

Evaluation of a Recombinant Yeast Cell Estrogen Screening Assay

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A wide range of chemicals with diverse structures derived from plant and environmental origins are reported to have hormonal activity. The potential for appreciable exposure of humans to such substances prompts the need to develop sensitive screening methods to quantitate and evaluate the risk to the public. Yeast cells transformed with plasmids encoding the human estrogen receptor and an estrogen responsive promoter linked to a reporter gene were evaluated for screening compounds for estrogenic activity. Relative sensitivity to estrogens was evaluated by reference to 17 β -estradiol (E₂) calibration curves derived using the recombinant yeast cells, MCF-7 human breast cancer cells, and a prepubertal mouse uterotrophic bioassay. The recombinant yeast cell bioassay (RCBA) was approximately two and five orders of magnitude more sensitive to E₂ than MCF-7 cells and the uterotrophic assay, respectively. The estrogenic potency of 53 chemicals, including steroid hormones, synthetic estrogens, environmental pollutants, and phytoestrogens, was measured using the RCBA. Potency values produced with the RCBA relative to E₂ (100) included estrone (9.6), diethylstilbestrol (74.3), tamoxifen (0.0047), α -zearalanol (1.3), equol (0.085), 4-nonylphenol (0.005), and butylbenzyl phthalate (0.0004), which were similar to literature values but generally higher than those produced by the uterotrophic assay. Exquisite sensitivity, absence of test compound biotransformation, ease of use, and the possibility of measuring antiestrogenic activity are important attributes that argue for the suitability of the RCBA in screening for potential xenoestrogens to evaluate risk to humans, wildlife, and the environment. **Key words:** antiestrogens, environmental contaminants, estrogen, metabolism, phytoestrogens, potency, receptor, recombinant yeast cell bioassay, uterotrophic. *Environ Health Perspect* 105:734–742 (1997)

There is considerable concern within the scientific community and media that the increasing occurrence of endocrine-related abnormalities in humans and wildlife may be associated with exposure to environmental pollutants capable of mimicking or modulating the action of natural hormones. Appreciation that many ubiquitous compounds of natural and anthropogenic origins are estrogenic has led to the hypothesis that exposure to such compounds *in utero* may be involved in falling sperm counts and disorders of the male reproductive tract (1). Although such estrogenic compounds are structurally heterogeneous (2), there are similarities including lipophilicity that facilitate accumulation in food-producing animals and potentiate risks associated with entry into the human food chain (3). In the United Kingdom these concerns have been given added impetus following a report by the Institute for Environment and Health (4), which made several recommendations aimed at investigating the risks of hormonal mimics; these included the need to develop robust and reliable assays capable of screening chemicals for estrogenic activity. Given that prediction of estrogenic potency through structural information alone is not yet possible, development of such generic assays is necessary because many existing assay systems are not sufficiently robust for

nonexpert operation or may not accurately reflect human potency.

Methods of detecting and assessing estrogenic compounds have been previously described (5) and critically evaluated (6). Biotransformation and consequent alteration of hormonal activity by test systems is an important consideration because phenolic metabolites produced by the action of cytochrome P450 enzymes can be more potent than the parent compounds (7,8). Clearly, although such metabolic effects are intrinsic to *in vivo* models, they may not be so readily reflected by *in vitro* test systems. Estrogen-sensitive human breast cancer cells, such as MCF-7 cells, express the human estrogen receptor and have been used to develop E-screen tests for chemicals (9) and contaminants in animal feeds (10). Uterotrophic assays such as the classical mouse uterine weight bioassay (11) have limitations (6), but are still preferred methods for many investigators (12). Uterotrophic assays also offer the potential to combine biochemical and histological analysis of estrogen-mediated events and further adaptation to measure antiuterotrophic activity (13).

The use of receptors linked to reporter genes in transformed cellular systems to detect biologically active xenobiotics has been proposed as a strategy to define

chemicals by their functional properties (2). This approach facilitates detection of not only receptor ligand binding but also response element occupancy and gene activation. The recombinant yeast cells used in the present report have recently been used in an ultrasensitive RCBA for the determination of prepubertal plasma estrogens (14). These cells contain an expression plasmid that includes the CUP1 metallothionein promoter fused to the human estrogen receptor cDNA and a reporter plasmid which contains two copies of the frog vitellogenin estrogen response element upstream of the yeast iso-1-cytochrome c promoter fused to the structural gene for β -galactosidase. Similar recombinant yeast cell systems have been used for the detection of xenoestrogens (15,16) and various estrogen receptor studies (17–19). The particular advantage of the transformed yeast cell line approach is that the cells are robust and substrate auxotrophy may be used to continuously select for estrogen sensitivity. Moreover, expression of human estrogen receptor and ability to automate suggest that these yeast cells have much to offer the analyst requiring an *in vitro* screening assay that affords some reflection of potential estrogenic activity in humans.

In the present study, bioassays were evaluated by comparison of the potency of test compounds relative to 17 β -estradiol (E₂) since this natural estrogen is the most commonly accepted positive control used in such *in vivo* and *in vitro* assays (20,21). Validation of the recombinant yeast cell bioassay (RCBA) is addressed through comparison of potency values with an alternative established *in vivo* bioassay and with potency values produced using similar *in vitro* assays described in the literature. Similar approaches to validation have been commonly used (6,20) and serve to identify some of the merits and limitations of an assay relative to other *in vitro* and *in vivo* methods. The metabolic fate of test compounds in the selected bioassay is another

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Table 1. Comparison of RCBA results and estrogenic potencies described in the literature

Source ^a	Compound tested	RP	RIE	RPL
1	Steroidal estrogens			
1	17 β -Estradiol	100	100	—
1	17 β -Estradiol-3(β -D-glucuronide)	0.32	89.6	—
1	17 β -Estradiol-3-sulfate	0.01	20.4	—
1	17 β -Estradiol-3-glucuronide-17-sulfate	0	0	—
1	17 α -Estradiol	5.25	100	49 ^b , 8 ^c
1	Estrone	9.6	100	5 ^d , 66 ^b , 8 ^c
1	Estriol	0.63	88	16 ^b , 14 ^c
	Steroids			
1	Cholesterol	0	0	—
1	Androstenedione	0	0	—
1	Testosterone	0.001	0.8	0.01 ^c
1	Androstenediol	0.023	31	—
1	Dehydroepiandrosterone	0.0018	1.2	—
1	Cortisol	0	0	0.01 ^c
1	Progesterone	0	0	0.01 ^c
1	D-Norgestrel	0.0004	3.1	—
	Synthetic estrogens			
1	17 α -Ethinylestradiol	88.8	100	191 ^b , 95 ^c
1	Mestranol	7.3	80	8 ^b
1	Diethylstilbestrol	74.3	100	246 ^b , 73 ^c , 73 ^e
1	Hexestrol	30.6	89	74 ^b , 55 ^c , 319 ^e , 5 ^f
1	Dienestrol	25.4	84	26 ^c , 102 ^e
	Triphenylethylene antiestrogens			
1	Tamoxifen	0.0047	46	<0.01 ^d
1	4-Hydroxytamoxifen	0.0073	56	—
1	Nafoxidine	0	0	0.01 ^d
	Fungal resorcylic acid lactones			
1	Zearalenone	0.26	91	1.4 ^c , 0.85 ^e , 0.5 ^f
1	α -Zearalenol	8.7	67	27 ^c
1	β -Zearalenol	0.066	58	0.43 ^c , 0.043 ^e , 0.07 ^f
1	α -Zearalanol (zearanol)	1.3	84	17 ^c , 47 ^e
1	β -Zearalanol	0.46		3.4 ^c
	Phytoestrogens			
2	Coumestrol	0.67	75	0.11 ^e
2	Equol	0.085	50	0.2 ^c
2	Daidzein	0.0013	5.4	0.00072 ^e , 0.01 ^f
2	Formononetin	0.0056	23	0.0004 ^e
2	Bochanin A	0.0091	61	0.0048 ^e , 0.0005 ^f
2	Genistein	0.049	61	0.08 ^e , 0.011 ^f
	4-Alkylphenols			
3	4-Nonylphenol (technical grade)	0.005	57	0.003 ^g
4	4-Nonylphenol (straight chain)	0.0022	38.3	—
3	4-Octylphenol	0.003	22.3	0.03 ^g
3	4-tert-Octylphenol	0.00036	43	—
	Organochlorines			
1	DDT	0.00003	0.8	—
6	<i>o,p'</i> -DDT	0.00011	2.4	—
6	<i>o,p'</i> -DDE	0.00004	1.0	—
5	2,3,7,8-Tetrachlorodibenzo- <i>p</i> -dioxin	0.26	5.5	—
3	Methoxychlor	0.0033	55	—
6	4'-Chloro-4-biphenylol	0.06	73.3	0.025 ^h
6	2'-Chloro-4-biphenylol	0.0037	32	—
6	2',5'-dichloro-4-biphenylol	0.62	65	0.198 ^h
6	2',4',6'-Trichloro-4-biphenylol	1.0	67.6	2.38 ^h
6	2',3',4',5'-Tetrachloro-4-biphenylol	0.82	77	1.05 ^h
6	3,3',5,5'-Tetrachloro-4,4'-biphenyldiol	0.016	83.5	0.074 ^h
	Other contaminants			
1	Bisphenol A	0.005	51	0.003 ^g , 0.05–0.02 ⁱ
1	Di- <i>n</i> -butylphthalate	0	0	—
6	Butylbenzylphthalate	0.0004	5.3	0.0003 ^g
1	Bis(2-ethylhexyl)phthalate	0	0	—
1	Butylhydroxytoluene	0	0	—

Abbreviations: E₂, 17 β -estradiol; RP, relative potency compared to E₂ (100) by molar mass determined with the recombinant yeast cell bioassay; RIE, relative inductive efficiency indicates the ratio between maximal (continued, next column)

important element of assay validation that is addressed because this helps to provide a better understanding of the significance of experimental observations.

Materials and Methods

Source of chemicals. [³H]Coumestrol (518 GBq/mmol; custom synthesis), di-*n*-butyl [carboxyl-¹⁴C]phthalate (962 MBq/mmol), [monoethyl-³H]diethylstilbestrol (3.4 TBq/mmol), 4-*n*-nonyl[ring 2,6(*n*)-³H]phenol (1.96 TBq/mmol; custom synthesis), [6,7-³H]17 β -estradiol (1.89 TBq/mmol), [³H]testosterone (3.88 TBq/mmol), and α -[³H]zearanol (1.92 TBq/mmol; custom synthesis) were obtained from Amersham International (Little Chalfont, U.K.). Radiochemical purity was checked by radio-HPLC using the conditions described below and found to be greater than 96% in all cases. Sources of chemicals tested for estrogenic potency are provided in Table 1.

Transformed yeast cells. Recombinant estrogen sensitive yeast cells (14) were stored in 30% glycerol at -80°C and grown as required at 30°C on selective media agar plates containing 0.17% yeast nitrogen base (without amino acids and ammonium sulfate), 0.5% (NH₄)₂SO₄, 2% dextrose, L-leucine (60 μ g/ml), L-histidine (20 μ g/ml) and 2% Bacto agar (Difco, East Molsey, U.K.).

RCBA of estrogens. Procedures that were followed were similar to those described previously (14). Briefly, a small colony of

(Table 1 continued) β -galactosidase activity achieved with the test compound and that of E₂ $\times$ 100; RPL, relative potency values compared to 17 β -estradiol derived using molar mass units sourced from the literature as indicated by the superscript. Where necessary, data were normalized relative to E₂ (100). Maximum concentration of 2,3,7,8-tetrachlorodibenzodioxin tested was 100 nM.

^aSources of chemicals: 1, Sigma Chemical Company, Poole, Dorset, U.K.; 2, Apin Chemicals Ltd., Abingdon, Oxon, U.K.; 3, Lancaster Synthesis Ltd., Morecambe, Lancashire, U.K.; 4, Aldrich Chemical Company, Gillingham, Dorset, U.K.; 5, Promochem Ltd., Welwyn Garden City, Herts, U.K.; 6, Mallinckrodt Baker, Milton Keynes, Bucks, U.K.

^bCompetitive binding radioreceptor assay; data from Korenman (21).

^cCompetitive binding radioreceptor assay; data from Arts and Van Den Berg (22).

^dE₂ receptor processing; data from Gyling and Leclercq (23).

^eMCF-7 cell proliferation assay; data from Welshons et al. (10).

^fHeLa cell cotransfection assay; data from Miksicek (20).

^gRelative proliferative potency in E-screen assay; data from Sonnenschien, et al. (24).

^hCompetitive binding radio-receptor assay; data from Korach et al. (8).

ⁱCompetitive binding radioreceptor assay–progesterone receptor induction; data from Krishnan et al. (25).

yeast cells was sampled from an agar plate and incubated for 18 hr at 30°C by continuous shaking in 2 ml selective media broth. The yeast cells were collected by centrifugation and diluted in 1 M mannose to an optical density of 0.1 (A_{630}). Yeast cell suspension was dispensed (20 μ l) into wells of 96-well plates (Nunc; Life Technologies, Paisley, U.K.) containing 75 μ l of 50 μ M copper sulfate in selective media and incubated with 25 μ l calibration standard. Calibration standards were prepared in ethanol at a concentration of 10 mM and serially diluted ($\times 10$) in selective media broth. After an 18-hr incubation at 30°C, the yeast cell density in the wells was estimated from the optical density at 630 nm using a Dynatech microtiter plate reader (Dynatech Labs, Billingshurst, U.K.). Induction of β -galactosidase activity was measured by addition of 100 μ l Z buffer (60 mM Na_2HPO_4 , 40 mM NaH_2PO_4 , 10 mM KCl, 50 mM mercaptoethanol, 1 mM MgSO_4) containing 2 mg/ml *o*-nitrophenyl- β -D-galactopyranoside (ONPG), 0.1% sodium dodecyl sulfate, and 250 units/ml lyticase (Sigma Chemical Co., Poole, Dorset, U.K.) to each well, and quantified by determination of *o*-nitrophenol production expressed as $(\text{OD}_{410\text{nm}} / \text{OD}_{630\text{nm}}) \times 1000/\text{min}$. Wells were read at 6 min and at 4 hr according to the level of induced β -galactosidase activity.

RCBA of antiestrogens. β -Galactosidase activity induced by E_2 (100 fM–100 nM) in the presence of the potent antiestrogen 4-hydroxytamoxifen (1 nM–10 μ M) was investigated with recombinant yeast cells as described above.

Estrogenic potency of test compounds in RCBA. A range of chemicals was tested for estrogenic activity in at least two separate experiments using the RCBA. Calibration curves of E_2 and test compounds were produced by analysis of standards (100 fM–10 nM for E_2 and potent estrogens and 10 pM–10 μ M for weak estrogens) in selective media for estrogenic activity. Each calibration standard of a test compound was analyzed in 8 wells (1 column) and assay blanks (vehicle only) in 16 wells (2 columns) of a 96-well microtitre plate. Each test compound was assayed in a separate 96-well microtitre plate and the wells were sealed with film plate sealers. β -Galactosidase activity in blank wells (typically $\text{OD}_{410\text{nm}}$ 0.080 $\pm$ 0.012 and 0.109 $\pm$ 0.009 at 6 min and 4 hr, respectively) was subtracted from wells containing standard concentrations of test chemicals. The relative potency of test compounds was determined from the concentration of E_2 and test compound that provided 50% induction of β -galactosidase activity (EC_{50}) and calculated by dividing the E_2 EC_{50} by the test compound $\text{EC}_{50} \times 100$. The relative potency of weak estrogens that failed to induce β -galac-

tosidase to 50% of the total E_2 activity was determined from the concentration of E_2 and test compound that provided the same induction of β -galactosidase activity and calculated similarly. The relative inductive efficiency (RIE) was determined as the ratio of maximal β -galactosidase activity induction with test compound to $\text{E}_2 \times 100$.

Uterotrophic assay. Prepubertal 18-day-old CFLP female mice (Harlan U.K. Ltd., Huntingdon, U.K.) were injected subcutaneously with 0.1 ml test compound dissolved in corn oil at the doses shown in Table 2, on three consecutive days, essentially as described by Rubin et al. (11). In each assay, five concentrations of E_2 (calibrant) and three concentrations of test compound were injected using seven animals per dose. Mice were sacrificed on the fourth day and

the weights of the animal and uteri recorded; a sample of vagina was fixed in phosphate buffered saline containing 1% formaldehyde and processed for routine histological evaluation. Results are expressed as the mean ratio of uterine weight to body weight. The relative potency of test compounds was determined by interpolation from calibration curves of the molar dose of E_2 and test compounds that provided similar increases in ratios of uterine wt/body wt $\times 100$. Care and treatment of the mice was in accordance with the Animals (Scientific Procedures) Act 1986 of the United Kingdom and supervised by a veterinary surgeon.

Human breast cancer cell proliferation assay. MCF-7 human breast cancer cells were maintained in Eagles minimal essential medium (EMEM) supplemented with 5%

Table 2. Effect of test compounds on prepubertal mouse uterine weight and comparison with the RCBA

Compound	Dose	Uterine wt/body wt ratio (mean $\pm$ SD)	Relative uterotrophic potency (molar)	Relative potency (RCBA)
Control	0	0.097 $\pm$ 0.030	—	—
17 β -Estradiol	5 ng	0.172 $\pm$ 0.028**	100	100
	10 ng	0.247 $\pm$ 0.030 [#]	—	—
	25 ng	0.405 $\pm$ 0.114 [#]	—	—
	50 ng	0.420 $\pm$ 0.121 [#]	—	—
	100 ng	0.554 $\pm$ 0.126 [#]	—	—
Diethylstilbestrol	0.5 ng	0.076 $\pm$ 0.029	234	74.3
	5 ng	0.137 $\pm$ 0.034**	—	—
	50 ng	0.715 $\pm$ 0.260**	—	—
4-Nonylphenol (technical grade, Sigma)	0.5 mg	0.086 $\pm$ 0.030	0.00036	0.005
	1 mg	0.094 $\pm$ 0.021	—	—
	5 mg	0.351 $\pm$ 0.192*	—	—
	20 mg ^a	—	—	—
4-Octylphenol	0.5 mg	0.074 $\pm$ 0.034	0	0.003
	1 mg	0.097 $\pm$ 0.034	—	—
	5 mg	0.085 $\pm$ 0.036	—	—
Coumestrol	1 μ g	0.104 $\pm$ 0.026	0.024	0.67
	10 μ g	0.132 $\pm$ 0.009 [#]	—	—
	100 μ g	0.436 $\pm$ 0.182 [#]	—	—
Benzylbutylphthalate	0.05 mg	0.086 $\pm$ 0.024	0	0.0004
	0.5 mg	0.093 $\pm$ 0.031	—	—
	5 mg	0.108 $\pm$ 0.027	—	—
Dibutylphthalate	0.05 mg	0.119 $\pm$ 0.036	0	0
	0.5 mg	0.101 $\pm$ 0.028	—	—
	5 mg	0.091 $\pm$ 0.014	—	—
α -Zearalanol	1 μ g	0.139 $\pm$ 0.029**	0.026	1.3
	10 μ g	0.190 $\pm$ 0.035 [#]	—	—
	100 μ g	0.411 $\pm$ 0.27*	—	—
Bisphenol A	0.05 mg	0.098 $\pm$ 0.014	0	0.005
	0.5 mg	0.092 $\pm$ 0.021	—	—
	5 mg ^a	—	—	—

Abbreviations: RCBA, recombinant yeast cell bioassay; SD, standard deviation. Potency values determined with the murine uterotrophic assay are presented with the RCBA for comparison. Statistical significance is relative to untreated controls.

^aAnimals from this treatment group were withdrawn from the study due to toxic effects of the dose.

* $p < 0.05$; ** $p < 0.01$; [#] $p < 0.001$.

(v/v) fetal calf serum (FCS), 20 mM HEPES, 2 mM glutamine nonessential amino acids, 0.075% sodium bicarbonate, and 600 ng/ml insulin. The proliferative effect of E_2 was determined by seeding 5×10^4 cells into replicate 25-cm² flasks (four per E_2 concentration) in EMEM supplemented as above. After 3 days incubation, the seeding medium was removed and the monolayers were washed with Earle's balanced salt solution (phenol red-free) and replaced with medium containing phenol red-free EMEM, dextran charcoal stripped FCS, supplemented as above with the exception of insulin. After a further 3-day incubation period to enable the MCF-7 cells to return to basal estrogen non-stimulated status (10) the cultures were treated with medium containing E_2 (1–10,000 pM dissolved in ethanol vehicle) and incubated for 5 days. The cell number per flask was determined by counting cell nuclei as described previously (26).

Biotransformation of steroids and xenoestrogens by yeast cells. Recombinant yeast cells were suspended at an appropriate dilution in minimal media so as to provide an optical density of 0.1 ($A_{630\text{ nm}}$); these and similar blank cultures (without yeast cells) were incubated at 30°C with 100 nM 37 KBq/ml tritium labeled coumestrol, E_2 , diethylstilbestrol, 4-nonylphenol, testosterone, and α -zearanol and 10 μ M [¹⁴C]di-*n*-butylphthalate for 18 hr. The medium was sampled after incubation periods of 1, 4, and 18 hr; it was then centrifuged and stored at -20°C prior to analysis by radio-HPLC.

Metabolite analysis. Samples of medium were analyzed for the presence of metabolites by reverse phase radio-HPLC using either a Hypersil BDS C18 3 μ m 100 $\times$ 4.6 mm (Life Sciences International, Runcorn, U.K.) or a Spherisorb ODS 5 μ m 250 $\times$ 4.6 mm (Jones Chromatography, Hengoed, U.K.) column (for di-*n*-butylphthalate only) and chromatographed at a flow rate of 1 ml/min. Conditions were selected to provide separation of parent compounds from metabolites using conditions similar to those described previously (27). Di-*n*-butylphthalate, testosterone, and E_2 analysis used an isocratic mobile phase consisting of 35% acetonitrile:65% water for 15 min and α -zearanol, coumestrol, and diethylstilbestrol analysis used a linear gradient mobile phase system consisting of 50:50 water methanol containing 0.1% acetic acid to 100% methanol, 0.1% acetic acid, over 25 min. 4-Nonylphenol analysis employed a linear gradient from 20:80 methanol:water to 100% methanol over 15 min, which was maintained for an additional 5 minutes.

The above studies were all conducted at the Central Veterinary Laboratory, United

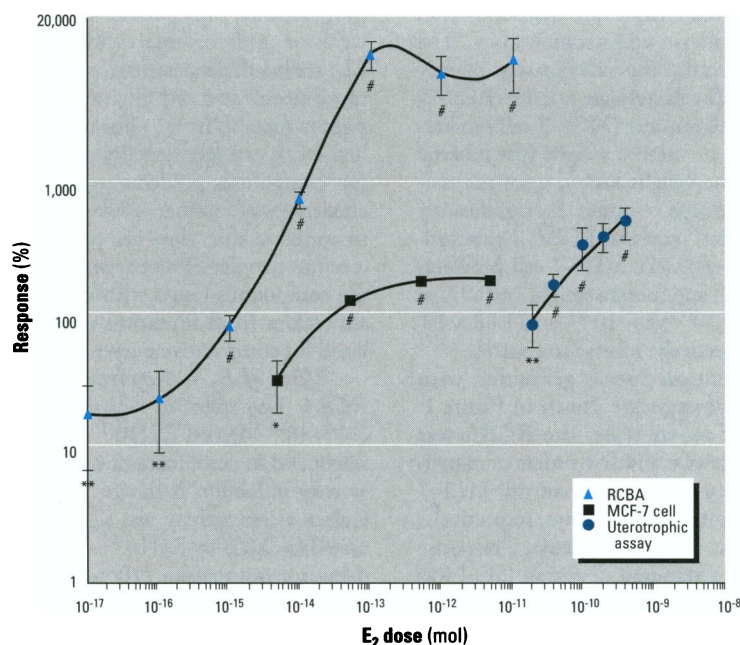


Figure 1. 17 β -Estradiol (E_2) calibration curves derived from the recombinant yeast cell bioassay (RCBA; $n = 8$), the MCF-7 human breast cancer cell proliferation assay ($n = 4$), and the prepubertal mouse uterotrophic assay ($n = 7$). Induction of galactosidase activity (RCBA), increase in cell number (MCF-7 cell proliferation assay), and uterine weight (prepubertal mouse uterine weight assay) were normalized to percentage response and are plotted against the total number of moles of E_2 added to the test system. Error bars represent ± 1 standard deviation.

* $p < 0.05$; ** $p < 0.01$; # $p < 0.001$ relative to values obtained in the absence of E_2 .

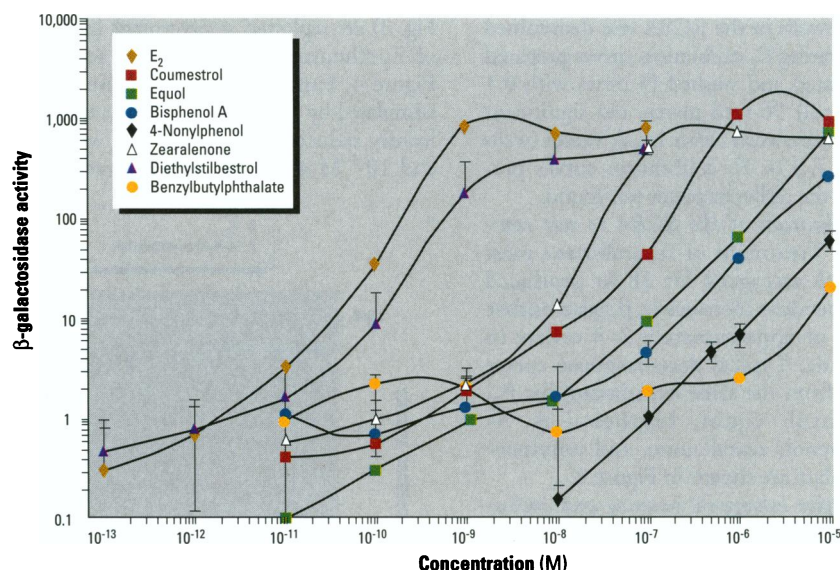


Figure 2. Typical calibration curves for 17 β -estradiol (E_2), coumestrol, equol, bisphenol A, 4-nonylphenol, zearalenone, diethylstilbestrol, and benzylbutylphthalate derived from the recombinant yeast cell bioassay. Error bars represent ± 1 standard deviation ($n = 8$).

Kingdom, between July 1995 and February 1997.

Results

Evaluation of bioassay sensitivity to E_2 The sensitivity of the RCBA, MCF-7 human breast cancer cell, and uterotrophic bioassays were compared by evaluation of respective E_2

calibration curves. Direct comparison of the two *in vitro* assays with the *in vivo* assay was enabled by conversion of the concentration units of E_2 used in the RCBA (100 fM–100 nM) and the MCF-7 proliferation (1–10,000 pM) bioassays to total moles added to the *in vitro* system during the assay because this could be equated with the quantity of

administered dose (mole per mouse) used in the *in vivo* mouse uterotrophic assay. The endpoints recorded for each bioassay, namely induction of β -galactosidase activity (RCBA), increase in cell number (MCF-7 cell proliferation assay), or uterine weight (prepubertal mouse uterine weight assay), were normalized to percentage response. E_2 significantly (Students *t*-test) increased RCBA β -galactosidase activity ($p < 0.01$), MCF-7 cell proliferation ($p < 0.05$), and uterine weight ($p < 0.01$) at total doses of 10^{-17} , 5×10^{-15} , and 1.83×10^{-11} moles respectively, relative to controls.

E_2 calibration curves generated with these three bioassays are shown in Figure 1. In terms of molar dose, the RCBA was approximately two and five orders of magnitude more sensitive to E_2 than the MCF-7 cells and uterotrophic bioassays, respectively. The dynamic range of percentage response to E_2 [mean $\pm$ standard deviation (SD)] was at least an order of magnitude greater for the RCBA (17.37 ± 10.8 – $7754 \pm 3370\%$) than either the MCF-7 proliferation (30 ± 13 – $175 \pm 25\%$) or uterotrophic (79 ± 28 – $472 \pm 130\%$) bioassays. E_2 calibration curves for the recombinant yeast cell and uterine weight bioassays without prior data normalization are shown in Figure 2 and Table 2, respectively. Potential interference of estrogenic residues leaching from the plastic microwells in the RCBA was determined by comparing E_2 calibration curves prepared in untreated and washed (5 times with 0.1 ml ethanol) 96-well plates. No significant difference between either blank values (wells without E_2) or E_2 calibration curves prepared in the different plates was found.

Application of the RCBA to test compounds. Treatment of recombinant yeast cells with estrogens for 18 hr produced dose dependent increases in β -galactosidase activity of approximately 2–3 orders in magnitude. Typical dose–response curves (not all from the same experiment) for E_2 , coumestrol, equol, bisphenol A, 4-nonylphenol, zearalenone, and butylbenzylphthalate are shown in Figure 2.

Relative estrogenic potency and inductive efficiency of test compounds in the RCBA. Dose–response curves for 53 chemicals were produced using the RCBA; from these, estrogenic potency and maximal β -galactosidase induction values relative to E_2 were determined (Table 1). The synthetic stilbenes diethylstilbestrol, hexestrol, and dienestrol and the synthetic steroid 17α -ethynylestradiol had the highest relative potencies and were of a similar order to E_2 . Environmental pollutants such as the 4-alkylphenols were among the weakest estrogens tested, with relative potencies 4–5 orders of magnitude less than E_2 . Steroids including cholesterol, androstenedione,

progesterone, and cortisol had either very weak or undetectable estrogenic activity. The triarylethylene antiestrogens nafoxidine, tamoxifen, and 4-hydroxytamoxifen [a potent metabolite of tamoxifen (28)] also had weak estrogenic activity. Many of the test compounds produced relative inductive efficiencies of less than 100%; this was taken to indicate that they are partial estrogen receptor agonists. Potency values of some of the compounds tested with the RCBA were also taken from literature sources and are listed for comparative purposes in Table 1.

Effect of E_2 + 4-hydroxytamoxifen in RCBA. The effect of 4-hydroxytamoxifen (10^{-9} – 10^{-5} M) and E_2 (10^{-13} – 10^{-7} M), both singly and in combination, on β -galactosidase activity induction is shown in Figure 3. β -Galactosidase activity was significantly stimulated ($p < 0.05$) by E_2 (10^{-13} – 10^{-7} M) and by 4-hydroxytamoxifen (10^{-8} – 10^{-5} M) compared with blank wells. The potential antagonist activity of 4-hydroxytamoxifen was investigated by addition of 4-hydroxytamoxifen (10^{-9} – 10^{-5} M) to E_2 calibration curves (10^{-13} – 10^{-7} M). β -Galactosidase activity stimulated by 1 nM E_2 was significantly reduced ($p > 0.05$) by inclusion of 4-hydroxytamoxifen (10^{-9} – 10^{-5} M) compared with 1 nM E_2 alone, indicating antagonism of E_2 binding; to clarify this point, these data (box, Fig. 3) are replotted as percentage inhibition of E_2 -stimulated β -galactosidase activity in Figure 4. Similarly, β -galactosidase activity stimulated by 10^{-8} and 10^{-7} M E_2 was significantly reduced by combination with 10^{-6} and 10^{-5} M 4-hydroxytamoxifen compared

with E_2 alone. 4-Hydroxytamoxifen displayed agonist activity when tested in isolation; as a result, β -galactosidase activity was significantly ($p > 0.05$) elevated by relatively high doses of 4-hydroxytamoxifen when tested with a low dose of E_2 (e.g., 10^{-13} M E_2 + 10^{-9} – 10^{-5} M 4-hydroxytamoxifen) when compared with E_2 alone (Fig. 3).

Determination of relative estrogenic potency using the mouse uterine weight assay. The uterotrophic effect and relative estrogenic potencies of E_2 , diethylstilbestrol, 4-nonylphenol, 4-octylphenol, coumestrol, α -zearalanol, bisphenol A, dibutylphthalates, and benzylbutylphthalates are shown in Table 2. 4-Nonylphenol, coumestrol, α -zearalanol, and diethylstilbestrol significantly ($p < 0.05$) increased uterine weight at the doses shown. After the first injection of the highest doses of 4-nonylphenol and bisphenol A, symptoms of acute toxicity were evident and, as a consequence, these mice were immediately withdrawn from the study and euthanized.

Xenobiotic biotransformation. Analysis of incubation medium revealed no evidence of xenobiotic metabolism following incubation of yeast cells with E_2 , coumestrol, dibutylphthalate, diethylstilbestrol, testosterone, and 4-nonylphenol. In the case of α -zearalanol, a single metabolite (5% of total) was detected in the medium after 18 hr (data not shown).

Discussion

High sensitivity is an important requirement for screening assays to enable detection of compounds of low potency that may be of

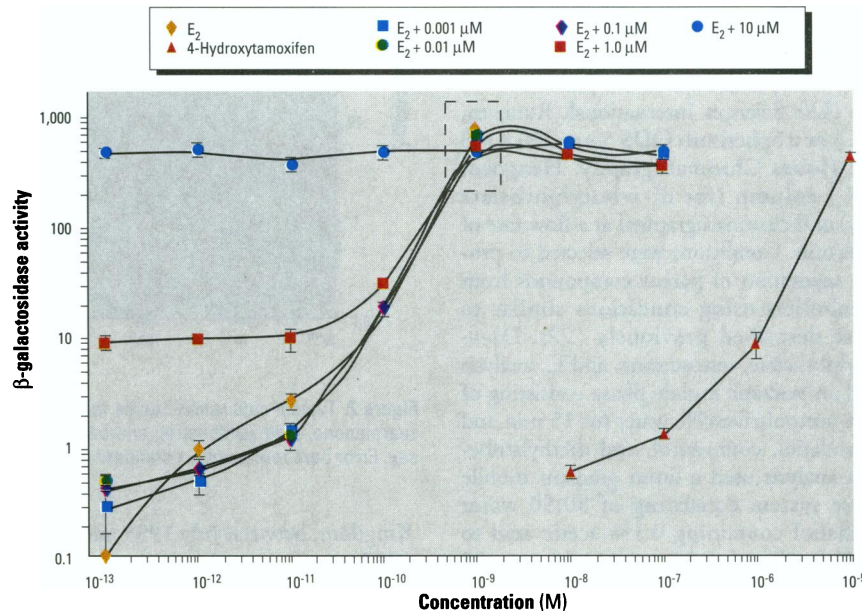


Figure 3. Calibration curves for 17β -estradiol (E_2) 100 fM–100 nM, 4-hydroxytamoxifen 10 pM–10 μ M, and E_2 (100 fM–100 nM) in the presence of 0.001, 0.01, 0.1, 1, and 10 μ M 4-hydroxytamoxifen with the RCBA. Error bars represent 1 standard deviation ($n = 8$). Box indicates data also presented in Figure 4 to illustrate the inhibition of E_2 (1 nM)-induced β -galactosidase activity by 4-hydroxytamoxifen (1 nM–10 μ M).

biological significance through chronic exposure and/or high abundance in the environment. The effect of E_2 on MCF-7 cell proliferation and uterotrophic activity found in the present study was similar to that observed in previous reports (10,11,29). The recombinant yeast cells were designed and engineered for exquisite sensitivity to estrogens (17,30,31); overexpression of human estrogen receptor, high amplitude frog vitellogenin estrogen response elements, and their tandem arrangement in the reporter plasmid all serve to amplify β -galactosidase production and hence sensitivity to E_2 (14). However, evaluation of the relative merits of these bioassays for estrogens cannot be based on sensitivity criteria alone; summary findings for the different assays used in this report and a similar recombinant yeast screen are presented for comparison in Table 3. Adaptation of reporter gene assays to the 96-well plate format enables automation on standard laboratory instrumentation and a marked reduction in sample size. In contrast, the prepubertal mouse uterotrophic assay requires a large mass of sample, is relatively labor intensive, requires the sacrifice of large numbers of animals, and is sensitive to the potentially diverse toxic effects of test compounds.

Interaction of activated receptor–ligand complex with human estrogen responsive elements and the opportunity to measure diverse estrogen-regulated gene products such as the progesterone receptor is a recognized advantage of cancer cell-based screens (24). Uterotrophic bioassays also offer the simultaneous measurement of other markers of estrogen action including vaginal and uterine cell proliferation, which may be used to provide definitive confirmation of estrogenic activity (32). One potential disadvantage of MCF-7 cells is that under certain conditions they may also express receptors for other classes of steroids (33), and other hormones including progestagens have been shown to increase breast cancer cell proliferation (34). Similarly, under certain circumstances uterotrophic assays are limited in selectivity for estrogens by sensitivity to androgens and progestagens (6). For these reasons, expression of single class of steroid receptor, as in the RCBA system, is highly desirable since this provides a system with appropriate sensitivity and specificity to a single class of hormone. Moreover, the genetic simplicity of yeast cells separates estrogen receptor signaling from the confounding affects of other signaling pathways in the cell.

Evaluation of the relative estrogenic potency of steroids from different hormone classes is necessary to demonstrate and estab-

lish assay selectivity for estrogens. This important element of assay validation also provides a reference scale for estrogenic activity found with the RCBA relative to compounds generally accepted to be either estrogens or without estrogenic activity. Many of the compounds tested with the recombinant yeast cell bioassay did not induce β -galactosidase activity to the same extent as E_2 . Similar observations have been made with the E-screen breast cancer cell proliferation assay and were interpreted as indicating that the xenobiotics tested were partial agonists (24). Evaluation of estrogenic activity with the RCBA in terms of relative potency alone can be misleading because several androgens (e.g., testosterone) have estrogenic potencies of a similar order to those of, for instance, the alkylphenols. Consideration must also be given to the relative inductive efficiency of β -galactosidase activity by test compounds. In this regard androgens were much weaker

estrogens than alkylphenols. Although androgens have uterotrophic (6) and mitogenic activity (35) at pharmacological doses *in vivo*, they are unlikely to be of physiological significance as estrogens because endogenous concentrations are several orders of magnitude less than used in such assays. A slight response to testosterone has been reported with a similar yeast screen for estrogens (15); indeed, the potency of testosterone found with the RCBA was an order of magnitude lower than that found with a competitive binding radioreceptor assay (22). Synthetic estrogens including 17 α -ethynylestradiol and the stilbenes were among the most potent estrogens tested comparable with E_2 . This contrasts with the relative estrogenic potency of most of the phytoestrogens and environmental pollutants, which were several orders of magnitude less potent than E_2 . Phytoestrogens have been ranked by potency in a HeLa cell

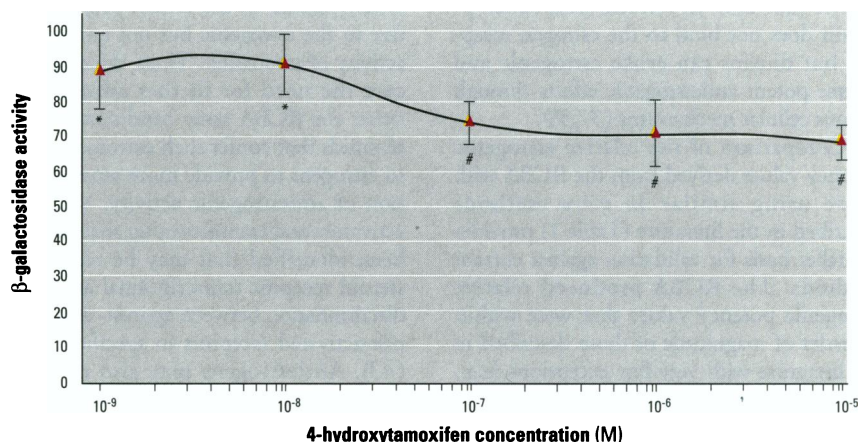


Figure 4. Percentage inhibition of 17 β -estradiol (E_2 , 1nM) induced β -galactosidase activity in recombinant yeast cells by 4-hydroxytamoxifen (1 nM–10 μ M). Error bars represent 1 standard deviation ($n = 8$). * $p < 0.05$; # $p < 0.001$.

Table 3. Comparison of assays used to determine estrogenic activity

Attribute	RCBA	Recombinant yeast screen ^a	MCF-7 breast cancer cells	Mouse uterotrophic assay
Sensitivity to E_2	100 fM	7.3 pM (2 ng/l)	1 pM	18 pmol
ER	Human	Human	Human	Mouse
Estrogen response elements	Frog	Not stated	Human	Mouse
Detection of antiestrogenic activity	Poor	Not tested	Good	Good
Assay endpoints	β -gal	β -gal	Cell proliferation, etc., e.g., PgR	Uterine weight, etc. (e.g., PgR) in other tissues (e.g., vagina)
Steroid receptors expressed	ER	ER	ER, PgR, AR, GR	ER, PgR, AR, GR
Detection of proestrogens	No	Not stated	Limited ?	Yes
Assay duration	18 hr	3–4 days	7 days	4 days
Automation in 96-well plate format	Yes	Yes	Yes	No

Abbreviations: RCBA, recombinant yeast cell bioassay; β -gal, β -galactosidase assay; ER, estrogen receptor; PgR, progesterone receptor; AR, androgen receptor; GR, glucocorticoid receptor.

^aRecombinant yeast screen described by Routledge and Sumpter (15).

cotransfection reporter gene assay (20) and were found to operate as complete agonists in this system. The phthalates and certain organochlorines (DDT, *o*'*p*'-DDT, and DDE) had little estrogenic activity in the recombinant yeast cell bioassay in conflict with other reports (9,36). Various factors including protein binding (16) and molecular permeation into the yeast cells (18), which have not been considered in the present study, cannot be excluded and may provide a partial explanation for these discrepancies. The transformed yeast cells used in the RCBA are a stable cell line and, as such, are robust and well suited for use by other laboratories in further studies, which may reveal the nature of these disparities and further identify strengths and limitations of this assay. Stimulation of β -galactosidase activity by 2,3,7,8-tetrachlorodibenzo-*p*-dioxin was an unexpected but reproducible finding and merits further investigation of the mechanism. Competitive binding studies have shown that 2,3,7,8-tetrachlorodibenzo-*p*-dioxin does not bind to the estrogen receptor, but dioxins can evoke estrogenic and diverse potent antiestrogenic effects through various cellular mechanisms (37,38).

Comparison of the relative estrogenic potency values derived from the RCBA with those using similar *in vitro* methods described in the literature (Table 1) provides a further basis for validation against current methods. The RCBA produced relative estrogenic potency values that were within an order of magnitude of those described in the literature with very few exceptions (e.g., daidzein). Indeed, variation of this order is apparent between literature values (see Table 1) both within and between laboratories according to the particular assay used (20,39). Both the RCBA and mouse uterotrophic assay indicate that 4-nonylphenol is a more potent estrogen than 4-octylphenol, contrary to other reports (24,40), although the potency of 4-nonylphenol obtained with the RCBA was very similar to the E-screen (24). Variation in the estrogenic potency of some 4-*tert*-nonylphenol preparations has been found (15), raising the possibility that the purity of chemicals may provide an explanation for the disparity between assays; alternatively, differences in biotransformation or those inherent to the assay system are plausible explanations. However, this comparative study demonstrates that the RCBA provides estimates of relative estrogenic potency that are reproducible and broadly similar with other *in vitro* methods. In addition to yeast cells, reporter gene assays have also been introduced into breast cancer cells and other cell lines and produced comparable potency values for alkylphenolic compounds (40).

Such reporter gene assays (19,20) and breast cancer cell proliferation assays (10,24) are typically sensitive to E_2 at concentrations of 1 pM (Table 3).

4-Hydroxytamoxifen is a metabolite of tamoxifen and a potent antiestrogen (28). Although the RCBA demonstrates the ability of 4-hydroxytamoxifen to significantly inhibit E_2 -induced β -galactosidase activity, the response was only weak. The incomplete suppression of E_2 -induced β -galactosidase activity observed may be due to the intrinsic estrogenicity of certain triarylethylene antiestrogens like tamoxifen (41). Given the relatively poor response to the potent antiestrogen 4-hydroxytamoxifen, it is anticipated that this system would not have sufficient sensitivity to detect weaker antiestrogens. More significant inhibition of E_2 -induced activity by triarylethylene antiestrogens has been demonstrated in reporter gene assays in mammalian cells (19,20,40) and breast cancer cell proliferation assays (42). A yeast cell system similar to the RCBA (18) was sensitive to the estrogenic but not antiestrogenic activity of tamoxifen. Thus, these data indicate the need for further adjustment of either the RCBA assay conditions or to the plasmids that confer such extreme sensitivity to estrogens to provide more sensitive detection of antiestrogenic activity. Various co-activators and transactivating sequences have been identified that may be required for steroid receptor transcriptional activity and discrimination between agonist- and antagonist-activated receptors in specific cell types (43). Antiestrogens may also operate by other cellular mechanisms including antagonism of calmodulin-regulated processes (42) to which steroid receptor reporter gene systems are likely to be insensitive. Assays with the potential to detect antiestrogens are of particular value because they may be important in evaluating certain classes of compounds such as the phytoestrogens and dioxins, which may have antiestrogenic activity and provide chemoprevention of certain cancers (38,44).

In vivo bioassays for estrogenic activity are of particular value because they provide a means of confirming the potential biological significance of *in vitro* findings (36). The uterotrophic assay used in the present study produced estrogenic potencies that were relatively imprecise because only three doses of test compound were assayed. Nonetheless, with the exception of diethylstilbestrol, lower potency values were derived using the uterotrophic assay compared with the recombinant yeast cell assay. Moreover, several compounds that have estrogenic activity *in vitro*, including the phthalates, 4-octylphenol, and bisphenol A, had no detectable uterotrophic activity. Histological evaluation

of vagina sections enabled further confirmation of estrogenic activity, as in all cases, uterotrophic activity was associated with cornification and thickening of the stratified squamous epithelium. Clearly, the biological significance of very weak estrogens detected with highly sensitive *in vitro* bioassays requires careful evaluation if activity *in vivo* is absent or found only at very high doses, which may exceed those giving rise to acute *in vivo* toxicity. Pharmacokinetic parameters (adsorption, metabolism, tissue distribution, and excretion) may have a significant influence on estrogenic activity *in vivo*, which cannot be readily accounted for in *in vitro* assays. These and other inherent differences between the RCBA and the murine uterotrophic assay will produce different potency values as found for diethylstilbestrol and other chemicals shown in Table 2. Chronic exposure *in vivo* may facilitate the bioaccumulation of pollutants and their metabolites and thereby give rise to estrogenic activity not evident in acute (3 day) uterotrophic studies. The current situation is further complicated by recent conflicting reports (45,46) relating to the potential for substantial synergistic interactions between xenoestrogens.

Biotransformation of xenobiotics *in vivo* is usually part of a process of deactivation and elimination, but proestrogens (e.g., certain phytoestrogens and organochlorines) may undergo metabolic activation to potent estrogens in various body compartments. Although the range of compounds tested for biotransformation in the RCBA was limited, these studies provide valuable information on the fate of test compounds in this type of system. The corollary to the absence of significant steroid and xenobiotic biotransformation by the recombinant yeast cells is that, in their present form, they could not be utilized for the detection of proestrogens. Proestrogen detection *in vitro* has been accomplished by co-incubation of metabolizing and estrogen detecting systems (28,47). In this regard, transformed yeast cells would be suitable for combination with an appropriate metabolizing system for the detection of species-specific proestrogens since there would be little interference from yeast cell xenobiotic metabolism. By contrast, other mammalian cell assay systems such as MCF-7 breast cancer cells will also have the potential for significant steroid and xenobiotic biotransformation; in isolation, such intrinsic metabolism may confound interpretation of results from a simple screen for estrogenic activity. Although cytochrome P450 activities are poorly maintained *in vitro* by isolated cells, steroid metabolizing enzymes, including aromatase, hydroxysteroid oxidoreductases, and estrogen sulfa-

tase, have been described in human breast cancer cells. Thus, a comprehensive knowledge of intrinsic metabolic activity is required before a test system can be fully validated and acceptable as a gold standard method.

Conclusion

The RCBA is a highly sensitive human estrogen receptor-based screening assay for the rapid detection of estrogens. The estrogenic potency and relative inductive efficiency of 53 chemicals has been evaluated with this system. Potency values produced with the RCBA were similar to literature values, but were generally higher than those produced with a murine uterotrophic assay. The mechanism through which TCDD stimulates β -galactosidase activity in the RCBA warrants further investigation. Because the RCBA cannot detect proestrogens and has limited sensitivity to antiestrogens, further development is required to broaden the application to the detection of such compounds. The RCBA may be suitable for use by regulatory bodies, such as the FDA in the United States and the Environment Agency in the United Kingdom, as a simple, rapid, and inexpensive method of evaluating intrinsic estrogenic activity of wide-ranging chemical classes and may provide a preliminary means of evaluating the risk presented by such chemicals to the public, wildlife, and the environment.

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