A Computational-Based Approach to Identify Estrogen Receptor α/β Heterodimer Selective Ligands[§]

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ABSTRACT

The biologic effects of estrogens are transduced by two estrogen receptors (ERs), ER α and ER β , which function in dimer forms. The ER α/α homodimer promotes and the ER β/β inhibits estrogen-dependent growth of mammary epithelial cells; the functions of ER α/β heterodimers remain elusive. Using compounds that promote ER α/β heterodimerization, we have previously shown that ER α/β heterodimers appeared to inhibit tumor cell growth and migration in vitro. Further dissection of ER α/β heterodimer functions was hampered by the lack of ER α/β heterodimer-specific ligands. Herein, we report a multistep workflow to identify the selective ER α/β heterodimer-inducing compound. Phytoestrogenic compounds were first screened for

ER transcriptional activity using reporter assays and ER dimerization preference using a bioluminescence resonance energy transfer assay. The top hits were subjected to in silico modeling to identify the pharmacophore that confers ER α/β heterodimer specificity. The pharmacophore encompassing seven features that are potentially important for the formation of the ER α/β heterodimer was retrieved and subsequently used for virtual screening of large chemical libraries. Four chemical compounds were identified that selectively induce ER α/β heterodimers over their respective homodimers. Such ligands will become unique tools to reveal the functional insights of ER α/β heterodimers.

Introduction

The biological effects of estrogenic compounds are mediated by two estrogen receptors (ERs), namely ER α and ER β . These receptors are expressed in a cell-type and tissue-specific manner; however, they can also colocalize within the same cell, and their presence varies based on different disease states (Leygue et al., 1998; Lau et al., 1999; Weihua et al., 2003; Powell et al., 2012; Nilsson and Gustafsson, 2013). Both ERs share a conserved nuclear receptor domain structure that encompasses a DNA binding domain, ligand-binding domain (LBD), a central hinge region, and two activation functional domains. The ligand binding to ER α or ER β induces a conformational change that leads to receptor dimerization,

where either homodimers (ER α/α or ER β/β) or heterodimers (ER α/β) can be formed.

The existence of the ER α/β heterodimer was first described 20 years ago using in vitro translated receptors and an estrogen response element (ERE) in a gel shift assay. Cowley et al. (1997) showed that ER heterodimers could bind to a consensus ERE and recruit coactivators in vitro. Similar observations were made by others (Pace et al., 1997; Tremblay et al., 1999). Pettersson et al. (1997) showed direct interaction between ER β and ER α in a glutathione *S*-transferase pull-down assay and binding of the heterodimer to DNA. Two dimerization domains were mapped to the DNA binding domain and LBD (Brzozowski et al., 1997; Pace et al., 1997). ER heterodimers were shown to form in a ligand-dependent and -independent manner in vitro (Pace et al., 1997). Recent technical advances confirmed the formation of the ER α/β heterodimer in vivo. Our laboratory developed a bioluminescence resonance energy transfer (BRET) assay to monitor ER dimerization in live cells (Powell and Xu, 2008). BRET assays revealed that the types of ER dimer pair being formed depend on the chemical characteristics of the ligand and its concentration (Powell and Xu, 2008). Moreover, ER α/β heterodimers have been detected in vivo using molecular imaging

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ABBREVIATIONS: BRET, bioluminescence resonance energy transfer; ChIP, chromatin immunoprecipitation; DMSO, dimethylsulfoxide; 3D, three-dimensional; E2, 17 β -estradiol; ER, estrogen receptor; ERE, estrogen response element; FBS, fetal bovine serum; GALAHAD, genetic algorithm with linear assignment of hypermolecular alignment of database; LBD, ligand-binding domain; RLuc, Renilla luciferase; YFP, yellow fluorescent protein.

techniques (Paulmurugan et al., 2011) and in breast cancer tissues using proximity ligation assay (Iwabuchi et al., 2017). Evidence also shows that the ER α /ER β heterodimer is transcriptionally active and may regulate a distinct set of genes from their respective homodimers (Tremblay et al., 1999).

In contrast to the established role that the ER α /ER α homodimer is a driver of estrogen-mediated cellular proliferation and ER β /ER β homodimers elicit an antiproliferative and proapoptotic effect, the function of ER α /ER β heterodimers in the biologic processes is the least understood. Unlike the ER α /ER α and ER β /ER β homodimers, where subtype-specific ligands for ER α and ER β aided in elucidating their function (Lindberg et al., 2003; Weihua et al., 2003), ligands that specifically induce ER α /ER β heterodimers have not been identified, largely due to the absence of a full-length crystal structure for the ER α /ER β heterodimer.

Indirect evidence suggesting that ER α /ER β heterodimers might have an antiproliferative role in breast cancer cells have previously been reported (Hall and McDonnell, 1999; Powell et al., 2012). Endoxifen, the primary metabolite of tamoxifen with growth inhibitory effects, stabilizes ER β and induces the formation of ER α /ER β heterodimers in cells expressing both ERs (Wu et al., 2011). Furthermore, high-throughput BRET assays identified a phytoestrogen (i.e., cosmosiin) that favors ER α /ER β heterodimer formation (Powell et al., 2012). This ER α /ER β heterodimer-inducing compound elicited antiproliferative effects in prostate and breast cancer cells. Although cosmosiin induces the formation of ER α /ER β heterodimers but not the pro-proliferative ER α /ER α homodimers, it is only effective at high concentrations (e.g., 10 μ M) and also slightly induces ER β /ER β homodimers (Powell et al., 2012). More potent and selective ER α /ER β heterodimer-inducing ligands are needed to elucidate the biologic functions of heterodimers.

Herein, we describe a multistep screening strategy (i.e., cell-based assays and in silico modeling) to identify ER α /ER β heterodimer-selective ligands. Reporter assays and BRET assays were employed to screen a small library of flavonoid-type phytoestrogenic compounds, from which a pharmacophore model was generated using the SYBYL GALAHAD (i.e., genetic algorithm with linear assignment of hypermolecular alignment of database) program (Tripos, St. Louis, MO). The pharmacophore model was subsequently used in a three-dimensional (3D) search query of two commercial chemical databases to identify new active structures. Four compounds were identified from the in silico screen that selectively induce ER α /ER β heterodimers. We showed that the representative compounds induce expression of putative ER α /ER β target genes by corecruiting ER α and ER β to the target gene promoter. Such ER α /ER β -selective compounds will be exploited to determine the biologic functions of ER α /ER β heterodimers, their downstream effectors, and target genes.

Materials and Methods

Cell Culture and Chemicals. Cell culture media were obtained from Invitrogen (Carlsbad, CA). HEK293 cells were maintained in Dulbecco's modified Eagle's medium supplemented with 10% Gibco fetal bovine serum (FBS) (Invitrogen) at 37°C and 5% CO₂. T47D-KBLuc cells were routinely maintained in RPMI 1640 medium supplemented with 10% FBS and 100 U/ml (1%) penicillin/streptomycin. Experiments were conducted in phenol-free media and dextran charcoal stripped FBS purchased from Hyclone (Logan, UT).

Compounds were dissolved in 100% dimethylsulfoxide (DMSO) and finally diluted in culture medium prior to the assay. 17 β -estradiol (E2) and ICI 182,780 were purchased from Sigma-Aldrich (St. Louis, MO). Thirty-one compounds used in our initial screening were a gift from the Lim Laboratory and have been previously described (Hyun et al., 2010; Hwang et al., 2011; Shin et al., 2011). They were chosen for screening based on their structural similarity to the lead compound cosmosiin, which was previously identified to induce ER α /ER β heterodimerization (Powell et al., 2012). Test compounds were purchased from ChemBridge (<http://www.chembridge.com>) and Maybridge (<http://www.maybridge.com>).

BRET Assays. Dimerization of ERs was measured by BRET assays as previously described in Powell and Xu (2008). Briefly, HEK293 cells were transfected with either a single BRET fusion plasmid (pCMX-ER α -RLuc or pCMX-RLuc-ER β) or cotransfected with Renilla luciferase (RLuc) and yellow fluorescent protein (YFP) BRET fusions (pCMX-ER α -RLuc + pCMX-YFP-ER β for ER α /ER β heterodimers; pCMX-ER α -RLuc + pCMX-ER α -YFP for ER α homodimers; or pCMX-RLuc-ER β + pCMX-YFP-ER β for ER β homodimers). Twenty-four hours post-transfection, cells were trypsinized and plated in a 96-well white-bottom microplate and incubated with ligands for 1 hour. Coelenterazine h (Promega, Madison, WI) was added in phosphate-buffered saline at a final concentration of 5 μ M, and 460 and 530 nm emission detection measurements were immediately taken at 0.1 second/wavelength read/well on a PerkinElmer Victor 3-V plate reader (Akron, OH). Similar assays were done using E2-binding defective mutants of the LBDs of ER α and ER β , ER α G521R-RLuc, and YFP-ER β G491R. Each compound was an independent experiment tested in a dose response with three biologic replicates per dose. For each condition (ER α /ER α , ER β /ER β , and ER α /ER β), two-way analysis of variance with random effect was conducted to obtain *P* values for each comparison of the individual compounds with DMSO controls. Then, these *P* values were adjusted by multiple comparisons analysis to control false discovery rate less than 0.05.

ER Luciferase Reporter Assays Using T47D-KBLuc Cells. T47D-KBLuc is a well-characterized cell line for the screening of estrogenic compounds (Wilson et al., 2004). These cells express both ER α and ER β and have been stably transfected with pGL2.TATA.luc.neo, which contains three estrogen responsive elements upstream of a luciferase reporter gene. Cells were seeded in 96-well plates at an initial concentration of 1×10^4 cells/well in RPMI 1640 phenol-free medium supplemented with 10% charcoal stripped FBS for 24 hours in 5% CO₂ atmosphere at 37°C. Cells were allowed to attach overnight and media were removed and replaced with media containing 10 μ M compound. Then, 10 nM E2 and 1% DMSO were used as positive and negative controls, respectively. The potent ER antagonist ICI 182,780 was used for counter-screen to determine ER specificity. Cells were incubated with compound for 18–24 hours at 37°C in 5% CO₂. Following incubation with compounds, luciferase was measured using the Bright-Glo Luciferase Assay System (Promega) on a PerkinElmer Victor 3-V plate reader. Luciferase activity was normalized according to protein concentration. Values were expressed as fold change over DMSO (mean value of induction as a multiple of the value of vehicle controls) and error bars represent S.D.

Quantitative Real-Time Polymerase Chain Reaction. Total RNA was extracted from the cells using Trizol reagent (Invitrogen). The first-strand cDNA was synthesized using the RevertAid First Strand cDNA Synthesis Kit (Thermo, Waltham, MA) according to the manufacturer's instructions. Quantitative polymerase chain reaction was conducted using SYBR Green dye (Roche Scientific, Basel, Switzerland) and a CFX96 instrument (BioRad, Hercules, CA). The following primer sequences (IDT, Coralville, IA) were used in this study:

BAG1-qRT-F: GCCCAAGGATTGCAAGCTG and
BAG1-qRT-R: CTGTGTCACTCGGCTAGG;
ATP6V0E1-qRT-F: CCTCACTGTGCCTCTCATTGT and
ATP6V0E1-qRT-R: AGCAAAGTGAACAGGTCACCA;
BAG-ChIP-F: AGGAAGCTCTGATAGAAGGCAG and
BAG-ChIP-R: AGAACAGTCCACAGGAAGAGGT; and
ATP6V0E1-ChIP-F: CCCCTGGCAGTTTCGTCAC and
ATP6V0E1-ChIP-R: TCTTGTTCATAATTTGACTTTGGAG.

Chromatin Immunoprecipitation (ChIP). Flag-tagged ER β was stably expressed in MCF7 cells by retroviral induction. MCF7-ER β cells were cultured in a 10-cm dish and crosslinked with 1% formaldehyde for 10 minutes at room temperature. Crosslink was quenched for 5 minutes at room temperature by the addition of glycine to a final concentration of 0.125 M. Anti-Flag antibody (Sigma-Aldrich) and anti-ER α (HC-20; Santa Cruz, Dallas, TX) were used for ChIP assays. ChIP assays were performed as described previously (Zeng and Xu, 2015; Zeng et al., 2016). The experiment was done in triplicate samples of biologic replicates. Statistical testing was performed using unpaired two-tailed Student's *t* test analysis. Experiments were repeated at least twice. A value of *P* < 0.05 was considered statistically significant.

Fluorescence Polarization Competition Ligand-Binding Assays. The binding affinity of ligands for ER α and ER β were measured using the PolarScreen ER Competitive Binding Assay Kit (Invitrogen). Purified ER α and ER β (30 and 20 nM), were incubated with serial dilutions of test compounds (1 mM to 10 nM) and fluorescein-labeled E2. Fluorescence polarization was measured using a Victor $\times$ 5 microplate reader (PerkinElmer). Approximate IC₅₀ values were determined by GraphPad Prism Software (GraphPad Software Inc., La Jolla, CA) from competitive binding curves.

Preparation of the Initial Ligands. All computational studies were done using the SYBYL molecular modeling package (Tripos) in a Stereo 3D Dell T5500 molecular graphics computer (Intel dual quad, Nvidia FX 4800 graphics). All of the structures used were built using the sketch module, energy minimized and prepared using SYBYL's ligand preparation module. The quick 3D parameter was used where 3D coordinates were generated and charges were neutralized.

Generation of Ligand-Based Pharmacophore Hypothesis and Virtual Screening. Pharmacophore hypothesis was generated using the GALAHAD module of the SYBYL software suite. There were seven compounds in the training set to generate the pharmacophore hypothesis. The GALAHAD module was run for 100 generations with a population size of 45, at least five molecules were required to hit for the program to consider it a pharmacophoric feature. Default values were used for all other settings. Between all of the models, the one with the best energy, sterics, and pharmacoric similarity values based on Pareto ranking was selected as the best model. For 3D virtual screening, the generated pharmacophore hypothesis model was converted into a 3D search query using the UNITY-3D module.

3D Virtual Screening of Two Commercial Databases. The selected pharmacophore model was validated and converted into a UNITY query for pharmacophore-guided virtual screening studies. The query was then used for screening two commercial chemical databases, Maybridge (<http://www.maybridge.com>) and Chembridge (<http://chembridge.com>), which were obtained from the ZINC public database. A flexible 3D search was executed and no filters or restrictions were applied. The UNITY module uses a conformationally flexible 3D-searching algorithm to result in rapid identification of molecules that match with the given pharmacophore. Compounds that had their chemical groups spatially overlap with the features of the pharmacophore model were captured as hits. Subsequent hits were then confirmed to match all seven key pharmacophoric features by visual analysis. Hits were then ranked by Qfit and by the integrated ranking features in SYBYL.

Results

Identification of ER Agonists Using the T47D-KBluc Reporter Cell Line. The T47D-KBluc breast cancer cell line is a well-characterized cell line that has an ERE-driven luciferase reporter stably integrated. It is considered a versatile cell system for screening estrogenic compounds because it expresses both ER α and ER β (Wilson et al., 2004). Using this cell line, 37 flavonoids in the subclass chemical compound library (Table 1) were screened due to their

structural similarity to cosmosiin, a previously identified ER α / β heterodimer-inducing compound (Powell et al., 2012). All compounds were tested at one final concentration of 10 μ M, because at this concentration even weak estrogenic compounds are able to activate ER transcriptional activity in T47D-KBLuc (Powell et al., 2012). With a 2-fold cutoff, 13 out of 37 compounds from three out of the four subclasses were identified as hits (Fig. 1; Supplemental Fig. 1). Seven compounds in the flavone subset were identified as hits. Compared with the DMSO control, compounds **3**, **5**, **6**, and **7** showed moderate activation, compounds **15** and **16** elicited almost 10-fold induction, and compound **17** elicited nearly 20-fold induction of ERE reporter. Three compounds (**18**, **23**, and **24**) in the flavanone subset and two compounds (**26** and **29**) in the isoflavone subset were retained as hits. Compound **28** is genistein, an isoflavone known to be an ER agonist that has been shown to induce all three ER dimer pairs (Powell and Xu, 2008). The 12 compounds were then subjected to a counter-screen in the presence of ER antagonist ICI 182,780. Cotreatment of ICI 182,780 completely ablated ERE-luciferase activation, demonstrating that their transcriptional response is ER dependent (Supplemental Fig. 2).

In vitro fluorescence polarization assays were used to determine the relative binding affinities of the phytochemicals that activated the ER in reporter assays. Fluorescence polarization assays are a competitive ligand-binding assay that measures the replacement of a fluorescein-labeled E2 by unliganded compounds from the LBDs of ER α and ER β . The dose-response curves for representative ligands and the relative binding affinities are shown in Fig. 1B with half-maximal inhibitory concentration (IC₅₀) ranging from 1.45 to 721 μ M.

Ligand-Induced ER Dimerization Measured by BRET Assay. The 12 compounds were subsequently tested for their ability to induce ER α / α , ER β / β , and ER α / β ER dimers in the BRET assay (Fig. 2, A–C). Of the flavones, compounds **3**, **5**, **6**, and **7** induced dimerization of ER α / α homodimers and ER α / β heterodimers at 10 μ M. Compound **15** induced all three dimer pairs at 1 μ M. Compound **16** induced ER α / α and ER β / β at 1 μ M and all dimer pairs at 10 μ M. Compound **17** induced ER β / β and ER α / β dimerization while restricting the induction of ER α / α homodimers at 1 μ M, but only induced the formation of ER α / β at 10 μ M (Fig. 2, A–C). From the flavanones, ER α / β heterodimers were selectively induced by compounds **23** and **24** at 10 μ M, whereas compound **18** induced ER β / β and ER α / β dimerization at 1 μ M (Fig. 2, A–C). Of the isoflavone subclass (Fig. 2, A–C), compound **26** induced ER α / β and ER β / β dimerization at 10 μ M, whereas compound **29** induced ER α / β and ER β / β dimerization at 1 μ M.

To determine if newly identified compounds from the reporter and BRET assays indeed activate ER target gene expression, we measured mRNA levels of two ER target genes, BAG1 and ATP6V0E1, after compound 29 treatment, using compound **28** (genistein) as a positive control. These compounds were selected since they exhibited the highest activities in the reporter assay (Fig. 1A). Because most breast cancer cell lines do not express ER β , we constructed MCF7 cells that stably express Flag-tagged ER β (Fig. 2D). Treatment of Flag-ER β MCF7 cells with compounds **28** and **29** significantly increased the mRNA levels of BAG1 and ATP6V0E1 compared with the DMSO control (Fig. 2, E and F). BAG1 has been implicated to be an ER β / β -specific target

TABLE 1
Core structures and names of flavonoid compounds used in this study

Thirty-seven flavonoid compounds from four different subclasses: flavones, flavanones, isoflavones, and chalcones were previously tested.

Compound	Number	Nomenclature	R1	R2	R3	Molecular Weight
 Flavone	1	Flavone	H	H	H	222.24
	2	5-Hydroxyflavone	H	5-OH	H	238.24
	3	2',3'-Dihydroxyflavone	H	H	2',3'-Di-OH	254.24
	4	2',3'-Dimethoxyflavone	H	H	2',3'-Di-OCH3	282.29
	5	4'-Hydroxy-3'-methoxyflavone	H	H	4'-OH-3'-OCH3	268.26
	6	3',5'-Dihydroxyflavone	H	5-OH	3'-OH	254.24
	7	4'-Hydroxy-5-methoxyflavone	H	5-OCH3	3'-OH	268.26
	8	5-Methoxyflavone	H	5-OCH3	H	252.26
	9	4',5,7-Trimethoxyflavone	H	5,7-Di-OCH3	4'-OCH3	312.32
	10	8-Carboxyl-3-methoxyflavone	CH3	8-COOH	H	280.27
	11	5,6-Benzoflavone	H	5,6-Benzo	H	272.3
	12	7,8-Benzoflavone	H	7,8-Benzo	H	272.3
	13	2'-Methoxy- α -naphthoflavone	H	7,8-Benzo	2'-OCH3	302.32
	14	3,3',4',5',7-Pentahydroxyflavone-8-O-glucoside (Gossypin)	OH	5,7-Di-OH-8-O-Glucoside	3',4'-Di-OH	480.38
 Flavanone	15	3,4',5,7-Tetrahydroxyflavone (Kaempferol)	OH	5,7-Di-OH	4'-OH	286.24
	16	5,7-Dihydroxyflavone (Luteolin)	H	5,7-Di-OH	H	286.24
	17	3,7-Dihydroxyflavone	H	3,7-Di-OH	H	254.24
	18	4',5,7'-Trihydroxyflavanone (Naringenin)	H	5,7-Di-OH	4'-OH	272.25
	19	5-Methoxyflavanone	H	5-OCH3	H	254.28
	20	5,7-Dimethoxyflavanone	H	5,7-OCH3	H	284.31
	21	3,4',5',7'-Tetramethoxyflavanone	H	7-OCH3	3',4',5'-Tri-OCH3	344.36
	22	Flavanone	H	H	H	224.08
	23	7-Hydroxyflavanone	H	7-OH	H	240.25
	24	4-Hydroxyflavanone	H	4-OH	H	254.24
 Isoflavone	25	7-Methoxyisoflavone	H	7-OCH3	H	252.26
	26	4',6,7-Trihydroxyisoflavone (Demethyltaxasin)	H	6,7-OH	4'-OH	270.24
	27	4'-Hydroxy-6-methoxyisoflavone-7-D-glucoside (Glycitin)	H	6-OCH3-7-D-Glucoside	4'-OH	446.4
	28	5,7,4'-Trihydroxyisoflavone (Genistein)	H	5,7-Di-OH	4'-OH	270.24
	29	5,7-Dihydroxy-4'-methoxyisoflavone (Biochanin A)	H	5,7-Di-OH	4'-OCH3	284.26
	30	5,7,3',4'-Tetramethoxyisoflavone (Orobol)	H	5,7-Di-OCH3	3',4'-Di-OCH3	342.34
	31	2',3'',5'',7'-Trimethoxychalcone	—	2'-OCH3	3'',5'',7'-Di-OCH3	298.33
 Chalcone	32	3',3'',5'',7'-Trimethoxychalcone	—	3'-OCH3	3'',5'',7'-Di-OCH3	298.33
	33	3',3'',5'',7'-Trimethoxychalcone	—	3'-OCH3	3'',5'',7'-Di-OCH3	298.33
	34	2'',3'',5'',7'-Trimethoxychalcone	—	3'5'-Di-OCH3	2'OCH3	314.33
	35	3',3'',5'',7'-Tetramethoxychalcone	—	3'5'-Di-OCH3	3'',5'',7'-Di-OCH3	328.36
	36	2'-Hydroxy-4',4'-dimethoxychalcone	—	2'-OH-4'-OCH3	4'-OCH3	284.31
	37	2'-Hydroxy-3'',4',5''-trimethoxychalcone	—	2'-OH-4'-OCH3	3'',5'',7'-Di-OCH3	298.33

—, no chemical group considered.

Fig. 1. Transcriptional and ligand-binding assays of 37 flavonoid compounds. (A) T47D-KBLuc transcriptional assays showing ERE-luciferase reporter activity of 13 out of 37 flavonoid compounds from four different subclasses, revealed 13 phytoestrogenic compounds able to transcriptionally activate ER in a dose-dependent manner. The red line represents a 2-fold cutoff for positive hits. RLU, relative luciferase unit, normalized to β -gal control. Data are shown as mean $\pm$ S.D. (B) Relative ligand-binding affinity of 12 compounds to ER α or ER β .

gene and ATP6V0E1 has been implicated to be an ER α / β -specific target gene (Grober et al., 2011). Because compound **29** was found to induce ER α / β and ER β / β dimerization but not ER α / α homodimers at 1 μ M, we went on to examine if compound **29** differentially recruited ER α and ER β to the target gene promoters at a dose (1 μ M) that elicits dimer specificity. Chromatin immunoprecipitation (ChIP) and quantitative polymerase chain reaction analysis showed that compound **29** treatment increased ER β association at the promoters of both BAG1 and ATP6V0E1 genes compared with the DMSO control. In contrast, compound **29** was only able to increase ER α recruitment to ATP6V0E1 but not to the BAG1 promoter. This result is consistent with the classification of compound **29** as an ER β / β and ER α / β dimer inducer by BRET assay and that ATP6V0E1 is likely regulated by ER α / β

heterodimer versus BAG1, which is likely regulated by ER β / β homodimer (Fig. 2, G and H).

Of the tested compounds, only three selectively induced ER α / β heterodimerization at select concentrations (compounds **17**, **23**, and **24**); however, three other compounds **18**, **26**, and **29** preferentially induced ER α / β and ER β / β dimers over ER α / α homodimers. Interestingly, compounds **17**, **18**, **23**, **24**, **26**, and **29**, which induced ER α / β and ER β / β dimers, showed higher binding affinity for ER β than for ER α (Fig. 1B). These six compounds, together with the ER α / β heterodimer-inducing compound cosmosiin, constitute a lead heterodimer-selective compound library for pharmacophore development.

Pharmacophore Development Using the GALAHAD Module in SYBYL. The structures of 37 compounds from the initial data set (Table 1) were built into the SYBYL software

Fig. 2. BRET assays in HEK293 cells show dimer selectivity of different flavonoid subclasses. (A–C) Fold change of BRET ratios when cells were treated with indicated compounds: (A) $ER\alpha/\alpha$, (B) $ER\beta/\beta$, and (C) $ER\alpha/\beta$ (10 nM E2 was used as a positive control). Each compound represents an individual experiment; those that induced dimer interaction at a threshold value of $P < 0.05$ were considered statistically significant. Fold change is relative to the negative control DMSO. Data are shown as mean $\pm$ S.D. of three biologic replicates. *Indicates compounds that significantly induced dimerization as determined by two-way analysis of variance. (D) Western blot analysis of Flag-tagged ERβ in MCF7-Flag-ERβ cells. (E and F) Relative ATP6V0E1 and BAG1 mRNAs levels in MCF7-Flag-ERβ cells treated with indicated compounds. (G) Compound 29-induced recruitment of ERβ to the BAG1 and ATP6V0E1 promoters in MCF7-Flag-ERβ cells shown by ChIP assays. (H) The enrichment of ERα on the BAG1 and ATP6V0E1 promoters in MCF7-Flag-ERβ cells after compound 29 treatment shown by ChIP assays. *Indicates statistically significant $P < 0.05$, ** $P < 0.01$.

platform using the sketch module, where hydrogens were added to every structure and energy minimized, and saved as Mol2 files. All structures were then converted into a 3D conformation for each input structure.

Compounds **17**, **18**, **23**, **24**, **26**, and **29** plus **cosmosiin** were used as the training set (Fig. 3) to build a pharmacophore model in the GALAHAD module. Ligands were flexibly aligned to each other completely independent of a template. This generates a molecular alignment based on the pharmacophoric features of the final conformations of the training set. Twenty pharmacophore models were generated by GALAHAD; each of the models represents a different trade-off among competing criteria (Supplemental Table 1). These models contained the same number of features and specificity. The most significant pharmacophore hypothesis models are characterized by assessing the relation between maximizing pharmacophore consensus, maximizing steric consensus, and minimizing conformer potential energy (Caballero, 2010). Within each set of hypotheses, models were first ranked by Pareto surface score (sterics vs. energy), of where the best model has the lowest energy and the highest steric score, as illustrated in the upper left-hand corner of Fig. 3B. Concerning the energy and pharmacophoric similarity criteria, the best model with low energy and high hydrogen-bond score lies

in the upper left-hand corner of the graph in Fig. 3B. Finally, the best model judged by the pharmacophoric similarity and sterics scores lies at the upper-right corner of Fig. 3B (bottom). In Fig. 3B, the ideal model in each ranking is depicted by an open circle. Taking all models into consideration, Model 6 (represented by a red diamond in Fig. 3) had a balanced consensus ranking in all three criteria, and thus was chosen as the best GALAHAD model (Fig. 3C).

Model 6 is comprised of one conformer for each molecule in the training set. All conformers aligned represent low-energy conformations of the molecules, and the final alignment shows a satisfactory superimposition of the pharmacophoric points. Model 6 contains seven key features, including three hydrophobes, three acceptor atoms, and one donor atom. The pharmacophore model clearly shows the importance of the hydrophobic center that is essential in the ER pharmacophores for $ER\alpha$ and $ER\beta$ -selective ligands (Anstead et al., 1997; Brzozowski et al., 1997). The pharmacophore model was validated for its ability to identify $ER\alpha/\beta$ heterodimer-selective compounds from the full data set (Supplemental Table 2).

3D Virtual Screening of the ChemBridge and Maybridge Databases Identified 167 Compounds as Potential Hits. The pharmacophore model was converted into a 3D search query using SYBYL's UNITY-3D module. The search

Fig. 3. Generation and selection of a pharmacophore hypothesis model of ER α / β heterodimer-inducing ligands. A ligand-based pharmacophore hypothesis was generated using GALAHAD. (A) Structures and bioactivity values of the training set chemicals used to generate ligand-based pharmacophore. The structures of the six lead compounds (cosmosiin, two isoflavones, four flavanones, and a flavone) identified from the cell-based assays. (B) Plot of the different criteria used to select the best model. Plot of the energy, sterics, Mol. QRY and H_Bond values for GALAHAD models with selected four ligands that contribute to the consensus feature. (A) Sterics vs. energy (B) Pharmacophore similarity vs. energy (C) Pharmacophore similarity vs. sterics. The open circle represents the ideal best scoring for each condition. The red diamond represents model 6. (C) Selected pharmacophore hypothesis GALAHAD model. GALAHAD assumes pharmacophore/shape and alignments from sets of ligand molecules, to generate a pharmacophore hypothesis that can be used for a 3D search query. GALAHAD models were derived by using the ligands in the training set, which contains seven features identified by GALAHAD represented by blue, green, and purple spheres. The three hydrophobes are centered in the benzopyran and phenyl rings. The three acceptor atoms are in green and a donor atom is in purple.

query was then used to screen the commercial chemical databases from ChemBridge and Maybridge. Both chemical libraries were retrieved from the Zinc Database (<http://zinc.docking.org/>), a free database of commercially available libraries for virtual screening. Flexible 3D screening with no restrictions of both databases was performed using the UNITY tool (Fig. 4A). A total number of 900 initial molecules were identified as hits, many of which contained different chemical scaffolds.

The hits were manually inspected to ensure all chemical groups from the compounds spatially overlapped with the corresponding features of the pharmacophore model. After visual inspection, 81% of the hits failed to match all seven pharmacophoric features, and thus were discarded. The 167 remaining hits contained 19 different core scaffolds that matched the spatial arrangements of our pharmacophore hypothesis. Subsequently, the hits were ranked using SYBYL's integrated ranking features (Supplemental Tables 3 and 4), among which the top 22 hits were purchased and further characterized (Supplemental Table 5).

Validation and Characterization of Hits. The hits were confirmed to activate ER transcription in T47D-KBLuc reporter assay at 10 μ M final concentration (Supplemental Fig. 3) and to induce dimerization in BRET assays (Supplemental Table 6). The ability of compounds to induce all three ER dimer pairs were tested at increasing doses between 1 and 20 μ M in BRET assays (data not shown). Four compounds selectively induced the ER α / β heterodimer but not the ER α / α or ER β / β homodimer at specific concentrations (Fig. 4B). The lowest concentration at which these four compounds selectively induce ER α / β heterodimer is 1 μ M.

The binding affinity of compounds 4, 6, 9, and 10 to ER α and ER β were measured by in vitro fluorescence polarization assay (Fig. 4C). The relative binding affinities are calculated as IC₅₀ values. Compounds 9 and 10 elicit the highest binding affinity. The IC₅₀ values for compound 9 to ER α and ER β were 1.4 and 2.0 μ M, respectively. The IC₅₀ values for compound 10 to ER α and ER β were 1.9 and 3.2 μ M,

Fig. 4. 3D search query of two commercially available databases, the Chembridge and the Maybridge databases, which together have over a million chemicals, resulted in a refined hit list of 167 compounds. (A) Represents a schematic of the 3D virtual screening of the ChemBridge and Maybridge databases. (B) Dose-response data of BRET assays in HEK293 cells, illustrating dimerization profile of selected hits. Data are shown as mean $\pm$ S.D. of three biologic replicates. Data are normalized to DMSO control. (C) Measurement of compound binding to ER α and ER β using in vitro fluorescence polarization competition binding assays.

respectively. Although like compound **29**, which was used to build the pharmacophore model, compounds **9** and **10** induced ER α / β heterodimer at 1 μ M, but they elicited improved binding affinity to ER α and ER β (Fig. 4C), suggesting that

in silico modeling expedites identification of stronger ER agonists with similar heterodimer specificity. Thus, compounds **9** and **10** would be better compounds to pursue for probing ER α / β heterodimer functions.

ER α Is the Dominant Heterodimeric Partner in the Presence of Selective ER α / β Heterodimer Compounds.

We previously reported that E2 induces heterodimer formation by binding to ER α (Powell and Xu, 2008). To examine whether the selective heterodimer-inducing compounds also induce heterodimer via binding to ER α , BRET assays were performed with a combination of wild-type and mutant ER α and ER β constructs (Powell and Xu, 2008). The expressed mutant proteins contained a single mutation in the LBD (ER α G521R and ER β G491R) of receptors that ablate ligand binding (Tremblay et al., 1999; Powell and Xu, 2008). A combination of wild-type and mutant ER α and ER β fusion proteins were used: ER α G521R-RLuc with wild-type YFP-ER β , YFP-ER β G491R with wild-type ER α -RLuc, wild-type YFP-ER β with wild-type ER α -RLuc, and ER α G521R-RLuc with YFP-ER β G491R. As has been previously reported for E2 (Powell and Xu, 2008), ligand-binding competent ER α LBD but not ER β LBD is required for heterodimerization in the presence of compounds, reinforcing that ER α is the dominant partner for heterodimerization (Fig. 5).

Discussion

Current ER-positive breast cancer therapies target ER α , either using selective ER modulators to inhibit ER α transcriptional activity or selective ER degraders to reduce ER α protein levels. However, the therapeutic potential of ER β in breast cancer has been poorly investigated. Our previous studies using ER α / β heterodimer-selective ligands show that ER β , via heterodimerization with ER α , can antagonize the pro-proliferative effects of ER α , rendering the heterodimer as a novel preventive or therapeutic target for hormone-dependent diseases. However, few ER α / β heterodimer-inducing selective compounds have been discovered and they generally elicit only weak binding affinities to ERs. Therefore, the goal of this study

was to combine computational and experimental approaches to identify compounds with improved binding affinity and dimerization specificity.

Emerging biochemical evidence supports the formation of ER α / β heterodimers when two receptors are coexpressed (Cowley et al., 1997; Pettersson et al., 1997); in particular, ER α / β heterodimers were recently detected in breast tissue using proximity ligation assay (Iwabuchi et al., 2017). However, the functions of ER α / β heterodimers remain elusive due to the lack of a crystal structure and heterodimer-specific compounds. Uncovering the biologic function of the ER α / β heterodimer is important for understanding ER signaling and designing ER-targeted therapeutics based on receptor dimerization status. The main distinction of heterodimer-inducing compounds from the existing selective ER modulators and degraders is that they target different steps in ER activation. ER heterodimer compounds target ER dimerization, a step between the ligand binding and the receptor association with chromatin. In our previously published report, we have shown that selective ER modulators such as tamoxifen, raloxifene, and the full ER antagonist ICI 182,780 do not interfere with the formation of all three dimer pairs (Powell and Xu, 2008). Although more studies are needed to demonstrate that the ER α / β heterodimer indeed serves as a therapeutic target, the concept of inducing ER β to pair with ER α , thus antagonizing ER α 's proliferative function, is distinct from the existing breast cancer therapies to target ER α alone.

We reason that identifying and improving chemical probes would be an essential step toward understanding the biologic role of ER α / β heterodimers. Natural phytoestrogens often elicit higher binding affinity to ER β than to ER α (Kuiper et al., 1997, 1998). Several phytoestrogens showing slight selectivity for ER α / β heterodimers were found to be antiproliferative in cancer cell lines coexpressing ER α and ER β (Powell and Xu, 2008; Powell et al., 2010, 2012). However, the slight selectivity

Fig. 5. Mutant ER α and ER β LBDs reveal ER α as the dominant heterodimeric partner in the presence of selective ER α / β heterodimer compounds. (A) Heterodimerization of the wild-type ER α and ER β . (B) Mutation in the ER β LBD does not affect heterodimerization with ER α . (C) Heterodimerization of mutant ER α with mutant ER β . (D) No dimerization is observed between mutant ER α and wild-type ER β . Data are shown as mean $\pm$ S.D. * Indicates statistically significant <0.05 .

and low potency of these compounds prevent definitive elucidation of the functions of ER α / β heterodimers. Pharmacophore-based techniques and virtual screening have successfully been employed for the discovery of ER subtype-selective ligands (Huang et al., 2015). Herein, using a combination of cell-based assays (i.e., reporter and BRET), pharmacophore modeling, and virtual screening, we identified four ER α / β heterodimer-selective ligands (Table 2) with improved efficacy compared with cosmosiin, a previously identified compound with slight preference for the ER α / β heterodimer. The main hurdle in identifying ER α / β heterodimer-selective ligands lies in the lack of a crystal structure. Ligand binding is necessary but insufficient for the formation of ER α / β heterodimers. Previous studies suggest that ligand binding is essential to induce a conformational change of ER to accommodate helix 12 in functional dimers. In this process, ER α and ER β appear to play separate roles such that ligand-bound ER α is the dominant partner in heterodimer formation. We have shown that ER β subtype-specific ligands promote the formation of ER β / β homodimers but ER α subtype-specific ligands could induce both ER α / α homodimers and ER α / β heterodimers (Powell and Xu, 2008). Because of the lack of protein crystal structures needed to build a structure-based pharmacophore model for a virtual ligand screen, we combined a multistep screening strategy with a ligand-based pharmacophore model to identify ER α / β

heterodimer-selective ligands. We confirmed that ligand binding is necessary but insufficient for inducing ER dimerization. Furthermore, the formation of ER homo- versus heterodimers appears to be ligand concentration dependent (Figs. 2 and 4). Our results also showed that a ligand must induce a conformational change in ER α in a manner such that it preferentially selects the other ER subtype as a partner (Fig. 5). Finally, we characterized the estrogenic activity and dimerization ability of 59 compounds, leading to the identification of four ER α / β heterodimer-selective ligands. To our knowledge, building a pharmacophore model to identify the chemical features responsible for induction of ER α / β heterodimers is unprecedented. Thus, the more selective and potent compounds identified in this study will serve as useful probes to elucidate ER α / β heterodimer functions in vitro and in vivo.

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Authorship Contributions

Participated in research design: Coriano, Liu, Xu, Lim.
Conducted experiments: Coriano, Liu, Sievers, Wang.
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Wrote or contributed to the writing of the manuscript: Coriano, Liu, Xu.

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TABLE 2

The structural arrangement of 5 ER α / β heterodimer-selective compounds

Compound Number	Structure	BRET
4		α/β at 10 μ M
6		α/β at 1 μ M
9		α/β at 1 μ M
10		α/β at 1 μ M

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