

# Estrogen-Like Activity of Perfluoroalkyl Acids *In Vivo* and Interaction with Human and Rainbow Trout Estrogen Receptors *In Vitro*

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The objectives of this study were to determine the structural characteristics of perfluoroalkyl acids (PFAAs) that confer estrogen-like activity *in vivo* using juvenile rainbow trout (*Oncorhynchus mykiss*) as an animal model and to determine whether these chemicals interact directly with the estrogen receptor (ER) using *in vitro* and *in silico* species comparison approaches. Perfluorooctanoic (PFOA), perfluorononanoic (PFNA), perfluorodecanoic (PFDA), and perfluoroundecanoic (PFUnDA) acids were all potent inducers of the estrogen-responsive biomarker protein vitellogenin (Vtg) *in vivo*, although at fairly high dietary exposures. A structure-activity relationship for PFAAs was observed, where eight to ten fluorinated carbons and a carboxylic acid end group were optimal for maximal Vtg induction. These *in vivo* findings were corroborated by *in vitro* mechanistic assays for trout and human ER. All PFAAs tested weakly bound to trout liver ER with half maximal inhibitory concentration (IC<sub>50</sub>) values of 15.2–289 μM. Additionally, PFOA, PFNA, PFDA, PFUnDA, and perfluorooctane sulfonate (PFOS) significantly enhanced human ER $\alpha$ -dependent transcriptional activation at concentrations ranging from 10–1000 nM. Finally, we employed an *in silico* computational model based upon the crystal structure for the human ER $\alpha$  ligand-binding domain complexed with E2 to structurally investigate binding of these putative ligands to human, mouse, and trout ER $\alpha$ . PFOA, PFNA, PFDA, and PFOS all efficiently docked with ER $\alpha$  from different species and formed a hydrogen bond at residue Arg394/398/407 (human/mouse/trout) in a manner similar to the environmental estrogens bisphenol A and nonylphenol. Overall, these data support the contention that several PFAAs are weak environmental xenoestrogens of potential concern.

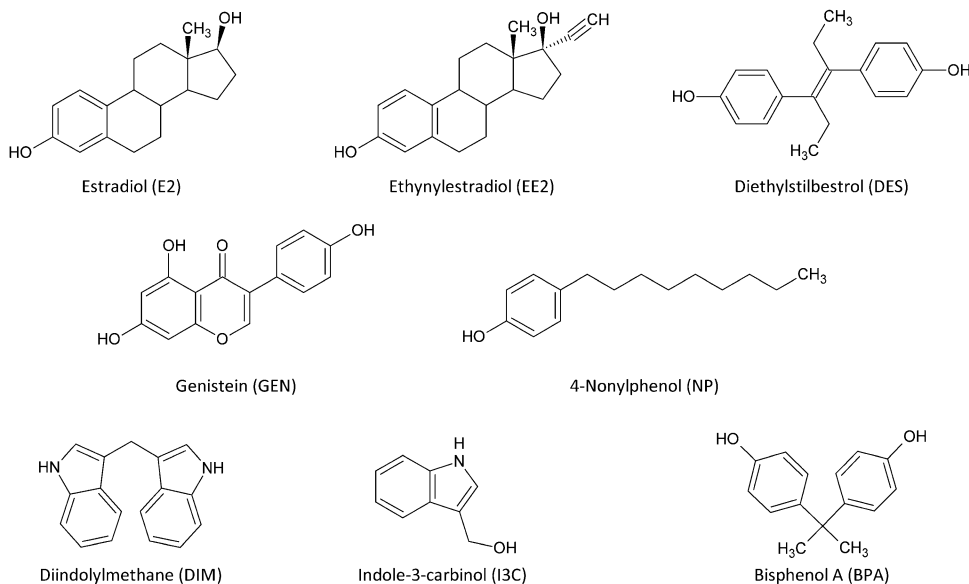
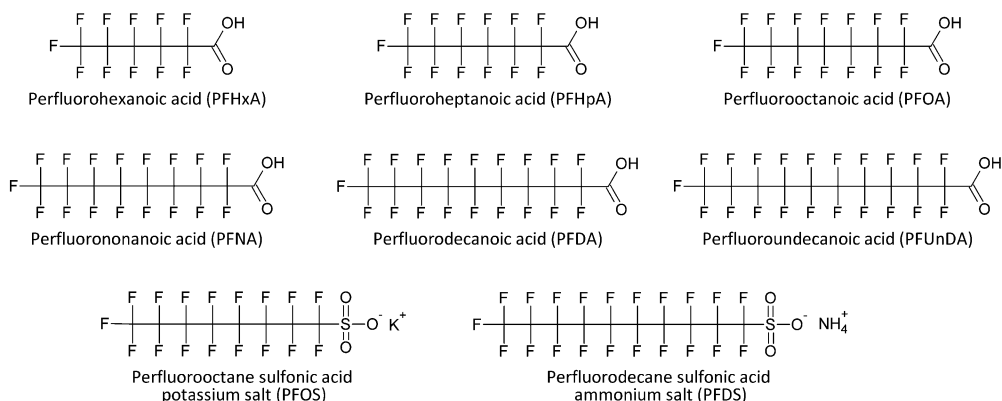
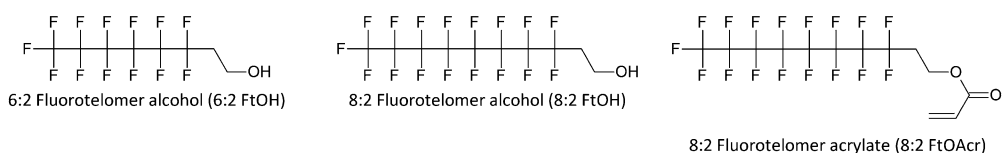
**Key Words:** perfluoroalkyl acid; estrogen; perfluorooctanoic acid; perfluorooctane sulfonate; vitellogenin; molecular docking.

The widespread industrial and commercial use of polyfluorinated chemicals (PFCs) as surfactants and surface protectors for paper and textile coatings, polishes, food

packaging, and fire-retardant foams has led to the pervasive presence of these chemicals in the environment, wildlife, and humans (see reviews by Calafat *et al.*, 2007; Houde *et al.*, 2006). The general structure of PFCs resembles that of fatty acids in that each compound has a hydrophobic polyfluorinated carbon tail of varying length and a functional end group, which provides the basis for classification (Fig. 1). Perfluoroalkyl acids (PFAAs) and fluorotelomers comprise the two major structural groups of PFCs. Human blood levels of perfluorooctanoic acid (PFOA) and perfluorooctane sulfonate (PFOS), the two most commonly studied PFCs, are about 4 and 20 ppb, respectively, although national survey data suggest that these levels have decreased in recent years (Calafat *et al.*, 2007). Levels of PFOS and PFOA in wildlife tend to be higher, in the range of tens to thousands ppb (Kannan *et al.*, 2002). Other PFCs commonly detected in biological samples include perfluorononanoic acid (PFNA), perfluorodecanoic acid (PFDA), perfluoroundecanoic acid (PFUnDA), and perfluorododecanoic acid (PFDoDA).

The toxicology and toxicokinetics of PFOA, as an example PFC, have been thoroughly reviewed by Kennedy *et al.* (2004). PFOA does not accumulate in fatty tissues because of its dual lipophobic and hydrophobic chemical properties but instead binds to blood proteins and is distributed primarily to liver, plasma, and kidney. The measured biological half-life of PFOA varies among species ranging from hours in female rat to days in dogs or rainbow trout (Hanhijarvi *et al.*, 1988; Martin *et al.*, 2003). The estimated half-life of PFOA in humans is nearly 4 years, pointing to a lower capacity for elimination of the compound compared with other species (Olsen *et al.*, 2007). Finally, PFOA is not metabolized or defluorinated *in vivo*, although some fluorotelomer compounds may be metabolized to PFAAs in rodents (Martin *et al.*, 2005; Nabb *et al.*, 2007).

PFOA and other PFCs are members of a large group of chemicals called peroxisome proliferators (PPs), which

**A Estradiol and other xenoestrogens****B Perfluoroalkyl acids (PFAAs)****C Fluorotelomers (FTs)****FIG. 1.** Structures for known xenoestrogens (A), select PFAAs (B), and fluorotelomer (C) compounds are shown.

increase the abundance of hepatic peroxisomes and induce peroxisomal and mitochondrial enzymes involved with  $\beta$ -oxidation, cytochrome P450 fatty acid  $\omega$ -oxidation, and cholesterol homeostasis (reviewed in Bosgra *et al.*, 2005) via ligand-dependent activation of the hepatic PP-activated receptor  $\alpha$  (PPAR $\alpha$ ) (Holden and Tugwood, 1999). Prolonged exposure to PPs, including PFOA, can result in hepatocarcinogenesis (Abdellatif *et al.*, 1991), although marked species differences in susceptibility to PPs have been

observed. Whereas rodents are highly susceptible, humans and nonhuman primates are insensitive or nonresponsive (Lai, 2004), possibly due to low expression of PPAR $\alpha$  in liver (Palmer *et al.*, 1998). Recent evidence points to PPAR $\alpha$ -independent mechanisms of action for increased risk of liver cancer by PPs, suggesting that the assumption that PPAR $\alpha$  agonists pose little risk for human hepatocarcinogenesis may be unsound (reviewed in Guyton *et al.*, 2009). For example, chronic exposure to the

environmental PPAR $\alpha$  agonist di(2-ethylhexyl) phthalate increased liver cancer incidence in PPAR $\alpha$  knockout mice (Ito *et al.*, 2007).

Estrogens have long been known to be associated with cancer in estrogen-responsive tissues including the breast, ovary, and uterus as complete carcinogens, promoters, or possible chemoprotective agents (Cavalieri *et al.*, 2000; Eliassen and Hankinson, 2008; Ho, 2003; Persson, 2000). Moreover, estrogens also influence carcinogenesis in other tissues not usually considered to be estrogen responsive. Experimental studies investigating hepatic cancer in mammalian animal models have implicated estrogens as tumor promoters and in some cases complete carcinogens (reviewed in De Maria *et al.*, 2002; Yager *et al.*, 1991), although the higher rate of hepatic cancer in males suggests the potential for a protective role of estrogens against liver cancer (Villa, 2008; Yeh and Chen, 2010). Several studies in our laboratory using the rainbow trout cancer model have also demonstrated that estrogen exposure promotes hepatocellular carcinogenesis in rainbow trout (Núñez *et al.*, 1989; Orner *et al.*, 1995).

Previously, we reported that PFOA may act via an alternative mechanism of action, not involving peroxisome proliferation, to increase risk of liver cancer in rainbow trout, an animal model that mimics human insensitivity to PPs (Tilton *et al.*, 2008). Dietary exposure to PFOA significantly increased expression of the sensitive estrogen biomarker protein vitellogenin (Vtg) and induced an overall hepatic gene expression profile highly similar to that of 17 $\beta$ -estradiol (E2). We concluded that the cancer-enhancing effects of PFOA in trout were due to novel mechanisms related to estrogen signaling rather than the typical PP response observed for this chemical in rodent models (Tilton *et al.*, 2008). Studies in other model organisms support the concept that certain PFCs may have estrogen-like activity. Liu *et al.* (2007) reported that PFOA, PFOS, and 6:2 fluorotelomer alcohol (6:2FtOH) increased Vtg expression in cultured tilapia hepatocytes, and Wei *et al.* (2007) showed that PFOA increased Vtg expression and caused disruption of gonad development in rare minnows (*Gobiocypris rarus*). Evidence for estrogen-like activity of some fluorotelomers has also been shown *in vitro* (Ishibashi *et al.*, 2007; Maras *et al.*, 2006).

Though several laboratories have shown estrogenic effects of certain PFCs *in vitro* and *in vivo*, it is yet unknown whether any of these chemicals interact directly with the estrogen receptor (ER). This receptor is generally considered to be quite promiscuous because dozens of structurally diverse exogenous chemicals, including phytochemicals, hydroxylated polychlorinated biphenyls, pesticides, and other industrial compounds, have been shown to bind to the receptor with varying degrees of affinity (Blair *et al.*, 2000; Matthews *et al.*, 2000; Olsen *et al.*, 2005). The optimal chemical structure for ligand binding to the ER is a four-ring arrangement with opposing hydroxyl end groups, although long-chain compounds, such as nonylphenol, have been shown to effectively bind to the protein as well (Matthews *et al.*, 2000; Sivanesan *et al.*, 2005).

In this present study, we determined the relationship between PFC structure and estrogen-like activity using the Vtg induction bioassay in rainbow trout to test the hypotheses that PFCs structurally similar to PFOA are estrogenic and that the Vtg response depends largely on chemical structure. We also assessed the interaction of PFAAs with the ER using direct steroid receptor-binding assays for trout liver ER and a gene reporter assay for human ER $\alpha$ . Finally, we employed computational modeling to determine the molecular interaction of these compounds with the ligand-binding domains (LBDs) of human, mouse, and trout ER $\alpha$  proteins.

## MATERIALS AND METHODS

### Chemicals

A complete list of PFCs investigated in this study, including their common names, abbreviations, and registry numbers for reference, is provided in Supplementary table 1. Perfluoroalkyl carboxylic acid and sulfonate compounds were all purchased from Sigma-Aldrich (St Louis, MO), except for perfluorodecane sulfonate (PFDS), which was graciously donated by Dr Jennifer Field (Oregon State University). Fluorotelomer alcohol and acrylate compounds were obtained from SynQuest Labs (Alachua, FL). E2, genistein (GEN), diethylstilbestrol (DES), ethynylestradiol (EE2), indole-3-carbinol (I3C), 3,3'-diindolylmethane (DIM), 4-nonylphenol (NP), and all other reagents were purchased from Sigma-Aldrich or other general laboratory suppliers and were of the highest purity available. [2,4,6,7-<sup>3</sup>H]-17 $\beta$ -estradiol (70 Ci/mmol) was purchased from Perkin Elmer (Waltham, MA).

### Animal Care

Mt Shasta strain rainbow trout were hatched and reared at the Sinnhuber Aquatic Research Laboratory (SARL) at Oregon State University in Corvallis, OR. Fish were maintained in flow-through 375-l tanks at 12°C with activated carbon water filtration on a 12:12 h light:dark cycle. Two weeks prior to experimental treatments, fish were fed a maintenance ration (2% of body weight [bw]) of Oregon Test Diet (OTD), a semipurified casein-based diet with menhaden oil as the lipid. All experimental diets described below were administered for 14 days, and feeding occurred 5 days per week. On day 15, trout were euthanized with an overdose (250 ppm) of tricaine methanesulfonate and necropsied. Body and liver weights were recorded to calculate the liver somatic index (LSI = liver weight/bw  $\times$  100), and blood samples from each donor animal were obtained. The plasma fraction of each blood sample was obtained by centrifugation at 850  $\times$  g and then frozen at -80°C. All procedures for treatment, handling, maintenance, and euthanasia of trout used in this study were approved by the Oregon State University Institutional Animal Care and Use Committee.

### In Vivo Experimental Procedures

**Experiment 1—subchronic dietary exposure to PFCs.** A variety of structurally diverse PFCs were chosen to test *in vivo* in juvenile rainbow trout, including perfluoroalkyl carboxylic acids with 6–14 fluorinated carbons, two perfluoroalkyl sulfonates, and three fluorotelomers. Eleven-month-old juvenile trout (initial bw approximately 70 g) were distributed to holding tanks 1 week prior to the start of feeding experimental diets. Test chemicals were dissolved in dimethylsulfoxide (DMSO) vehicle ( $\leq$  0.5 ppm) and added directly to the oil portion of the custom trout OTD diet. PFCs were administered at a diet concentration of 250 ppm calculated with respect to diet wet weight (approximately 5 mg/kg bw/day) to six fish per treatment group. Additionally, PFOA, PFNA, PFDA, and PFUnDA were selected for testing in a simple mixture study. Two additional dietary groups at 5 and 50 ppm (0.1 and 1 mg/kg

bw/day) were added to the experiment for these four compounds so that three concentrations in total were tested for each individual chemical (5, 50, and 250 ppm). Mixture treatments were prepared by combining equal mass amounts of each test compound at each concentration level. Thus, mixture A consisted of 5 ppm each PFOA, PFNA, PFDA, and PFUnDA for a total perfluoroalkyl carboxylic acid concentration of 20 ppm. Similarly, mixture B was made by combining 50 ppm each chemical for a total diet concentration of 200 ppm; finally, mixture C was prepared by combining 250 ppm each compound for total concentration of 1000 ppm. The control group ( $N = 24$ , six fish in four replicate tanks) was fed OTD diet with no modifications; the vehicle control group was fed diet with 0.5 ppm DMSO ( $N = 6$ ), and 5 ppm E2 was included as a positive control ( $N = 12$ , six fish in two replicate tanks). The additional fish were included in the E2 positive and the control groups to account for multiple sampling days.

**Experiment 2—dose-dependent induction of Vtg by PFOA and PFDA.** The goal of this experiment was to determine the dose-dependent impact of dietary exposure to PFOA and PFDA on trout blood Vtg levels. PFOA and PFDA were selected as test compounds for this experiment because of their common detection in wildlife and humans and the observed robust induction of Vtg by both chemicals in experiment 1. Each experimental group consisted of a single tank of 5-month-old individual juvenile trout with an initial approximate bw of 30 g. The control group ( $N = 16$ ) was fed OTD diet with no modifications, and 5 ppm E2 diet was used as a positive control ( $N = 16$ ). Eight trout per treatment group were fed experimental diets containing either PFOA or PFDA at concentrations of 0.026, 0.128, 0.64, 3.2, 16, 80, 400, or 2000 ppm calculated with respect to diet wet weight (range of 0.52  $\mu\text{g}/\text{kg}$  bw/day up to 40  $\text{mg}/\text{kg}$  bw/day). In this study, the test compounds were added directly to the oil portion of the diet, which was gently heated ( $< 50^\circ\text{C}$ ) to aid in chemical dissolution.

#### Vtg ELISA

Blood plasma Vtg was measured following an ELISA method previously described (Shilling and Williams, 2000) with modifications as outlined by Benninghoff and Williams (2008). For this study, the interassay coefficient of variance for the Vtg ELISA was 7.1% with an assay detection limit of 6.25 ng/ml.

#### Measurement of Palmitoyl Coenzyme A Oxidation and Catalase Activity

Peroxisomal fractions of liver tissues were prepared by differential centrifugation, and catalase activity was assessed following the methods described by Orner *et al.* (1995). Beta-oxidation of palmitoyl coenzyme A (CoA) was measured using a spectrophotometric assay as previously described (Mitchell *et al.*, 1985).

#### PFOA and PFDA Blood Content by Liquid Chromatography-Tandem Mass Spectrometry

Complete methods for the determination of PFOA and PFDA content in trout blood plasma samples are provided in the Supplementary materials including details on the instrumentation, quality control, sample preparation, liquid chromatography-tandem mass spectrometry/LC-MS/MS conditions, and quantitation. Briefly, measurement of PFOA and PFDA was performed at the 3M Medical Department, and this laboratory was blinded to any information on the effects of PFOA or PFDA on Vtg expression in trout. A Shimadzu High Performance Liquid Chromatography (HPLC) system free of contaminating polytetrafluoroethylene-based materials was used in conjunction with an API 5000 triple stage quadrupole mass spectrometer. Samples were prepared by solid phase extraction with Waters Oasis HLB cartridges. PFOA and PFDA standard materials were the free acid linear isoforms, and internal dual  $^{13}\text{C}$ -labeled PFOA and PFDA standards were used for quantitation.

#### Trout Liver ER-Binding Studies with PFCs

**Preparation of trout liver cytosol.** Juvenile trout were fed control OTD diet ( $N = 24$ ) or supplemented with 5 ppm E2 ( $N = 24$ ). Livers were pooled in groups of six, homogenized in two volumes of ice-cold TEMS buffer (10mM

Tris-HCl, 1mM EDTA, 12mM monothioglycerol, 20mM sodium molybdate, 10% vol/vol glycerol, and 0.1mM phenylmethanesulfonyl fluoride), and centrifuged at  $100,000 \times g$  for 15 min at  $4^\circ\text{C}$ . Two volumes of ice-cold TEMS buffer containing 5% wt/vol activated charcoal and 0.5% wt/vol dextran were added to the supernatant, which was mixed, incubated on ice for 15 min, and then centrifuged at  $100,000 \times g$  for 90 min at  $4^\circ\text{C}$ . The upper lipid layer was removed, and the remaining supernatant was spun at  $100,000 \times g$  for an additional 30 min. Again, the lipid layer was aspirated, and the resulting supernatant (the cytosol fraction) was collected, aliquoted, and stored at  $-80^\circ\text{C}$  until use.

**ER saturation binding assays.** Saturation analyses were performed by incubating 200  $\mu\text{l}$  of charcoal-stripped cytosol (5 mg protein per milliliter) with 25  $\mu\text{l}$  of [ $^3\text{H}$ ]-E2 (final concentration 0.1–25nM) in TEMS buffer for 24 h at  $12^\circ\text{C}$ . ER saturation binding assays were performed with separate cytosol preparations from control and E2-exposed trout. Nonspecific binding was determined by incubating cytosol with [ $^3\text{H}$ ]-E2 and DES (10–2.5 $\mu\text{M}$ ) and was defined as the fraction of total binding displaced by 100-fold excess DES. Free steroid was separated from bound by incubation with 500  $\mu\text{l}$  of TEMS with 2.5% wt/vol activated charcoal and 0.25% wt/vol dextran for 30 min on ice followed by centrifugation at  $3000 \times g$  for 10 min. Bound [ $^3\text{H}$ ]-E2 was measured in the supernatant by liquid scintillation counting (Beckman Instruments Inc., Fullerton, CA).

**ER competitive binding assays.** Competitive binding assays were performed by incubating 200  $\mu\text{l}$  charcoal-stripped cytosol (5 mg protein per milliliter, 200 microliters per assay tube) extracted from livers of E2-exposed trout, 25  $\mu\text{l}$  of [ $^3\text{H}$ ]-E2 in TEMS buffer at a final concentration of 4nM (near saturating concentration), and 25  $\mu\text{l}$  of cold competitor for 24 h at  $12^\circ\text{C}$ . Competitor stock solutions were prepared in TEMS buffer with DMSO (5% final assay concentration). Bound [ $^3\text{H}$ ]-E2 was measured as described above. Competitors were assayed according to the anticipated affinity of the putative ligand (1pM–10 $\mu\text{M}$  for strong binders and 10 $\mu\text{M}$ –100mM for weak binders). Maximum nonspecific binding was determined using 100 $\mu\text{M}$  E2 and was subtracted from all total count values to determine specific binding.

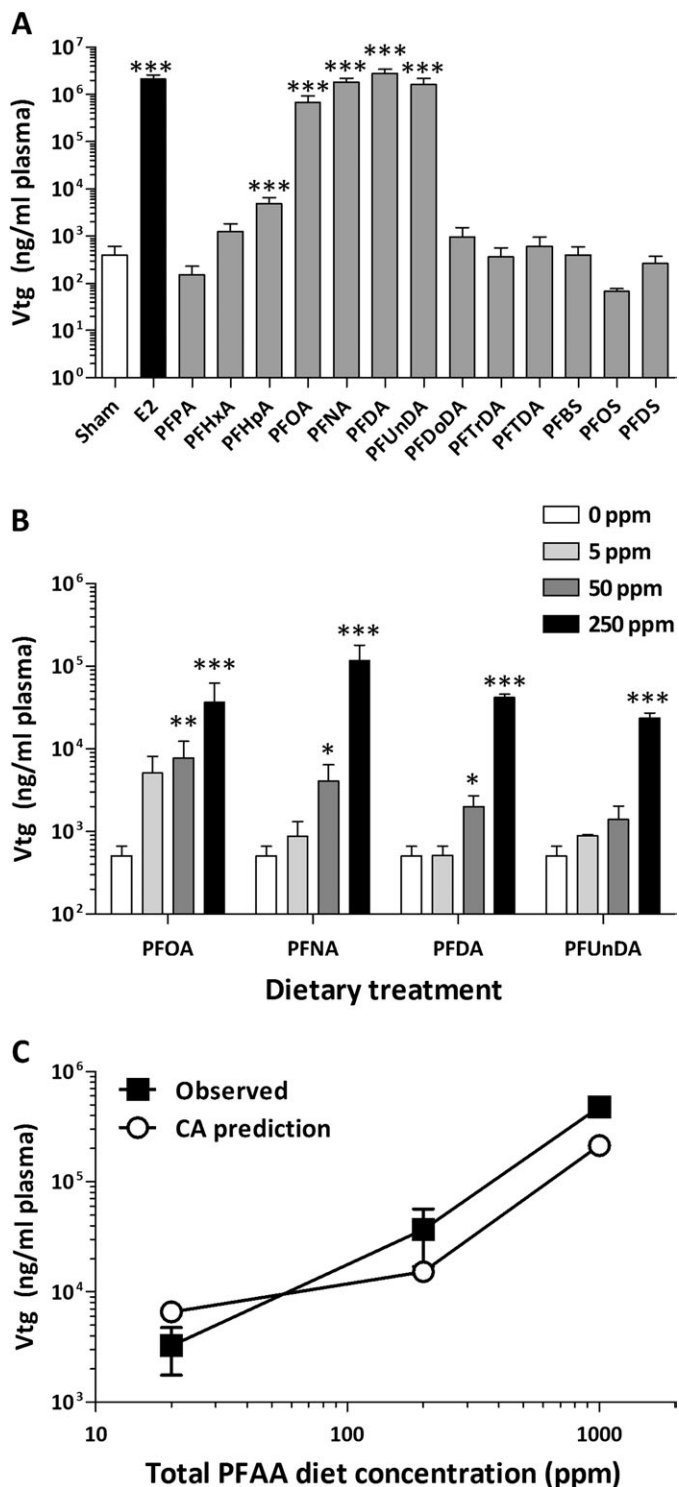
#### Human ER $\alpha$ Gene Reporter Assay

Human embryonic kidney (HEK-293T) cells were cultured in Dulbecco's modified Eagle's medium (DMEM) with L-glutamine (Mediatech, Manassas, VA) supplemented with 10% fetal bovine serum (FBS, Tissue Culture Biologicals, Seal Beach, CA) and 10,000 U/ml penicillin-streptomycin (Mediatech) in a humidified, 5%  $\text{CO}_2$  atmosphere. HEK-293T cells were seeded in 96-well microplates at 10,000 cells per well in 100  $\mu\text{l}$  phenol red-free DMEM with 10% charcoal-stripped FBS. Cells were cotransfected 24 h after seeding with the following: 70 ng of the XTEL luciferase reporter plasmid containing a consensus estrogen-responsive element (ERE) sequence from the *Xenopus* Vtg promoter (graciously donated by Dr Michael Garabedian, New York University School of Medicine) and 10 ng of the human ER $\alpha$  expression vector (Kolluri *et al.*, 2003), an independent  $\beta$ -galactosidase reporter driven by a cytomegalovirus (CMV) promoter and pcDNA for DNA input normalization (Invitrogen, Carlsbad, CA). Each well received 0.2  $\mu\text{l}$  of PLUS reagent with 0.2  $\mu\text{l}$  of lipofectamine (Invitrogen). Twenty-four hours after transfection, cells were treated with E2, PFOA, PFNA, PFDA, PFUnDA, PFOS, or 8:2FtOH at concentrations ranging from 1 to 1000nM for 24 h. Cells were then lysed and measured for reporter gene activity as previously described (Bisson *et al.*, 2009). Raw luminescence data were normalized by  $\beta$ -galactosidase activity to account for any variability in transfection efficiency; then, values for ER $\alpha$  gene reporter activity were expressed as fold induction with respect to vehicle-only cells.

#### In Silico Modeling of PFC Interaction with Human, Mouse, and Trout ER $\alpha$ Proteins

**Molecular modeling.** The 3D coordinates of the human ER $\alpha$  (hER $\alpha$ ) LBD complexed with the natural agonist E2 were taken from the crystal structures available in the Protein Data Bank (accession 1ERE) (Brzozowski *et al.*, 1997).





**FIG. 2.** Effects of subchronic dietary exposure to select PFAAs and fluorotelomers on plasma Vtg concentration. (A) Mean blood plasma Vtg values are shown on a logarithmic scale + SEM ( $N = 18$  for control,  $N = 12$  for E2, and  $N = 6$  for all other treatments). The diet concentration was 250 ppm (approximately 5 mg/kg bw/day) for all perfluoroalkyl compounds and 5 ppm for E2, the positive control. (B) For individual chemical treatments, fish were fed diets containing 0, 5, 50, or 250 ppm of PFOA, PFNA, PFDA, and PFUnDA. Mean blood plasma Vtg values are shown on a logarithmic scale +

Following protein sequence alignment of the LBD regions for the mouse *Esr1* and trout *era1* proteins (UniProt accessions P19785 and P16058, respectively) using ClustalW2 (EMBL-EBI), homology models for mouse  $ER\alpha$  (m $ER\alpha$ ) and rainbow trout  $ER\alpha$  (r $ER\alpha$ ) proteins were built based upon 1ERE as the 3D-template with Molsoft ICM v3.5-1p (Molsoft, L.L.C., La Jolla, CA). Rainbow trout have two isoforms of  $ER\alpha$ , *era1* and *era2* (Nagler *et al.*, 2007); because *era1* is the  $ER\alpha$  isoform abundantly expressed in the liver, we selected this sequence for constructing the homology model. The percent identities of the mouse *Esr1* and trout *era1* protein sequences to that of the human  $ER\alpha$  were 89 and 46%, respectively. All models were energetically refined in the internal coordinate space with Molsoft ICM v3.5-1p (Cardozo *et al.*, 1995).

**Molecular docking.** The receptor is represented by five types of interaction potentials: (1) van der Waals potential for a hydrogen atom probe, (2) van der Waals potential for a heavy-atom probe (generic carbon of 1.7 Å radius), (3) optimized electrostatic term, (4) hydrophobic terms, and (5) loan-pair-based potential, which reflects directional preferences in hydrogen bonding. The energy terms are based on the Merck Molecular Force Field to account for solvation free energy and entropic contribution (Totrov and Abagyan, 2001). Modified intermolecular terms such as soft van der Waals and hydrogen bonding, as well as a hydrophobic term, are added. Conformational sampling is based on the biased probability Monte Carlo procedure, which randomly selects a conformation in the internal coordinate space and then makes a step to a new random position independent of the previous one, but according to a predefined continuous probability distribution. It has also been shown that after each random step, full local minimization greatly improves the efficiency of the procedure. In the ICM-virtual ligand screening (VLS) procedure (Molsoft ICM v3.5-1p), the ligand scoring is optimized to obtain maximal separation between the binders and nonbinders. Each compound is assigned a score according to its fit within the receptor; this ICM score accounts for continuum and discrete electrostatics, hydrophobicity, and entropy parameters (Abagyan *et al.*, 1994; Totrov and Abagyan, 1997, 2001).

#### Data Analysis

For *in vivo* experiments, an individual animal consisted of a single biological replicate, and the sample size is indicated in each figure legend or table. For most experiments, a one-way ANOVA test with Dunnett's multiple comparisons *post hoc* test was employed for statistical analyses of the data (Prism 5, GraphPad Software, LaJolla, CA). For analyses of Vtg data, all values were  $\log_{10}$  transformed to equalize variance prior to statistical analyses. A significant effect of an experimental treatment was inferred when  $p < 0.05$ .

## RESULTS

### Experiment 1—Subchronic Dietary Exposure to PFCs

The objective of this experiment was to determine the impact of subchronic dietary exposure to structurally diverse PFCs, including PFAAs and fluorotelomers, on Vtg expression and to determine whether a structure-activity relationship was evident. As expected, short-term dietary exposure to 5 ppm E2 caused

SEM ( $N = 18$  for control and  $N = 6$  for all other treatments). (C) The mixture treatments were prepared by adding equal amounts of each constituent chemical to achieve total concentrations of 20, 200, and 1000 ppm as described in the "Materials and Methods" section. Closed squares represent the measured mean blood plasma Vtg values  $\pm$  SEM for the mixture treatments ( $N = 6$ ). Open circles represent the predicted level of Vtg response at each concentration calculated using a simple CA model. \* $p < 0.05$ , \*\* $p < 0.01$ , and \*\*\* $p < 0.001$  compared with control (for B, within a diet group) as determined by one-way ANOVA with Dunnett's multiple comparisons *post hoc* test.

a large > 10,000-fold increase in plasma Vtg levels (Fig. 2A) compared with control fish. Exposure to small perfluoroalkyl carboxylic acids with six or seven fluorinated carbons did not significantly alter plasma Vtg levels, whereas medium and long-chain perfluoroalkyl carboxylic acids with 8–14 fluorinated carbons significantly induced Vtg expression by 7- to 190-fold. PFOA, PFNA, PFDA, and PFUnDA were the most potent inducers of Vtg, although the level of response was substantially lower than that of E2. The perfluoroalkyl sulfonates PFOS and PFDS induced a minor three- to fourfold increase in blood Vtg levels. Overall, the fluorotelomers tested had little effect on plasma Vtg levels, with only one compound, 6:2FtOH, causing a modest eightfold increase.

The perfluoroalkyl carboxylic acids PFOA, PFNA, PFDA, PFUnDA, and PFDoDA all caused a significant increase in relative liver size, though the response to PFDoDA was unexpectedly high (Supplementary table 2). Although only the fluorotelomer compound 6:2FtOH was effective in increasing Vtg expression, 8:2FtOH and 8:2FtOAc caused a significant increase in relative liver weight. Overall, for this dietary screening study, there was a significant correlation between Vtg response and LSI with a Spearman's correlation  $r$  value of 0.62 ( $p = 0.022$ ).

Four perfluoroalkyl carboxylic acids, PFOA, PFNA, PFDA, and PFUnDA, were tested at diet concentrations ranging from 5 to 250 ppm and also in a quaternary mixture (20–1000 ppm). For each individual chemical, a significant concentration-dependent induction of Vtg expression was observed ( $p < 0.0001$  by one-way ANOVA) (Fig. 2B). Bonferroni *post hoc* tests revealed that PFOA, PFNA, and PFDA exposures at 50 and 250 ppm caused significant increases in plasma Vtg levels, whereas the response to PFUnDA was only significant at 250 ppm. To calculate the predicted response to the PFOA/PFNA/PFDA/PFUnDA mixture, a simple concentration addition (CA) model was employed, where the response for each individual compound was summed at each concentration level (Fig. 2C). For example, at the middle exposure, Vtg levels of 7726 ng/ml for PFOA + 4083 ng/ml for PFNA + 2008 ng/ml for PFNA + 1410 ng/ml for PFUnDA yielded a CA prediction model value of 15,227 ng/ml for the quaternary mixture. At the lowest concentration level, the observed response to the carboxylic acid mixture was about half of that predicted by the CA model. Alternatively, at the medium and higher diet concentrations, the mixture treatment induced Vtg at levels 1.8- and 2.2-fold higher, respectively, than the predicted response. As these *in vivo* data sets were limited in the number of exposure levels examined, a nonlinear regression analysis for curve comparisons was not performed.

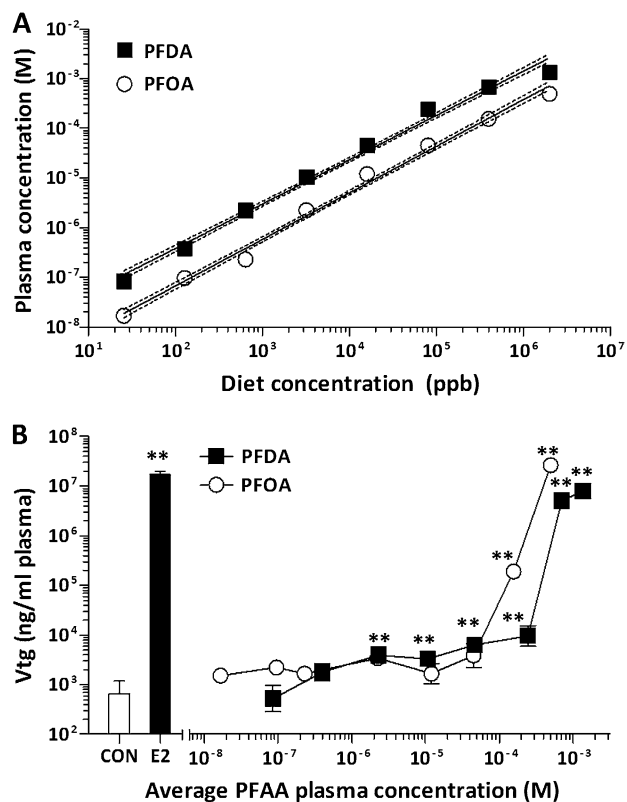
Finally, palmitoyl CoA oxidation and catalase activity were measured as markers of peroxisome proliferation; a subset of the compounds tested in experiment 1 were selected for

enzyme activity analyses, including those chemicals that strongly induced Vtg as well as representative compounds from each structural group. None of the test agents significantly increased peroxisomal palmitoyl CoA oxidation or catalase activity in trout liver, whereas 6:2FtOH and 8:2FtOH apparently decreased catalase activity (Supplementary fig. 1).

#### *Experiment 2—Dose-Dependent Induction of Vtg by PFOA and PFDA*

**PFOA and PFDA levels in trout blood.** A strong linear dose-response relationship between the concentration of PFOA or PFDA administered in the diet and the amount of compound measured in trout blood plasma was observed (Fig. 3A) with linear regression  $R^2$  values of 0.86 and 0.95, respectively. However, at the very highest concentration of PFDA tested (2000 ppm), the measured blood concentration of PFDA was slightly lower than indicated by the linear model, suggesting that this or higher dietary concentrations may exceed the limits for PFDA absorption in trout. The lowest dietary concentration tested (25.6 ppb or 0.5  $\mu\text{g}/\text{kg}$  bw/day) resulted in blood levels of these compounds in the low nanomolar range:  $17.3 \pm 1.82\text{nM}$  for PFOA and  $85.8 \pm 4.79\text{nM}$  for PFDA, whereas the highest dietary concentrations (2000 ppm or 40 mg/kg bw/day) resulted in blood levels of  $508 \pm 35\mu\text{M}$  for PFOA and  $1.36 \pm 0.11\text{mM}$  PFDA.

**Dose-dependent induction of Vtg by PFOA and PFDA.** PFOA and PFDA caused a significant dose-dependent induction of Vtg expression in trout with a maximal response-similar to that of E2 ( $p < 0.0001$  by separate one-way ANOVA for PFOA and PFDA) (Fig. 3B). Bonferroni *post hoc* tests indicated that the lowest observed effect level (LOEL) for PFOA exposure was at the 80 ppm diet (1.6 mg/kg bw/day) level or  $35.4\mu\text{M}$  PFOA in blood plasma (Table 1). Alternatively, the LOEL for PFDA was much lower at 0.64 ppm (12.8  $\mu\text{g}/\text{kg}$  bw/day) or  $2.0\mu\text{M}$  PFDA in blood plasma. Although PFOA did not induce a significant increase in Vtg below the relatively high diet concentration of 80 ppm, the dose-response curve for this compound was substantially left shifted compared with that of PFDA, resulting in a substantially lower estimated effective concentration at 50% response ( $\text{EC}_{50}$ ) value,  $182\mu\text{M}$  for PFOA compared with  $360\mu\text{M}$  for PFDA. However, this lower  $\text{EC}_{50}$  value did not correlate with a lower PFOA dietary exposure, likely a result of the higher half-life and lower elimination rate of PFDA in trout compared with PFOA (Martin *et al.*, 2003). The corresponding amount of dietary PFOA to achieve the  $\text{EC}_{50}$  value of  $182\mu\text{M}$  is 458 ppm, whereas a dietary PFDA concentration of 214 ppm corresponds with its  $\text{EC}_{50}$  value of  $360\mu\text{M}$  (Table 1). A dose-dependent effect of increasing diet concentrations of PFOA and PFDA on relative liver size was less evident in this experiment, as only the highest diet concentrations significantly increased LSI values, 2000 ppm for PFOA and > 400 ppm for PFDA (Supplementary table 3).



**FIG. 3.** Dose-dependent effects of subchronic dietary exposure to PFOA and PFDA on chemical concentrations in blood plasma and Vtg expression. (A) Blood plasma concentrations of PFOA and PFDA following 2-week exposure to a broad range of diet concentrations (0.0256–2000 ppm) are shown ( $N = 8$ ) in a log-log plot. Results of linear regression analyses of nontransformed data are represented by the solid lines, whereas the 95% confidence intervals are represented by the dashed lines. (B) Mean blood plasma Vtg values are shown  $\pm$  SEM ( $N = 8$ ) for PFOA- and PFDA-exposed fish plotted according to the measured blood concentrations of PFOA and PFDA, respectively. \* $p < 0.05$  and \*\* $p < 0.01$  compared with the 0 ppm treatment group (CON) as determined by one-way ANOVA with Dunnett's multiple comparisons *post hoc* test.

#### Trout Hepatic ER-Binding Studies

Radiolabeled E2 binding in liver cytosol extract from E2-exposed rainbow trout showed a single class of high-affinity binding sites that saturated at 5–10 nM [ $^3\text{H}$ ]-E2 with an equilibrium dissociation constant ( $K_d$ ) value of  $4.4 \pm 0.7$  nM and measured binding capacity ( $B_{\text{max}}$ ) value of  $152 \pm 4.9$  pM (38 pmol/g protein) (Supplementary fig. 2). At the near saturating concentration of 4 nM [ $^3\text{H}$ ]-E2, a larger fraction of total binding was specific binding (about 75%) for the E2-exposed liver cytosol compared with only about 50% for the control cytosol. Therefore, all competitive binding experiments were conducted using the cytosol protein from E2-exposed fish because of the greater dynamic range in which to perform these assays.

In the radioligand competitive binding assays, the natural hormone E2 binds to the trout hepatic ER with an  $\text{IC}_{50}$  value of 13.9 nM (Fig. 4A, Table 2). The synthetic estrogen DES was

**TABLE 1**  
Dose-Response Curve Parameters for PFOA and PFDA  
Induction of Vtg

Parameter	PFOA		PFDA	
	Diet (ppm)	Blood level ( $\mu\text{M}$ )	Diet (ppm)	Blood level ( $\mu\text{M}$ )
LOEL	<b>80</b>	35.4	<b>0.64</b>	2.00
NOEL	<b>16</b>	7.83	<b>0.128</b>	0.48
$\text{EC}_{50}$	458	<b>182</b>	214	<b>360</b>
$\text{EC}_{10}$	141	<b>60.1</b>	118	<b>212</b>

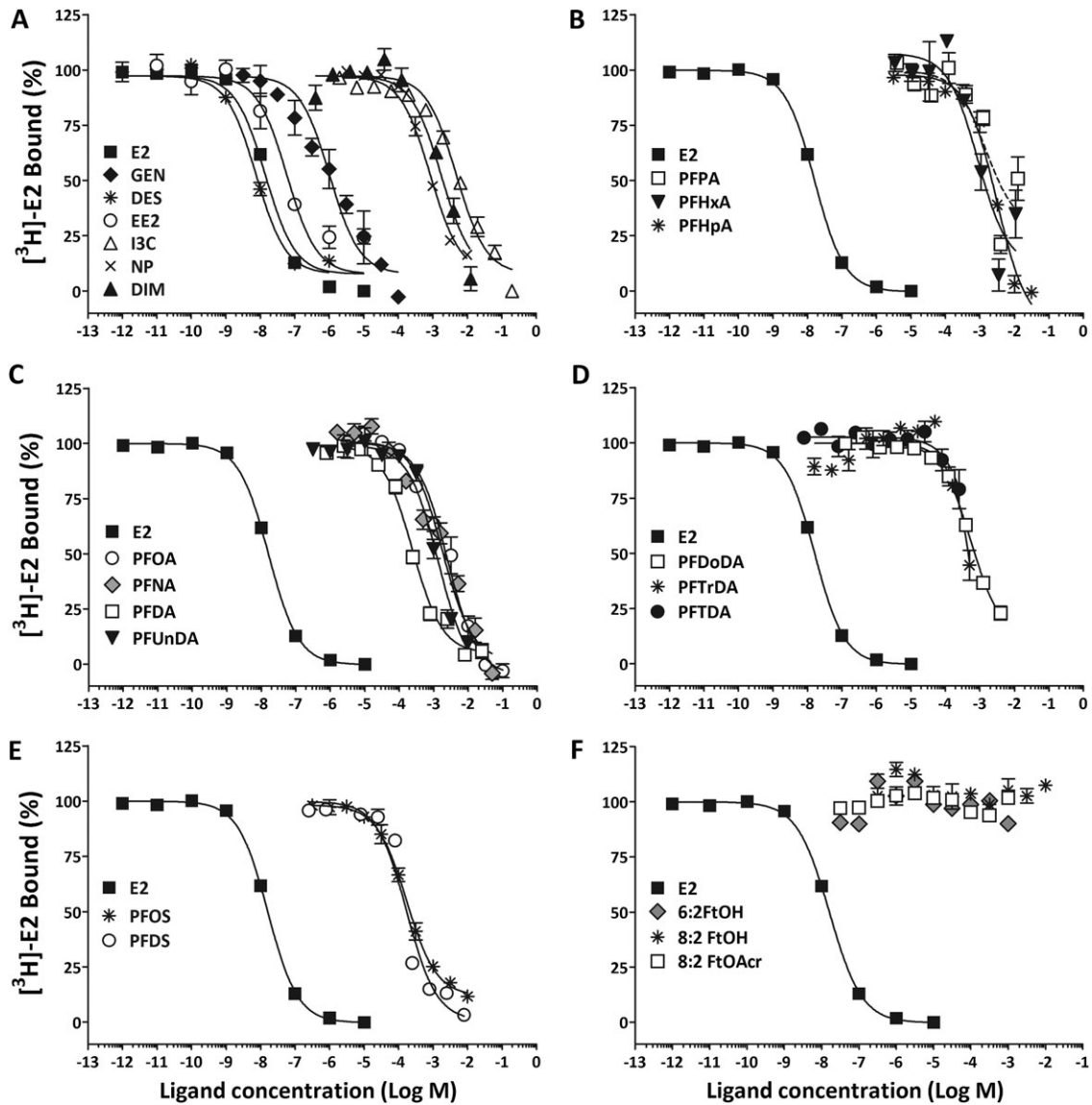
*Note.* Experimentally determined values (diet concentrations) are shown in bold for the LOEL and NOEL estimates, whereas the corresponding blood levels were extrapolated from the standard curve shown in Figure 3A (diet concentration vs. blood level). Experimentally calculated values (blood levels) are shown in bold for  $\text{EC}_{50}$  and  $\text{EC}_{10}$  estimates, whereas the corresponding dietary exposure level was extrapolated from the dose-response curve shown in Figure 3B (Vtg induction vs. blood level). NOEL, no observed effect level;  $\text{EC}_{10}$ , effective concentration at 10% response.

also a strong binder with a relative binding affinity (RBA) value nearly twice that of E2, whereas EE2 binding was about fivefold weaker. The dietary phytoestrogen GEN had moderate binding affinity for the trout liver ER, with an RBA value of about 1%. Though NP, DIM, and I3C all effectively displaced [ $^3\text{H}$ ]-E2 from the ER, these compounds were classified as weak-affinity ligands with  $\text{IC}_{50}$  values in the high micromolar range and RBA values well below 0.01%.

An apparent relationship between the PFAA carbon chain length and the strength of binding to the trout ER was evident (Fig. 4, Table 2). The shortest (perfluoropentanoic acid) and longest (PFDoDa, PFTrDA, and perfluorotetradecanoic acid) compounds did not completely displace E2 from the ER. Compounds of medium size (7–11 fluorinated carbons), including perfluorohexanoic acid (PFHpA), PFOA, PFNA, PFDA, and PFUnDA, had the highest affinities of the carboxylic acids for the ER and completely displaced E2 from the receptor. Because the calculated RBA values for these compounds were more than 10,000-fold lower than that of E2, they were classified as weak or very weak binders. PFDA had the highest affinity of the carboxylic acids tested for the trout liver ER with  $\text{IC}_{50}$  and RBA values comparable to that of the weak environmental estrogen NP. PFOS and PFDS were also weak ligands for the trout liver ER, with RBA values comparable to that of the similarly sized carboxylic acid PFDA. In contrast, none of the fluorotelomers tested displaced E2 from the receptor and were classified as nonbinders in this study (Fig. 4F).

#### Human ER $\alpha$ Gene Reporter Assay

For all test compounds evaluated, the presence of the hER $\alpha$ -expressing plasmid was required to increase gene reporter activity (Fig. 5), although the assay had



**FIG. 4.** Binding of known xenoestrogens and various PFCs to the trout hepatic ER. Competition assays were performed with 1 mg cytosolic protein in the presence of 4nM [<sup>3</sup>H]-estradiol and increasing concentrations of test ligands, including known xenoestrogens (A), perfluoroalkyl carboxylic acids of increasing carbon chain length (B–D), perfluoroalkyl sulfonic acids (E), and fluorotelomers (F). Symbols represent mean ± SEM specific binding for each test chemical.

a somewhat limited dynamic range because of fairly high basal activity of the gene reporter under the culture conditions described above. (For this reason,  $EC_{50}$  and relative activity values were not calculated for this assay.) Treatment of cells with 1–1000nM E2 caused a significant, concentration-dependent induction of hER $\alpha$  gene reporter activity, with a maximal response of nearly threefold observed. PFOA, PFNA, and PFDA significantly induced gene reporter activity up to 2.5-fold at concentrations of 100–1000nM. PFOS appeared to be the most effective of the PFAAs tested, with significant induction of gene reporter activity observed at 1nM. Alternatively, PFUnDA and 8:2FtOH did not significantly alter gene reporter activity.

#### Molecular Docking Studies with hER $\alpha$ , mER $\alpha$ , and rtER $\alpha$

For this comparative molecular docking study, we included human and trout ER $\alpha$  proteins as well as the mouse ER $\alpha$ , as most of the toxicological assessments for this compound class have been performed in mouse models. The LBD regions of the human, mouse, and trout ER $\alpha$  proteins are highly homologous, with perfect identity at amino acid sites previously recognized as key for hydrogen bonding with E2 (Supplementary fig. 3) (Brzozowski *et al.*, 1997; Marchand-Geneste *et al.*, 2006). The molecular models for human, mouse, and trout ER $\alpha$  proteins were constructed using the experimentally resolved crystal structure available for hER $\alpha$  and were energetically minimized in the internal



TABLE 2  
IC<sub>50</sub> and RBA Values for Competitive Ligands for the Trout Liver ER

Treatment	Abbreviation	IC <sub>50</sub> (M) <sup>a</sup>	RBA (%) <sup>b</sup>	Classification <sup>c</sup>
Xenoestrogens				
Estradiol	E2	1.39 × 10 <sup>-8</sup>	100	S
Diethylstilbestrol	DES	7.59 × 10 <sup>-9</sup>	186	S
Ethinylestradiol	EE2	5.89 × 10 <sup>-8</sup>	23.7	S
Genistein	GEN	1.01 × 10 <sup>-6</sup>	1.38	M
4-Nonylphenol	NP	8.42 × 10 <sup>-4</sup>	0.0017	W
Indole-3-carbinol	I3C	4.81 × 10 <sup>-3</sup>	0.0003	W
3,3'-Diindolylmethane	DIM	1.76 × 10 <sup>-3</sup>	0.0008	W
Perfluoroalkyl carboxylic acids				
Perfluoropentanoic acid	PFFA	2.89 × 10 <sup>-3</sup>	0.0005	VW
Perfluorohexanoic acid	PFHxA	1.22 × 10 <sup>-3</sup>	0.0011	W
Perfluoroheptanoic acid	PFHpA	1.78 × 10 <sup>-3</sup>	0.0008	VW
Perfluorooctanoic acid	PFOA	1.82 × 10 <sup>-3</sup>	0.0008	VW
Perfluorononanoic acid	PFNA	1.63 × 10 <sup>-3</sup>	0.0009	VW
Perfluorodecanoic acid	PFDA	2.34 × 10 <sup>-4</sup>	0.0060	W
Perfluoroundecanoic acid	PFUnDA	1.01 × 10 <sup>-3</sup>	0.0014	W
Perfluorododecanoic acid	PFDODA	6.51 × 10 <sup>-4</sup>	0.0021	W
Perfluorotridecanoic acid	PFTTrDA	5.22 × 10 <sup>-4</sup>	0.0027	W
Perfluorotetradecanoic acid	PFTDA	L	L	
Perfluoroalkyl sulfonates				
Perfluorooctane sulfonic acid potassium salt	PFOS	2.01 × 10 <sup>-4</sup>	0.0069	W
Perfluorodecane sulfonic acid ammonium salt	PFDS	1.52 × 10 <sup>-4</sup>	0.0092	W
Fluorotelomers				
6:2 Fluorotelomer alcohol	6:2FtOH	ND	ND	
8:2 Fluorotelomer alcohol	8:2FtOH	ND	ND	
8:2 Fluorotelomer acrylate	8:2FtOAc	ND	ND	

Note. ND, no significant displacement; L, less than 50% displacement.

<sup>a</sup>IC<sub>50</sub> is the competitor concentration that causes 50% displacement of [<sup>3</sup>H]-E2.

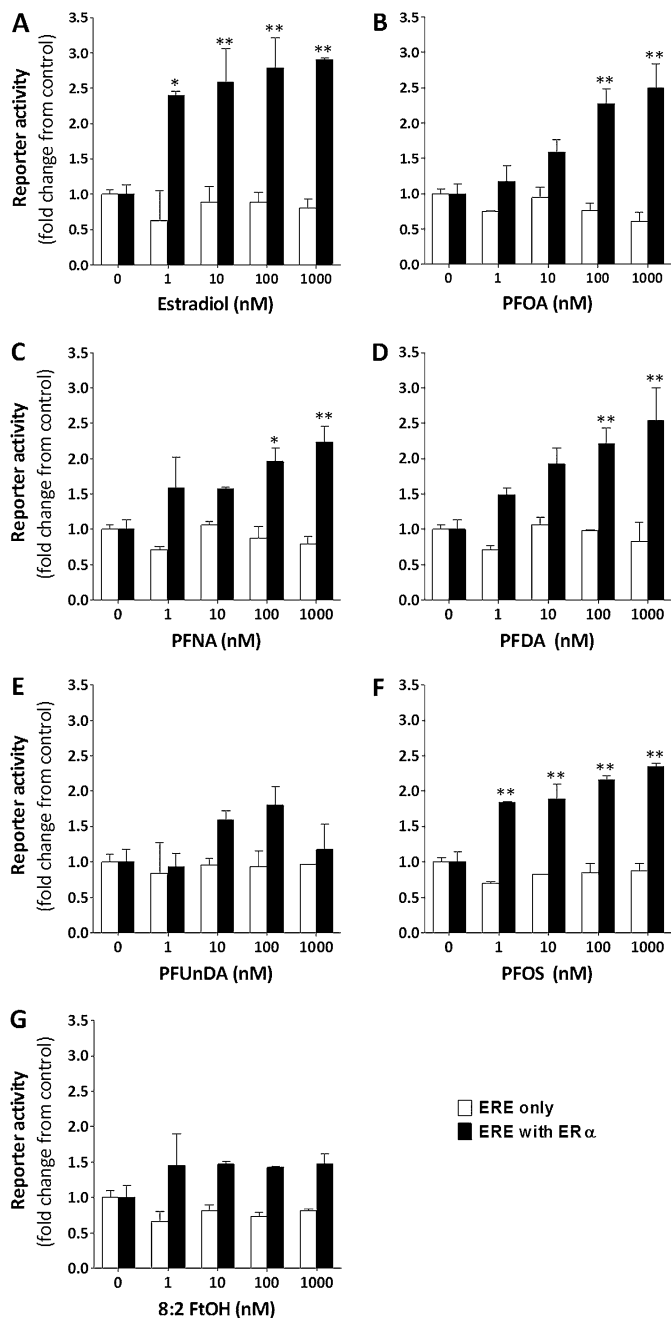
<sup>b</sup>RBA values were calculated by normalizing IC<sub>50</sub> values to that of E2 which was set at 100%.

<sup>c</sup>Letters indicate classification of RBA values into four broad groups indicating calculated affinity for trout liver ER as follows: S, strong binding (RBA > 10%); M, moderate binding (RBA 0.1–10%); W, weak binding (RBA 0.001–0.1%); and VW, very weak binding (RBA < 0.001%).

coordinate space as described in “Materials and Methods” section (Supplementary fig. 4). The root-mean-square deviation (RMSD) values for superimposition of the mER $\alpha$  and rtER $\alpha$ 1 homology models with the 3D crystal structure of hER $\alpha$  were 0.087 and 0.078 Å, respectively (SuperImpose Algorithm, ICM-Browser, Molsoft).

The natural ligand E2 docked within a 4-Å sphere around its 3D position in the crystal structure with favorable ICM scores of -38.68 (human), -37.75 (mouse), and -36.34 (trout) and a very low all atoms RMSD (< 0.151 Å for all three models) when compared with the crystallographic orientation. The hydrogen bond interactions observed in the crystal structure between the phenolic hydroxyl group of the A-ring and the hydroxyl group of the D-ring of the natural ligand and the side chains of residues Glu353/357/366 (hER $\alpha$ /mER $\alpha$ /rtER $\alpha$ 1) of helix 2, Arg394/398/407 of helix 3, and His524/528/537 of helix 8 were confirmed (Fig. 6). Sites of hydrophobic interactions with the natural ligand were similar to those previously identified (Marchand-Geneste *et al.*, 2006).

To explain the effects observed *in vitro* (radioligand-binding assay, cell-based assay) through ER $\alpha$  activation, PFHpA, PFOA, PFNA, PFDA, PFUnDA, PFOS, PFDS, 6:2FtOH, 8:2FtOH, and 8:2FtOAc were docked into the human, mouse, and trout models of the ER $\alpha$ -LBD binding pocket. In addition, DES (a strong ER $\alpha$  binder), GEN (a moderate binder), and NP and bisphenol A (BPA) (weak binders) were included in the docking study as reference compounds with known binding affinities toward the ER $\alpha$  (this study; Matthews *et al.*, 2000; Olsen *et al.*, 2005). The results of the molecular docking for all test ligands into the LBD of hER $\alpha$  are shown in Figure 7, mER $\alpha$  in Supplementary figure 5, and rtER $\alpha$ 1 in Supplementary figure 6. When considered by broad groupings (strong, moderate, weak, and very weak), the ranking by ICM score of the docked reference compounds correlated well with the difference in ER $\alpha$ -binding affinity and activity of the compounds (Table 3). The higher score for BPA in the mouse model is likely because of the third hydrogen bond interaction between the compound and the side chain of residue T351



**FIG. 5.** Transactivation of hER $\alpha$  by PFAAs. HEK-293T cells were transfected with (solid bar) or without (open bar) a human ER $\alpha$ -expressing plasmid, the XTEL luciferase reporter plasmid containing a consensus ERE sequence, and a  $\beta$ -galactosidase-containing plasmid for normalization of raw luminescence data. Twenty-four hours after transfection, cells were treated with increasing concentrations of E2 (A), PFOA (B), PFNA (C), PFDA (D), PFUnDA (E), PFOS (F), and 8:2FtOH (G). Values are the mean gene reporter activity expressed as fold change with respect to control (0nM treatment) ( $N = 3$  replicate experiments). Within the ERE + ER $\alpha$  data set, \* $p < 0.05$ , \*\* $p < 0.01$  as determined by one-way ANOVA with Bonferonni *post hoc* tests compared with 0nM.

(Supplementary fig. 5D). The moderate binder GEN and the weak binders BPA and NP docked into the human, mouse, and

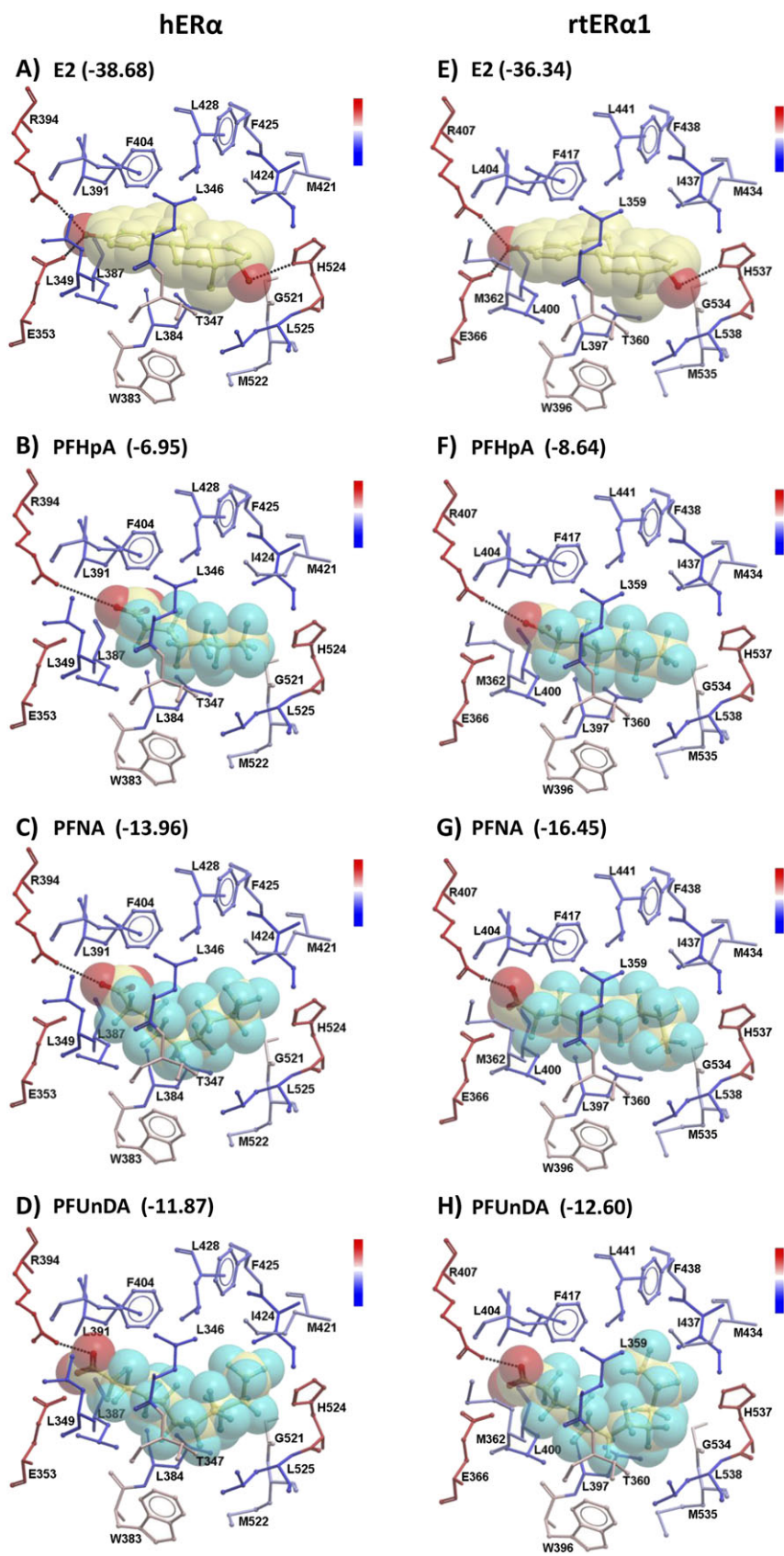
trout ER $\alpha$ -LBD binding pockets with lower ICM scores than E2 and DES because of loss of the classical triple hydrogen-bonding pattern observed with the two strongest binders (e.g., hER $\alpha$  in Figs. 7A–E). In the binding site, GEN establishes a triple hydrogen-bonding pattern like E2 and DES but with the carboxyl backbone of L387/391/400 instead; this bonding pattern is the likely cause for the improved docking score of GEN compared with BPA and NP, which form only two hydrogen bonds.

All PFAAs and fluorotelomer alcohol derivatives docked in all three ER $\alpha$  models with similar orientations. The negative charged oxygen of the carboxylic (PFHpA, PFOA, PFNA, PFDA, and PFUnDA) and the sulfonic (PFOS and PFDS) groups establishes a hydrogen bond interaction with the side chain of R394/398/407 but not with E353/357/366 (e.g., hER $\alpha$  in Figs. 7F–L). The loss of the classical hydrogen-bonding pattern present in E2, DES, and GEN docking poses is caused by the repulsion between the negative charge of the ligand (carboxylic and sulfonic groups) and the side chain of E353/357/366. Alternatively, 8:2FtOH and 6:2FtOH docked with higher scores (with the exception of 6:2FtOH in the mouse model) with the hydroxyl group interacting with both side chains of R394/398/407 and E353/357/366 (e.g., hER $\alpha$  in Figs. 7M and 7N). The fluorotelomer acrylate compound 8:2FtOAc displayed a similar hydrogen-bonding pattern as the carboxylic acids tested rather than the fluorotelomer alcohols, likely because of the positioning of the acrylate side chain near R394/398/407. Unlike several of the xenoestrogens studied, no hydrogen bonding was possible with H524/528/537 for the PFCs tested because of the orientation of the fluorinated carbon tail near this residue. The PFAA compounds with the best docking scores had fluorinated carbon chains of medium length, PFDA for hER $\alpha$  (–18.77) and PFNA for mER $\alpha$  and rER $\alpha$ 1 (–22.54 and –16.45, respectively), whereas shorter or longer compounds have lower scores (Table 3, Fig. 6). Space-filling models for the PFAAs showed that short carbon chain fluorinated chemicals (e.g., PFHpA) did not adequately fill the LBD pocket (Figs. 6B and 6F), whereas longer fluorinated carbon chain structures (e.g., PFUnDA) were nearly too large and required the fluorinated carbon tail to bend so that the ligand could fit into the binding pocket (Figs. 6D and 6H).

No major differences in the pattern of xenoestrogen and PFC docking to the LBD region of ER $\alpha$  were observed for the species studied, aside from the noted different site of hydrogen bonding for BPA in the mER $\alpha$  model. However, the ICM scores for PFAA docking to the mER $\alpha$  model were generally somewhat higher than those for the hER $\alpha$  or rER $\alpha$ 1 models (Table 3).

## DISCUSSION

This study details the first comprehensive screen of structurally diverse PFCs for estrogen-like activity in an animal model that mimics human insensitivity to peroxisome



proliferation, the presumed mechanism of action for PFAAs. As has been shown previously by our laboratory for PFOA (Tilton *et al.*, 2008), dietary exposure to PFAAs did not elicit a change in palmitoyl CoA oxidation or catalase activity, indicative of a peroxisome proliferation response. Rather, eight different PFAAs significantly increased expression of the estrogen biomarker protein Vtg, and perfluoroalkyl carboxylic acids were more estrogenic compared with the perfluoroalkyl sulfonates and fluorotelomers tested.

Additionally, we provide the first evidence that PFAAs interact directly with the ER to exert their estrogenic activity. Results from competitive steroid-binding assays show that multiple PFAAs effectively bind to the trout liver ER with a structure-activity relationship similar to that observed for *in vivo* Vtg induction in trout. Perfluoroalkyl carboxylic acids ranging in size from 7 to 13 carbons were identified as competitive binders for the rER, with the 10-carbon compound PFDA being the strongest binder of the chemical group. Although not considered strong ER agonists, these PFAAs had sufficient binding affinity for the ER that, in the absence of endogenous estrogen (as is the case in juvenile trout), the compounds are capable of eliciting a weak estrogen-like effect as measured by the Vtg bioassay. The rER is considered more promiscuous than in other species (Matthews *et al.*, 2000) and has been shown to weakly bind a wide variety of environmental xenobiotics; in this study, PFAA binding was comparable to some of these weak estrogens, such as nonylphenol and I3C. Interestingly, PFOS and PFDS showed the highest affinity of all the PFCs tested for this receptor, contradicting the fairly modest Vtg response observed for these sulfonate compounds *in vivo*. This discrepancy is not likely due to poor bioavailability of perfluoroalkyl sulfonates because PFOS is reported to have a four-fold greater half-life in trout than PFOA (Martin *et al.*, 2003). Currently, it is unknown why PFOS and PFDS are less estrogenic *in vivo* than would be predicted by their RBA for the trout ER. Moreover, results of steroid-binding assays for trout liver ER suggest that the observed difference in Vtg response between carboxylic acids and sulfonates is not likely the result of differing affinities for the ER. It should be noted, as well, that the ER-binding assay does not discriminate between the four trout ER isoforms, of which rER $\alpha$ 1 and rER $\beta$ 2 are highly expressed in trout liver (Nagler *et al.*, 2007). Finally, we observed that dietary PFOA, PFNA, PFDA, and PFUnDA induced a marked increase in Vtg expression *in vivo*, even though the binding affinity for these

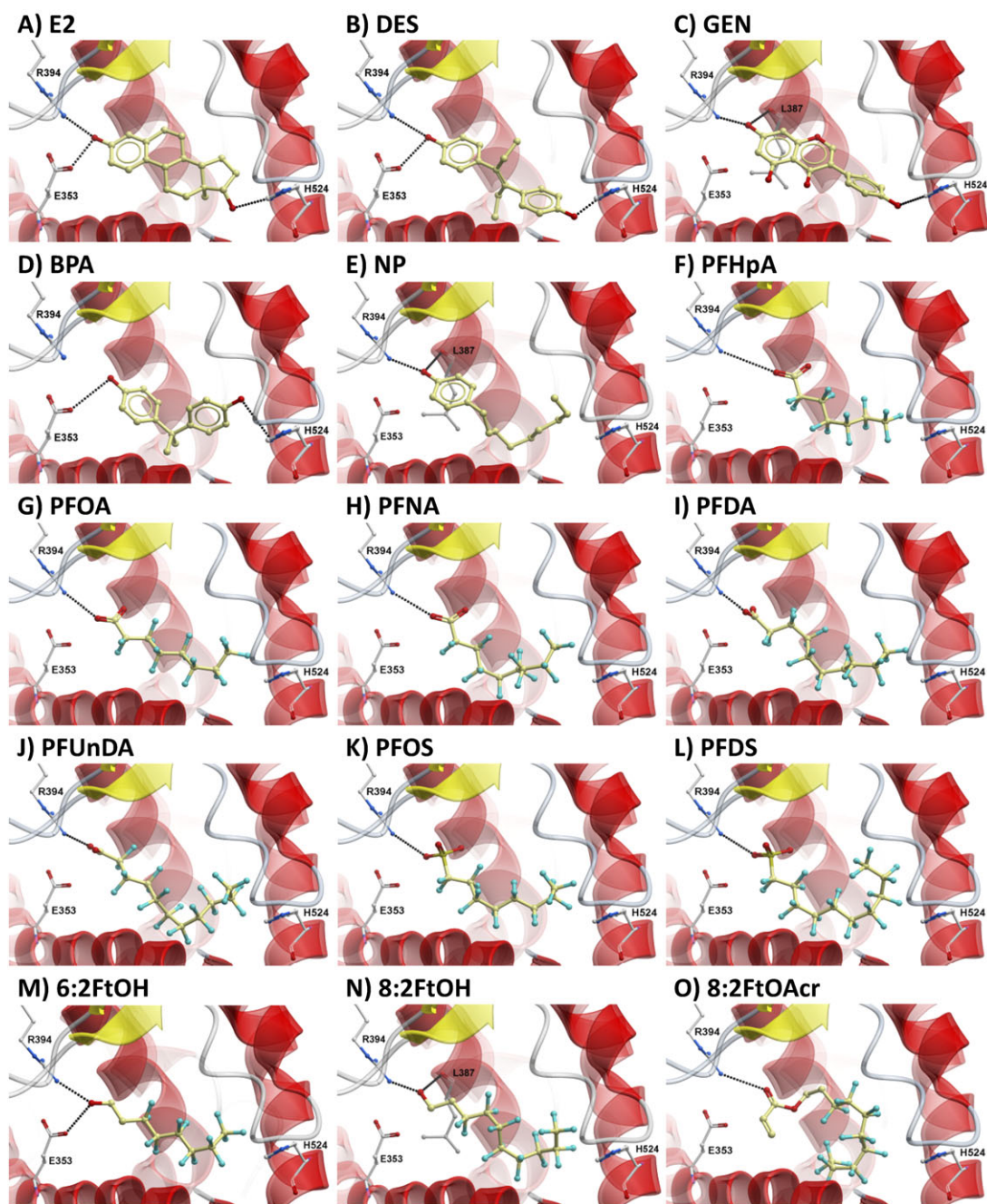
compounds was classified as weak or very weak. This biomarker assay integrates several factors that impact the biological response to PFCs, including chemical-specific patterns of assimilation, distribution, and elimination. The perceived discrepancy in the effectiveness of PFAAs *in vivo* versus *in vitro* could be explained by the apparent accumulation of these compounds in trout over time (14 days), which allowed for blood levels to achieve concentrations sufficient for PFAA binding to the trout ER and a robust Vtg response (Fig. 3).

In general, the fluorotelomers tested in this study were not potent estrogens *in vivo*, save for 6:2FtOH which caused a slight increase in Vtg expression. Similar to our observations in trout, Liu *et al.* (2007) showed that PFOA, PFOS, and 6:2FtOH significantly increased Vtg expression by tilapia hepatocytes *in vitro*, whereas 8:2FtOH was ineffective. However, subsequent reports showed that aqueous exposure to 6:2FtOH or 8:2FtOH caused significant induction of Vtg in male zebrafish, though this may be the result of changes in steroid metabolism rather than direct action of the chemicals at the ER (Liu *et al.*, 2009, 2010). On the other hand, Ishibashi *et al.* (2008) reported that waterborne 6:2FtOH and 8:2FtOH increased hepatic Vtg gene expression in male medaka and that this effect was likely linked to interaction with the medaka ER $\alpha$ . Interestingly, in this same study (Ishibashi *et al.*, 2008) as well as another report using human breast cancer cells (Maras *et al.*, 2006), the authors found no evidence for ER interaction with PFAAs. Overall, these differences suggest that the estrogenic response to PFAAs and fluorotelomers may depend on the route of exposure or the species studied.

This report also describes the first broad-scale dose-response study for estrogen-like activity of PFOA and PFDA, two PFAAs commonly detected in humans and wildlife. Overall, the dose-response data suggest that the compounds tested are weakly estrogenic, as fairly high diet levels were required to induce expression of the estrogen biomarker protein Vtg. Neither chemical caused a significant increase in Vtg expression at blood levels for PFOA or PFDA in trout that correspond to the levels of these compounds observed in the general human population, about 2–7 ppb (reviewed in Fromme *et al.*, 2009; Kannan *et al.*, 2004). However, a significant estrogenic response was observed at levels that may reflect certain human exposures, such as residents living near a contaminated site or occupationally exposed workers (Emmett *et al.*, 2006). It should be noted, however,

**FIG. 6.** Effect of fluorinated carbon chain length on perfluoroalkyl carboxylic acid docking orientation and efficiency for human or trout ER $\alpha$  proteins. The docked orientation of E2 is represented by the stick/space-filling model in the binding pocket of the receptor (A and E). Dockings of three representative perfluoroalkyl carboxylic acids with different fluorinated carbon chain length are shown, including PFHpA (seven carbons), PFNA (nine carbons), and PFUnDA (11 carbons), with hER $\alpha$  (B–D) or rER $\alpha$ 1 (F–H). Interacting residues of the LBDs are shown as sticks and are colored according to their hydrophobicity (inset scale) with hydrophobic residues colored blue and hydrophilic colored red. Hydrogen bonds are represented by black dotted lines between the donor (D) and the acceptor (A) and are defined as follows: Distance D–A: 2.8–3.2 Å; Angle D–H–A: 140–180°. Residue labels are numbered according to the indicated protein sequences, which are available in Supplementary figure 3. The corresponding ICM docking scores are indicated for each ligand-protein pair; scores for all PFAAs tested are available in Table 3.



hER $\alpha$ 

**FIG. 7.** *In silico* model showing docking of estrogens and select PFCs into the hER $\alpha$  LBD. Docking of test ligands into the hER $\alpha$  LBD was performed using Molsoft ICM, and the lowest ICM-scored poses for each calculated ligand-protein docking are shown. Ligands are colored by atom type (the carbon atoms in tan) and are displayed as ball and sticks. Relevant protein residues, including E353, R394, H524, and L387, are displayed as ball and sticks and colored by atom type: carbon atoms in gray, oxygen in red, sulfur in yellow, and fluorine in cyan. Hydrogen bonds are represented by black dotted lines between the donor (D) and the acceptor (A) and are defined as follows: Distance D–A: 2.8–3.2 Å; Angle D–H–A: 140–180°.

that the *in vivo* studies described above are short-term (14 day) experiments that do not precisely reflect the typical chronic exposure to these chemicals observed in humans. Moreover, humans and wildlife have detectable blood levels of multiple PFCs (Calafat *et al.*, 2007), which could have additive

or synergistic toxic effects depending on the mechanism of action. In this study, results of a simple *in vivo* mixture experiment with PFOA, PFNA, PFDA, and PFUnDA suggest that perfluoroalkyl carboxylic acids impact Vtg expression in at least an additive manner at higher exposures.

TABLE 3  
Molecular Modeling of Ligand Interaction with Human, Mouse, and Trout ER $\alpha$  LBDs

Ligand	ICM docking score			Hydrogen-bonding interactions			Classification <sup>a</sup>
	hER $\alpha$	mER $\alpha$	rtER $\alpha$ 1	HER $\alpha$	mER $\alpha$	rtER $\alpha$	
E2	-38.68	-37.75	-36.34	R394, E353, H524	R398, E357, H528	R407, E366, H537	S
DES	-38.34	-37.40	-37.25	R394, E353, H524	R398, E357, H528	R407, E366, H537	S
GEN	-30.10	-32.70	-31.63	R394, L387, H524	R398, L391, H528	R407, L400, H537	M
NP	-23.41	-24.49	-23.39	R394, L387	R398, L391	R407, L400	W
BPA	-18.77	-27.86	-17.34	E353, H524	R398, E357, T351	E366	M,W
PFHpA	-6.95	-9.63	-8.64	R394	R398, E357	R407	VW
PFOA	-11.84	-17.75	-11.70	R394	R398, L391	R407	VW
PFNA	-13.96	-22.54	-16.45	R394	R398	R407	VW,W
PFDA	-18.77	-20.05	-13.41	R394	R398	R407	VW,W
PFUnDA	-11.87	-17.85	-12.60	R394	R398	R407	VW,W
PFOS	-15.77	-19.06	-10.96	R394	R398	R407	VW,W
PFDS	-11.52	-14.79	-11.12	R394	R398	R407	VW
6:2FtOH	-20.48	-20.46	-20.30	R394, E353	R398	R407, E366	W
8:2FtOH	-18.57	-25.68	-18.06	R394, L387	R398	R407, L400	W
8:2FtOAc	-15.60	-14.50	-17.06	R394	R398	R407	W

Note. ICM scores are a measure of fit within the receptor-binding pocket, taking into account continuum and discreet electrostatics, hydrophobicity, and entropy parameters. The predicted amino acid residue locations for hydrogen bonding are indicated for each ligand-protein docking.

<sup>a</sup>Letters indicate classification of ICM docking scores into four broad groups indicating predicted affinity for ER $\alpha$  as follows: S, strong affinity (ICM scores > -35); M, moderate affinity (scores -25 and -35); W, weak affinity (scores -15 to -25); and VW, very weak affinity (scores > -15). Two classifications are given when scores for different receptors fit within multiple ranges.

Subsequent experiments sought to expand the findings in trout to humans using a reporter gene assay for hER $\alpha$ . We determined that certain PFAAs activate hER $\alpha$ -induced transcription, including the perfluoroalkyl carboxylic acids PFOA, PFNA, and PFDA and one sulfonate compound, PFOS, whereas 8:2FtOH was ineffective. Although our observations for PFC activation of hER $\alpha$  largely match those for ER binding and *in vivo* activity in trout, they follow a pattern opposite to that observed by Ishibashi *et al.* (2007); results of their studies using a yeast two-hybrid assay suggested that fluorotelomer alcohols, but not PFOA or PFOS, interacted with hER $\alpha$ . The apparent differences in observations of these two research groups might be explained by interassay variation associated with the cellular and molecular responses in the bioassays employed; one may reasonably expect that our reporter gene assay better represents the molecular context for ER activation in human cells. The estrogen-like activity of some fluorotelomers could be the result of metabolic production of polyfluorinated derivatives, such as 8:2 or 7:3 fluorotelomer acids (Nabb *et al.*, 2007), which have chemical structures and lengths similar to the estrogenic perfluoroalkyl carboxylic acids identified in this study. Moreover, substantial differences in the metabolism of fluorotelomers in human, rodent, and trout hepatocytes have been reported (Nabb *et al.*, 2007).

A major focus of the present study was to understand the potential molecular interactions of PFAAs with the LBD region of the ER, particularly because the chemical structures of PFAAs previously identified as estrogenic have little in common with the structure of the native hormone or other

environmental xenoestrogens. To this end, we employed an *in silico* molecular docking model to investigate docking of these putative ligands with hER $\alpha$ , mER $\alpha$ , and rtER $\alpha$ 1. From the results obtained and after considering the score ranking of the reference compounds E2, DES, GEN, BPA, and NP, the PFAAs and fluorotelomers examined were predicted to be *in vitro* weak binders and ER $\alpha$  activators. For the three species investigated, the overall pattern of xenoestrogen and PFC docking to the LBD region of ER $\alpha$  was fairly consistent, as all compounds docked into each model with similar orientations. Hydrogen bonding with the R394/398/407 (human/mouse/rainbow trout) residue was critical for all putative ligands, whereas more the estrogen-like compounds DES and GEN docked with the classic triple hydrogen bond pattern, with additional hydrogen bonds predicted at residues E353/357/366 and H524/528/537. For the PFAAs examined, docking scores for mER $\alpha$  were moderately more favorable than those for the human and trout receptors. However, it is uncertain whether this modest difference in computed affinity for mER $\alpha$  would translate into greater *in vivo* activity for PFAAs in mice, especially because these chemicals are reported to have quite different pharmacokinetic properties in rodents, humans, and trout (Kudo and Kawashima, 2003; Martin *et al.*, 2003).

The homology model developed for rtER $\alpha$ 1 agreed well with a similar computational model described by Marchand-Geneste *et al.* (2006), where the natural ligand E2 formed hydrogen bonds with active site residues E366, R407, and H537. Our docking model appropriately predicted PFAA interaction with the trout ER and *in vivo* estrogen activity, as the medium-sized compounds

PFNA and PFDA docked to the rER $\alpha$ 1 with the most favorable ICM scores of the carboxylic acids examined. However, the model was not predictive for fluorotelomer alcohols. Despite the higher computed docking scores for all the fluorotelomers examined *in silico*, none of these compounds showed activity *in vitro*, and only 6:2FtOH was slightly estrogenic *in vivo* in trout. Moreover, the computational models presented here do not perfectly align with prior studies predicting positive interaction of fluorotelomers, but not PFOA or PFOS, with human or medaka ER $\alpha$  (Ishibashi *et al.*, 2007, 2008). *In silico* modeling is a useful tool for predicting potential ligand-receptor interactions or, in the case of this study, a retrospective tool used to gain insight into ligand-receptor interaction. However, these computational models are not perfect predictors of *in vivo* or *in vitro* interactions. Other factors, such as metabolism, bioavailability, or other biophysical parameters, could account for prediction errors in translating results from computational modeling to *in vivo* or *in vitro* experimental systems.

In this study, we investigated molecular interactions with ER $\alpha$  isoforms only, primarily because of its high expression in the liver (Flouriot *et al.*, 1998), which is generally considered the primary target organ for PFC toxicity. Because the LBDs for ER $\alpha$  and ER $\beta$  isoforms are highly similar, it is reasonable to predict that *in silico* modeling with the ER $\beta$  crystal structure would generate similar docking results as described above for ER $\alpha$  (Supplementary fig. 7).

In conclusion, we present here the first comprehensive evaluation of PFAAs for estrogenic activity *in vivo* using an animal model that mimics human insensitivity to PPs. Multiple estrogenic PFAAs have been identified, and a structure-activity relationship was observed, where a fluorinated carbon chain length of 8–10 carbons and a carboxylic acid end group were optimal for maximal induction of plasma biomarker proteins. Perfluoroalkyl carboxylic acids, in particular PFOA, PFNA, PFDA, and PFUnDA, were all potent inducers of Vtg, although at fairly high dietary exposures. Moreover, this study provides the first evidence for direct interaction of PFAAs with the ER, including the persistent environmental pollutants PFOA and PFOS. As with the *in vivo* Vtg assay, a structure-activity relationship between ER interaction and fluorinated carbon chain length was evident, with medium-sized PFAAs having the highest binding affinity and effective concentration values and most favorable docking scores for hER $\alpha$ , mER $\alpha$ , and/or rER $\alpha$ 1. Collectively, these findings support the idea that several PFAAs could act as weak environmental xenoestrogens.

#### SUPPLEMENTARY DATA

Supplementary data are available online at <http://www.toxsci.oxfordjournals.org>.

*Supplementary materials and methods.pdf* includes experimental details for measurement of PFOA and PFDA in trout blood plasma by LC-MS/MS. *Supplementary tables.pdf* contains nine additional tables, including a list of PFCs

investigated in this report (Suppl. Table 1) and morphometric data from the *in vivo* experiments described above (Suppl. Tables 2 and 3). Supplementary Figures.pdf contains seven additional figures, including activity of peroxisome proliferator-responsive enzymes in trout liver (Suppl. Fig. 1), trout ER saturation analyses (Suppl. Fig 2), alignment of human, mouse and fish ERs (Suppl. Figs 3 and 7), the mouse and trout ER homology models (Suppl. Fig 4) and results of *in silico* docking of PFCs into the mouse and trout ER $\alpha$  LBDs (Suppl. Figs. 5 and 6).

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