

Protocol of Retinol Signaling Pathway C3H10T1/2 Cell-based Assay for High-throughput Screening

DOCUMENT: Retinol Signaling Pathway_TOX21_SLP_Version1.0
TITLE: Protocol of Retinol Signaling Pathway C3H10T1/2 Cell-based Assay for High-throughput Screening

ASSAY REFERENCES:

Assay Target	Cell Lines	Species	Tissue of Origin	Assay Readout	Assay Provider	Toxicity Pathway
Retinoic acid receptor	C3H10T1/2	Mouse	Mouse embryo	Luciferase reporter	OARSA/CFSAN/FDA	Retinol Signaling Pathway (RSP)

QUALITY CONTROL PRECAUTIONS:

1. -Maintain cells below 85% confluence
2. -Fetal Bovine serum used for cell culture and assay purpose is heat inactivated at 56 C for 30min
3. -Extra precautions to be taken for making Retinol as it is photosensitive and moisture absorbant

MATERIALS and INSTRUMENTS:

Supplies/Medium/Reagent	Manufacturer	Vender/Catalog Number
-Eagles Basal Medium: BME	-Invitrogen	-Invitrogen/21010
-Fetal Bovine Serum	-ATCC	-ATCC/30-2020
-L-Glutamine	-Invitrogen	-Invitrogen/25030
-Puromycin	-Invitrogen	-Invitrogen/A11138-03
-Penicillin & Streptomycin	-Invitrogen	-Invitrogen/15140
-Recovery Cell culutre Freezing Medium	-Invitrogen	-Invitrogen/12648
-0.05% Trypsin-EDTA	-Invitrogen	-Invitrogen/25300
-Retinol	-Sigma	-Sigma/95144
-ER 50891	-Tocris	-Tocris/3823
-Tetraoctyl ammonium bromide	-Sigma	-Sigma/294136
-1536-well white solid plates	-Greiner Bio-One	-Greiner Bio-One / 789173-F
-MULTIDROP COMBI	-Thermo Electron Corporation	-Thermo Electron Corporation
-BioRAPTR FRD	-Beckman Coulter	-Beckman Coulter
-ViewLux Plate Reader	-Perkin Elmer	-Perkin Elmer
-Amplite Luciferasse reporter gene assay kit	-AAT Bioquest	-AAT Bioquest/12520

PROCEDURE:

1. Cell handling:

1.1. Media Required:

Component	Growth Medium	Assay Medium	Thaw Medium	Freezing Medium
-Eagles Basal Medium: BME	-90%	-90%	-90%	-
-FBS (Heat inactivated)	-10%	-10%	-10%	-
-L-Glutamine	-2 mM	-2 mM	-2 mM	-
-Puromycin	-2 ug/mL	-	-	-
-Penicillin & Streptomycin	-100U/ml & 100ug/ml	-100U/ml & 100ug/ml	-100U/ml & 100ug/ml	-
-Recovery Cell culutre Freezing Medium	-	-	-	-100%

1.2. Thawing method

1.2.1 -Thaw a vial of cells in 9ml of pre-warmed thaw medium and then centrifuge

1.2.2 -Resuspend the pellet with the thaw medium and seed at 2 million cells per T-75 flask

1.3. Propagation method

1.3.1 -Trypsinize cells from the culturing flask and centrifuge and then resuspend cells in culture medium

1.3.2 -Passage cells at 1-1.5 million per T-225 flask

2. Assay Protocol

2.1 -Trypsinize cells from the culturing flask and centrifuge and then resuspend cells in assay medium at a density of 0.25×10^6 cells/mL

2.2 -Dispense 1000 cells/4uL/well into 1536-well tissue treated white/solid bottom plates using a 8 tip dispenser (Multidrop)

2.3 -Incubate the plates for an overnight (20hr) at 37C and 5% CO₂

2.4 -Transfer 23nL of compounds from the library collection (5.6nM to 92uM) and positive control through pintoole

2.5 -Compound transfer was followed by the addition of 1ul of 1uM (final concentration) Retinol (Retinol made fresh from the powder) or assay buffer using two different tips of a Bioraptr

2.6 -Incubate the plates for 6hr at 37C and 5% CO₂

2.7 -Add 5ul of Amplite(TM) Luciferase reagent using a single tip dispense (Bioraptr)

2.8 -Incubate the plates at room temperature for 30min

2.9 -Measure luminescence (exposure time = 60sec) by ViewLux plate reader

3. Assay Performance

RSP	Online Validation Antagonist (Mean ± SD)
IC50	0.03 ± 0.01 μM (n = 27)
S/B	5.48 ± 0.79
CV (%)	13.15 ± 0.69* (n = 18)

Z'	0.42 ± 0.10
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*CV values shown represent average of all assay plates excluding the top 3 compound concentration plates.