this heterodimer (Fig. 3).

It is well known that the PPAR γ /RXR α heterodimer is permissive because it can be activated by ligands for either partner (38). Using a series of GAL4-receptor LBD chimeric receptors and cotransfection with the RXR α LBD, we showed that 2,4-DTBP specifically activated the RXR α subunit of the PPAR γ -RXR, LXR α -RXR α , and TR β -RXR heterodimers (Figs. 3 and 4; Figs S1, S2 (30)). Previously, studies showed that activating the RXR α in the PPAR γ /RXR α heterodimer played an important role in mediating the obesogenic effect of some environmental chemicals. For example, we found that the agrochemical fludioxonil induced adipogenesis in 3T3-L1 preadipocytes and in MSCs via the PPAR γ /RXR α heterodimer by activating RXR α but not PPAR γ (20). Moreover, we demonstrated that signaling through the RXR α subunit of the PPAR γ /RXR α heterodimer was critical to commit MSCs to the adipocyte lineage (22). Combining the results of human MSC adipogenesis assay and receptor activation assays, we inferred that 2,4-DTBP is a potential obesogen that acts via RXR α in the context of the PPAR γ /RXR α heterodimer.

Unlike the PPAR γ /RXR α heterodimer, TR β /RXR α was thought to be a nonpermissive heterodimer (23). However, RXR α can act as a “nonsilent” partner of TR in particular cellular environments (high ratio of coactivator to corepressor), allowing activation of TR signaling by RXR agonists (44, 45). The well-studied environmental obesogen TBT can act as an RXR α agonist to potentiate TR-regulated gene expression and resultant morphological programs directed by thyroid hormone signaling *in vitro* and *in vivo* (46). Our studies showed that 2,4-DTBP activated the TR β /RXR α heterodimer by activating RXR α . Overall, our results demonstrated that the 2,4-DTBP activated the RXR α component of PPAR γ /RXR α , LXR α /RXR α , and TR β /RXR heterodimers.

We next solved the crystal structure of 2,4-DTBP bound to RXR α and identified its specific mode of binding (Fig. 5). The structure explained the RXR binding and activation properties of 2,4-DTBP and allowed us to predict the activity of various congeners. Notably, we predicted that 2,4,6-TTBP would be a more potent activator of RXR, and receptor activation assays confirmed this prediction (Fig. S3 (30)). In a recent study, Kodama et al explored the structure–activity relationship of 1,3-bis-tert-butyl monocyclic benzene derivatives for RXR activation by *in vitro* and *in silico* analyses. Their results also

indicated that addition of a bulky substituent at the 6-position of 2,4-DTBP is favorable for agonistic activity toward human RXR (47).

Because 2,4-DTBP received little attention previously, limited information was available about its toxicity. 2,4-DTBP elicited hepatic toxicity (related to centrilobular hypertrophy of hepatocytes which resulted in heavier livers) and renal toxicity (tubular basophilia and an increased incidence of proteinaceous and granular casts) in rats after oral administration at 300 mg/kg/day for 28 days (48). 2,4-DTBP also increased cholesterol and phospholipid in female rats at the dose of 300 mg/kg/day. Susceptibility of newborn rats to 2,4-DTBP was 4 to 5 times higher than that of young rats. In a 1-generation fertility and repeated dose toxicity study, parental generation rats were administered 2,4-DTBP orally for 4 weeks before mating and throughout mating, gestation, and lactation. No treatment-related gross lesions or impairment of reproductive capability were found in the parental generation animals treated with 2,4-DTBP at up to 300 mg/kg/day. However, dietary administration of 2,4-DTBP at 300 mg/kg daily for 13 weeks to F1 generation rats elicited a toxic effect on the growth rate, which was secondary to reduced diet palatability (49). Based on several *in vitro* assays, 2,4-DTBP was shown to be a possible EDC. 2,4-DTBP bound to the rainbow trout estrogen receptor (50). Using an effect-directed analysis-based approach to identify EDCs in multi-contaminated river sediment, 2,4-DTBP was identified as a new environmental human estrogen receptor ligand by using gas chromatography coupled with mass spectrometry (13).

The molecular mechanisms underlying these observations were unclear. Our data showing that 2,4-DTBP disrupted nuclear receptor function via RXR α and previous studies linking 2,4-DTBP to estrogen receptor signaling (13) provided potential clues to identifying mechanisms underlying these toxic effects. There is considerable evidence linking EDCs and obesity and other health problems (10, 51, 52). Many EDCs exert their effects by disrupting the functions of nuclear receptors including PPARs, RXR α , TRs, LXRs, and RARs (10, 53). PPAR γ , RXRs, LXR α , TR β , and RARs play important roles in many physiological and pathological processes such as development, differentiation, and growth (54). The ability of 2,4-DTBP and related chemicals to activate RXR α and RXR α heterodimers implicates it as an EDC. Comprehensive *in vivo* and *in vitro* toxicology studies are warranted to evaluate the potential effects of 2,4-DTBP and related chemicals.

2,4-DTBP concentrations in human urine samples were in the range of 7.3 to 130 ng/mL (0.03 μ M–0.63 μ M). 2,4-DTBP promoted adipogenesis in human MSCs at concentrations of \sim 10 μ M, which is somewhat higher, but not so far from the concentrations observed in limited studies of human urine. Receptor activation studies showed that 2,4-DTBP and 2,4,6-TTBP activated RXRs in the low micromolar to submicromolar range, which is an environmentally relevant level according to the limited human biomonitoring data (18). Considering the ability of tert butyl phenols to target RXR α in the context of multiple receptor heterodimers, as well as their ability to activate ERS, additional large-scale surveys on the burden to the human body of 2,4-DTBP are warranted.

Taken together, the data presented here demonstrated that 2,4-DTBP belongs to a family of compounds whose potential endocrine-disrupting and obesogenic effects can be strongly modulated by their chemical composition. That the United States, and presumably other populations, can be exposed to

endocrine active chemicals such as DTBPs in the absence of specific knowledge of these effects by regulatory agencies means that the population is vulnerable to such chemicals and that current regulatory approaches are not providing protection from their endocrine-disrupting effects. Structure–activity studies such as the present one could help the rational development of safer antioxidants.

Acknowledgments

We thank Dr. Jane Muncke (Food Packaging Forum Foundation) for suggesting 2,4-DTBP as a compound of interest. We acknowledge experimental assistance from the staff of the European Synchrotron Radiation Facility (Grenoble, France) during crystallographic data collection.

Funding

Supported by grants from the National Institutes of Health (ES023316, ES031139) to B.B., by the European Union's Horizon 2020 research and innovation program under grant agreement GOLIATH (825489) to P.B., W.B., and B.B., by the ANSES TOXCHEM (2018/1/020) to P.B. and W.B., by the European Union's Horizon 2021 research and innovation PARC (101057014) programs to P.B. and W.B., and by a grant from China Scholarship Council to X.R. This work was supported by the French Infrastructure for Integrated Structural Biology (FRISBI) ANR-10-INBS-05.

Author Contributions

X.-M.R., R.C.C., Y.H., Y.K., M.G., C.C., and W.B. performed experiments. Xi.-M.R., R.C.C., W.B., P.B., and B.B. designed experiments and interpreted data. X.-M.R., W.B., P.B., and B.B. wrote the manuscript. W.B., P.B., and B.B. acquired funding and supervised the experiments. All authors have given approval to the final version of the manuscript.

Disclosures

B.B. is a named inventor on US patents 5,861,274, 6,200,802, 6,815,168, and 7,250,273 related to PPAR γ . All other authors declare they have nothing to disclose.

Data Availability

Original data generated and analyzed during this study are included in this published article or in the data repositories listed in References. The atomic coordinates and structure factors have been deposited in the PDB under accession code 7NKE (29) and data collection and refinement statistics can be found in Table S1 (30). Supplemental Figures S1–S3 and Table S1 can be found online at Figshare (30).

References

- Hales CM, Fryar CD, Carroll MD, Freedman DS, Ogden CL. Trends in obesity and severe obesity prevalence in US youth and adults by sex and age, 2007–2008 to 2015–2016. *JAMA*. 2018;319(16):1723–1725.
- WHO. Obesity and overweight; 2018. Accessed August 16, 2022. <https://www.who.int/news-room/fact-sheets/detail/obesity-and-overweight>.
- Picon-Ruiz M, Morata-Tarifa C, Valle-Goffin JJ, Friedman ER, Slingerland JM. Obesity and adverse breast cancer risk and

- outcome: mechanistic insights and strategies for intervention. *CA Cancer J Clin*. 2017;67(5):378–397.
- WHO. *Obesity and Overweight Fact Sheet No. 311*. World Health Organization, 2015.
- Stierman B, Afful J, Carroll MD, *et al*. National health and nutrition examination survey 2017–March 2020 prepandemic data files development of files and prevalence estimates for selected health outcomes. National Health Statistics Reports (NHSR No. 158). <http://dx.doi.org/10.15620/cdc:106273>
- Heindel JJ, Blumberg B. Environmental obesogens: mechanisms and controversies. *Annu Rev Pharmacol Toxicol*. 2019;59(1): 89–106.
- Turcot V, Lu Y, Highland HM, *et al*. Protein-altering variants associated with body mass index implicate pathways that control energy intake and expenditure in obesity. *Nat Genet*. 2018;50(1): 26–41.
- Janesick AS, Blumberg B. Obesogens: an emerging threat to public health. *Am J Obstet Gynecol*. 2016;214(5):559–565.
- Heindel JJ, Howard S, Agay-Shay K, *et al*. Obesity II: establishing causal links between chemical exposures and obesity. *Biochem Pharmacol*. 2022;199:115015.
- Heindel JJ, Blumberg B, Cave M, *et al*. Metabolism disrupting chemicals and metabolic disorders. *Reprod Toxicol*. 2017;68:3–33.
- ChemView. United States Environmental Protection Agency; 2016. Accessed August 22, 2022. <https://chemview.epa.gov/chemview>
- Liu R, Lin Y, Ruan T, Jiang G. Occurrence of synthetic phenolic antioxidants and transformation products in urban and rural indoor dust. *Environ Pollut*. 2017;221:227–233.
- Creusot N, Budzinski H, Balaguer P, Kinani S, Porcher JM, Ait-Aïssa S. Effect-directed analysis of endocrine-disrupting compounds in multi-contaminated sediment: identification of novel ligands of estrogen and pregnane X receptors. *Anal Bioanal Chem*. 2013;405(8):2553–2566.
- Sebok A, Vasani Zsigrai A, Helenkár A, Záray G, Molnár-Perl I. Multiresidue analysis of pollutants as their trimethylsilyl derivatives, by gas chromatography-mass spectrometry. *J Chromatogr A*. 2009;20(12):2288–2301.
- Zhao F, Wang P, Lucardi RD, Su Z, Li S. Natural sources and bioactivities of 2,4-Di-tert-butylphenol and its analogs. *Toxins (Basel)*. 2020;12(1):35.
- Yang Y, Hu C, Zhong H, Chen X, Chen R, Yam KL. Effects of ultraviolet (UV) on degradation of irgafos 168 and migration of its degradation products from polypropylene films. *J Agric Food Chem*. 2016;64(41):7866–7873.
- Choi SJ, Kim JK, Kim HK, *et al*. 2,4-Di-tert-butylphenol From sweet potato protects against oxidative stress in PC12 cells and in mice. *J Med Food*. 2013;16(11):977–983.
- Liu R, Mabury SA. Unexpectedly high concentrations of 2,4-di-tert-butylphenol in human urine. *Environ Pollut*. 2019;252(Pt B):1423–1428.
- Chamorro-García R, Kirchner S, Li X, *et al*. Bisphenol A diglycidyl ether induces adipogenic differentiation of multipotent stromal stem cells through a peroxisome proliferator-activated receptor gamma-independent mechanism. *Environ Health Perspect*. 2012;120(7):984–989.
- Janesick AS, Dimastrogiovanni G, Vanek L, *et al*. On the utility of ToxCast™ and ToxPi as methods for identifying new obesogens. *Environ Health Perspect*. 2016;124(8):1214–1226.
- Grün F, Watanabe H, Zamanian Z, *et al*. Endocrine-disrupting organotin compounds are potent inducers of adipogenesis in vertebrates. *Mol Endocrinol*. 2006;20(9):2141–2155.
- Shoucri BM, Martinez ES, Abreo TJ, *et al*. Retinoid X receptor activation alters the chromatin landscape to commit mesenchymal stem cells to the adipose lineage. *Endocrinology*. 2017;158(10): 3109–3125.
- Forman BM, Umehono K, Chen J, Evans RM. Unique response pathways are established by allosteric interactions among nuclear hormone receptors. *Cell*. 1995;81(4):541–550.

24. Balaguer P, Boussioux AM, Demirpence E, Nicolas JC. Reporter cell lines are useful tools for monitoring biological activity of nuclear receptor ligands. *Luminescence*. 2001;16(2):153-158.
25. Kabsch W. XDS. *Acta Crystallogr D Biol Crystallogr*. 2010;66(2):125-132.
26. Evans P. Scaling and assessment of data quality. *Acta Crystallogr D Biol Crystallogr*. 2006;62(1):72-82.
27. Murshudov GN, Skubak P, Lebedev AA, et al. REFMAC5 for the refinement of macromolecular crystal structures. *Acta Crystallogr D Biol Crystallogr*. 2011;67(4):355-367.
28. Emsley P, Cowtan K. Coot: model-building tools for molecular graphics. *Acta Crystallogr D Biol Crystallogr*. 2004;60(12):2126-2132.
29. Carivenc C, Bourguet W. Crystal structure of human RXRalpha ligand and binding domain in complex with 2,4-Di-tert-butylphenol and a coactivator fragment. March 2, 2022 ed. Worldwide PDB Protein Data Bank. March 2, 2022. <https://doi.org/10.2210/pdb7nke/pdb>
30. Ren X-M, Chang RC, Huang Y, et al. 2,4-Di-tert-butylphenol induces adipogenesis in human mesenchymal stem cells by activating retinoid X Receptors. Figshare. Deposited January 18, 2023. <https://doi.org/10.6084/m9.figshare.21919368>
31. Kassotis CD, Vom Saal FS, Babin PJ, et al. Obesity III: obesogen assays: limitations, strengths, and new directions. *Biochem Pharmacol*. 2022;199:115014.
32. Mohajer N, Du CY, Checkcinco C, Blumberg B. Obesogens: how they are identified and molecular mechanisms underlying their action. *Front Endocrinol (Lausanne)*. 2021;12:780888.
33. Tontonoz P, Spiegelman BM. Fat and beyond: the diverse biology of PPARgamma. *Annu Rev Biochem*. 2008;77(1):289-312.
34. Kirchner S, Kieu T, Chow C, Casey S, Blumberg B. Prenatal exposure to the environmental obesogen tributyltin predisposes multipotent stem cells to become adipocytes. *Mol Endocrinol*. 2010;24(3):526-539.
35. Ebisawa M, Umemiya H, Ohta K, et al. Retinoid X receptor-antagonistic diazepinylbenzoic acids. *Chem Pharm Bull (Tokyo)*. 1999;47(12):1778-1786.
36. Kahlen JP, Carlberg C. Allosteric interaction of the 1alpha, 25-dihydroxyvitamin D3 receptor and the retinoid X receptor on DNA. *Nucleic Acids Res*. 1997;25(21):4307-4313.
37. Aranda A, Pascual A. Nuclear hormone receptors and gene expression. *Physiol Rev*. 2001;81(3):1269-1304.
38. Evans RM, Mangelsdorf DJ. Nuclear receptors, RXR, and the big bang. *Cell*. 2014;157(1):255-266.
39. Collins JL, Fivush AM, Watson MA, et al. Identification of a non-steroidal liver X receptor agonist through parallel array synthesis of tertiary amines. *J Med Chem*. 2002;45(10):1963-1966.
40. Santin EP, Germain P, Quillard F, et al. Modulating retinoid X receptor with a series of (E)-3-[4-hydroxy-3-(3-alkoxy-5,5,8,8-tetramethyl-5,6,7,8-tetrahydronaphthalen-2-yl)phenyl]acrylic acids and their 4-alkoxy isomers. *J Med Chem*. 2009;52(10):3150-3158.
41. le Maire A, Grimaldi M, Roecklin D, et al. Activation of RXR-PPAR heterodimers by organotin environmental endocrine disruptors. *EMBO Rep*. 2009;10(4):367-373.
42. Creusot N, Tapie N, Piccini B, et al. Distribution of steroid- and dioxin-like activities between sediments, POCIS and SPMD in a French river subject to mixed pressures. *Environ Sci Pollut Res Int*. 2013;20(5):2784-2794.
43. Li X, Pham HT, Janesick AS, Blumberg B. Triflumizole is an obesogen in mice that acts through peroxisome proliferator activated receptor gamma (PPARγ). *Environ Health Perspect*. 2012;120(12):1720-1726.
44. Castillo AI, Sánchez-Martínez R, Moreno JL, Martínez-Iglesias OA, Palacios D, Aranda A. A permissive retinoid X receptor/thyroid hormone receptor heterodimer allows stimulation of prolactin gene transcription by thyroid hormone and 9-cis-retinoic acid. *Mol Cell Biol*. 2004;24(2):502-513.
45. Fattori J, Campos JL, Doratioto TR, et al. RXR Agonist modulates TR: corepressor dissociation upon 9-cis retinoic acid treatment. *Mol Endocrinol*. 2015;29(2):258-273.
46. Mengeling BJ, Murk AJ, Furlow JD. Trialkyltin retinoid-X receptor agonists selectively potentiate thyroid hormone induced programs of *Xenopus laevis* metamorphosis. *Endocrinology*. 2016;157(7):2712-2723.
47. Kodama S, Matsumoto S, Takamura Y, et al. Structural characterization of 1,3-bis-tert-butyl monocyclic benzene derivatives with agonistic activity towards retinoid X receptor alpha. *Toxicol Lett*. 2022;373:76-83.
48. Hirata-Koizumi M, Hamamura M, Furukawa H, et al. Elevated susceptibility of newborn as compared with young rats to 2-tert-butylphenol and 2,4-di-tert-butylphenol toxicity. *Congenit Anom*. 2005;45(4):146-153.
49. ECHA. Registration profile for 2,4-di-tert butylphenol; 2018. Accessed August 22, 2022. Available from, as of July 27, 2018: <https://echa.europa.eu/registration-dossier/-/registered-dossier/14828/14827/14821>.
50. Tollefsen KE, Julie Nilsen A. Binding of alkylphenols and alkylated non-phenolics to rainbow trout (*Oncorhynchus mykiss*) hepatic estrogen receptors. *Ecotoxicol Environ Saf*. 2008;69(2):163-172.
51. Gore AC, Chappell VA, Fenton SE, et al. EDC-2: the endocrine Society's Second scientific statement on endocrine-disrupting chemicals. *Endocr Rev*. 2015;36(6):E1-E150.
52. Papalou O, Kandaraki EA, Papadakis G, Diamanti-Kandaraki E. Endocrine disrupting chemicals: an occult mediator of metabolic disease. *Front Endocrinol*. 2019;10:112.
53. Hall JM, Greco CW. Perturbation of nuclear hormone receptors by endocrine disrupting chemicals: mechanisms and pathological consequences of exposure. *Cells*. 2019;9(1):13.
54. Yen PM. Classical nuclear hormone receptor activity as a mediator of complex biological responses: a look at health and disease. *Best Pract Res Clin Endocrinol Metab*. 2015;29(4):517-528.