

## Predictor<sup>TM</sup> Assay Setup Guide on the Tecan Infinite<sup>®</sup> F500 Microplate Reader

NOTE: The Tecan Infinite<sup>®</sup> F500 Microplate Reader was tested for compatibility with Invitrogen's Predictor<sup>TM</sup> hERG FP Assay (PV5365) using controls provided within the assay kit and the known hERG channel blockers astemizole and terfenadine. The following document is intended to demonstrate setup of this instrument. For more detailed information and technical support of Invitrogen assays please call 1-800-955-6288, select option "3", then extension 40266. For more detailed information and technical support of Tecan instruments or software, please contact Tecan at 1-888-798-0538 or [info@tecan.com](mailto:info@tecan.com).

### A. Recommended Optics

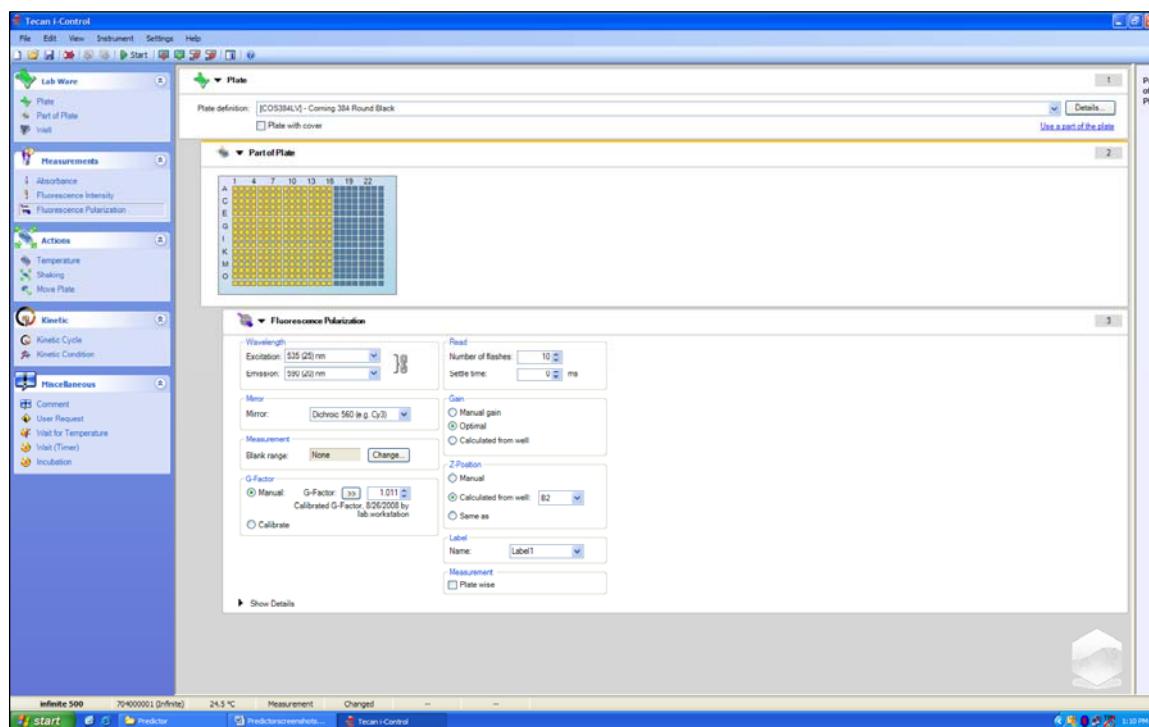
Tecan part number	wavelength (nm)	diameter (mm)
Excitation Filter *contact Tecan	535/25	12.5
Emission Filter 1 *contact Tecan	590/20 (S-pol)	12.5
Emission Filter 2 *contact Tecan	590/20 (P-pol)	12.5
Dichroic Mirror *contact Tecan	560 (Cy3)	

### B. Instrument Setup

1. Make certain plate reader is turned on, and open up Tecan i-Control software on computer.

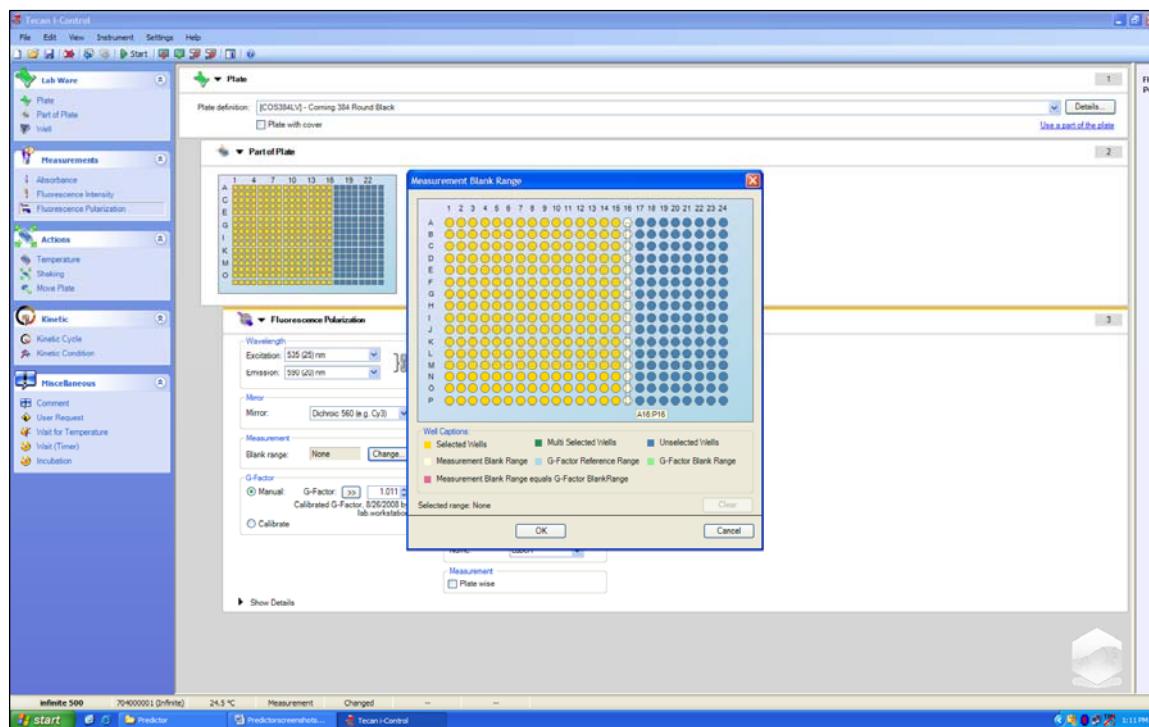
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3. At this point, a secondary settings window will open (below). Select the portion of the plate you wish to read by dragging across with your mouse to highlight selected wells, and in the new Fluorescence Polarization tab below, select your excitation and emission settings and dichroic from the drop-down lists. Select "Optimal" for the Gain, and select a well with tracer in it for Z-positions (well B2 in this example). When finished, under "Measurement" select the "Change..." tab.



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4. A new popup window will appear. Select your Measurement Blank Range wells, in this case Column 16. When finished, select “OK”.

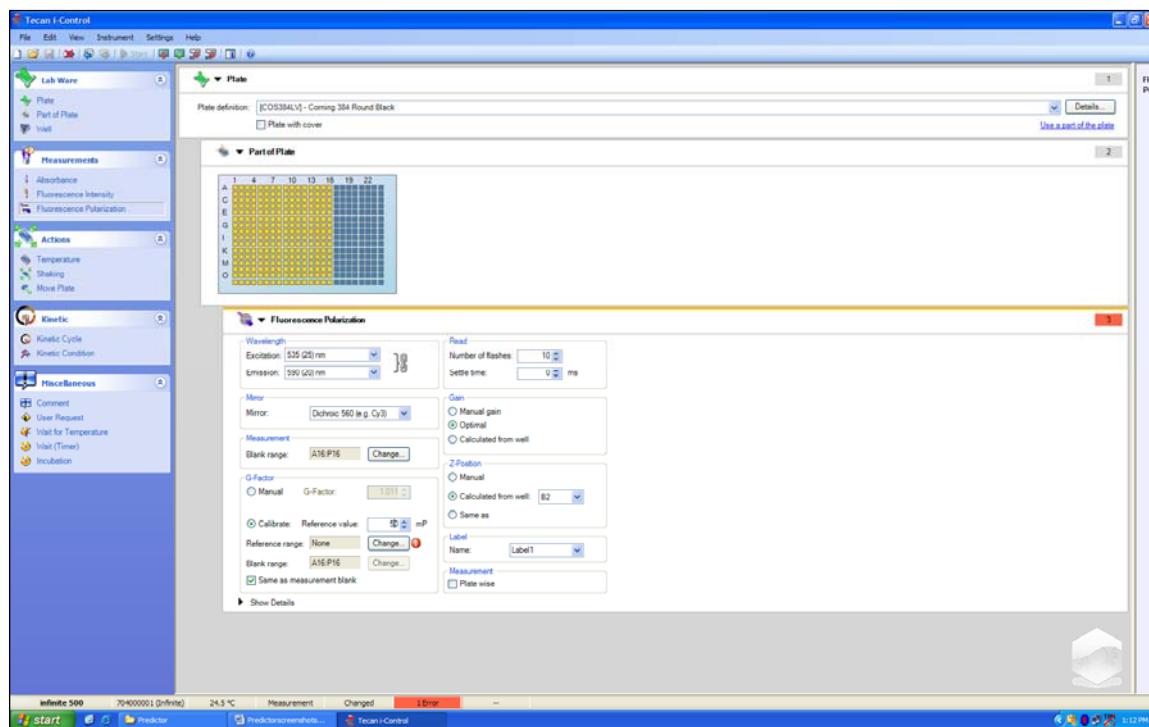


Have a question? Contact our Technical Support Team

NA: 800-955-6288 or INTL: 760-603-7200 Select option 3, ext. 40266 Email: drugdiscoverytech@invitrogen.com

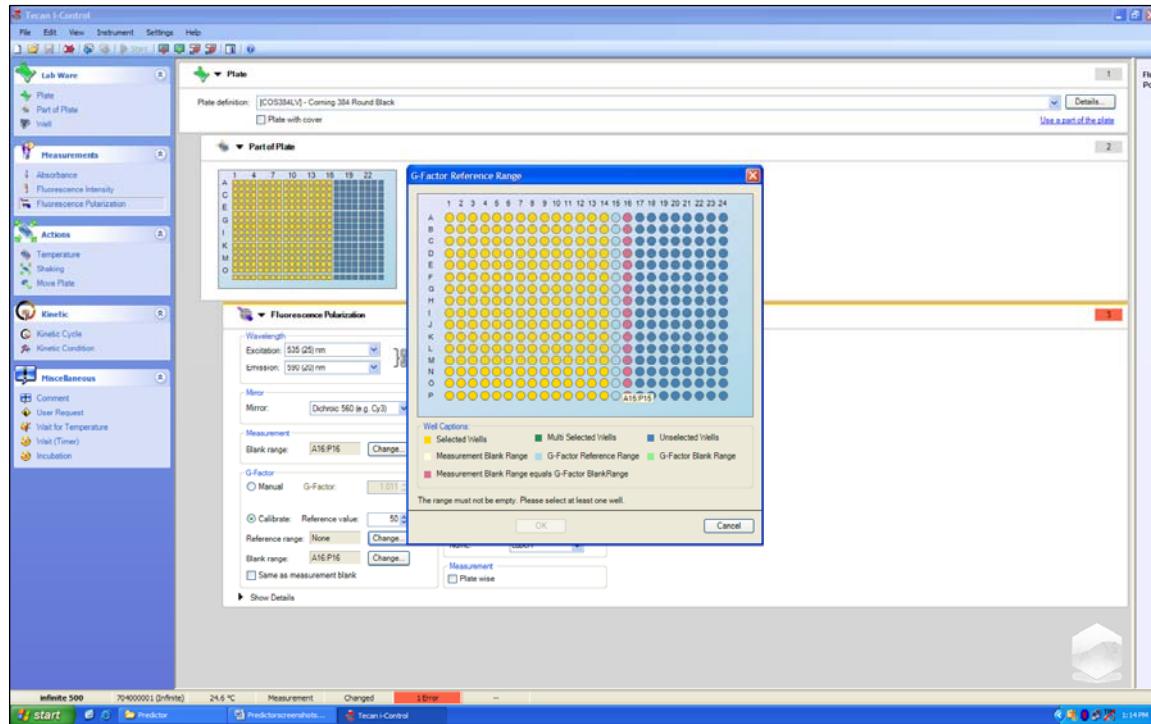
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5. You will return to the main assay setup window. Under “G-Factor”, select the “Calibrate” button, then select the “Change” tab.



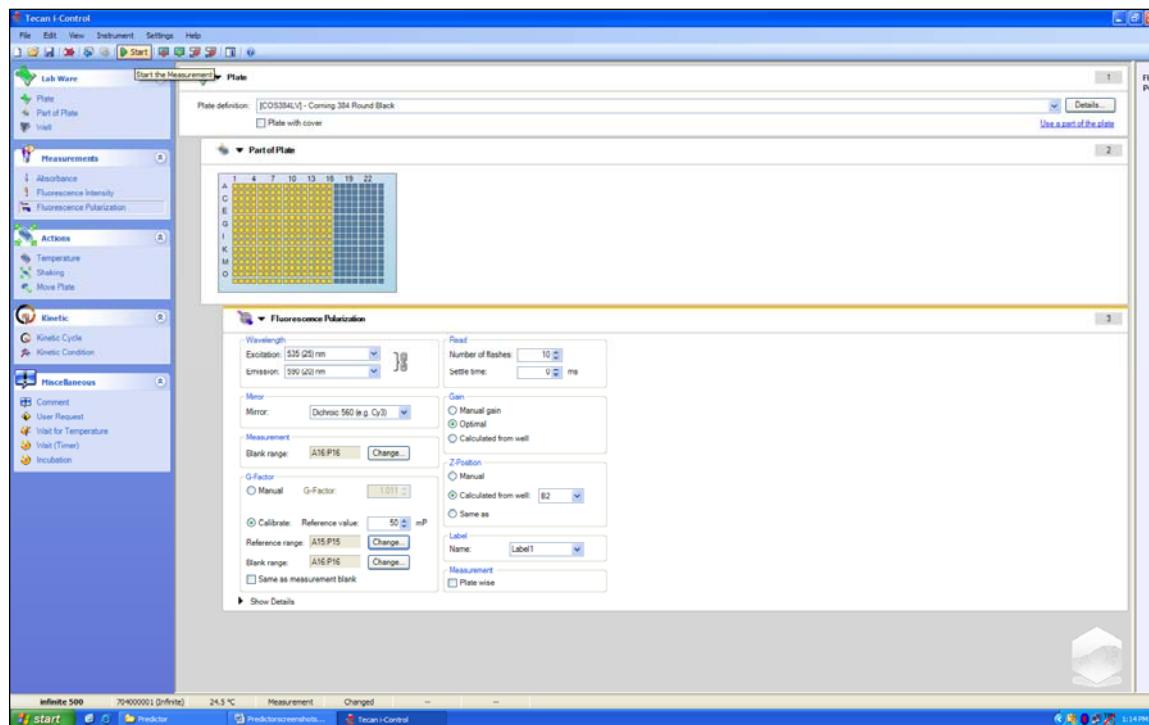
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6. Once again, a popup window will appear. Select the G-Factor Reference Range (the Free Tracer wells in Column 15 in this case). Select “OK” when finished.



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7. You will return to the main setup window a final time. Under “G-Factor” set the Reference Value to 50mP to set the mP value for free tracer. When finished, select “Start” from the toolbar at the top to begin reading your plate.



## C. Predictor™ hERG FP Assay

NOTE: The following is a sample assay performed for demonstration purposes. The instrument settings above would be sufficient for any Predictor™ assay or other Invitrogen red FP assay, the information below is provided as representative data. Assays were run in 20  $\mu$ l in 384-well untreated low-volume polystyrene plates (Corning #3677). In order to demonstrate the correction of Polarization Interference at high concentrations of compound described in the Predictor protocol Section 4.5, we prepared dose-response curves of both astemizole and terfenadine with and without saturating E-4031. Also, all FP data was background-subtracted using wells containing membranes but no tracer. Note background subtraction is not required, but is strongly advised and can increase overall assay window by 10-20%.

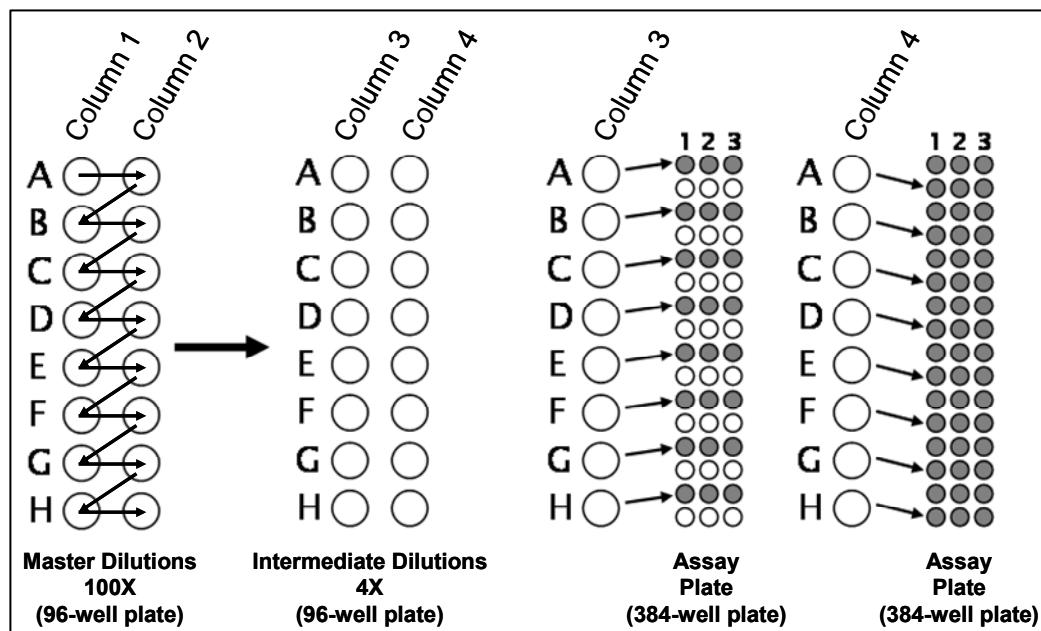
1. Thaw all reagents as directed in protocol. Thaw compounds (Astemizole and Terfenadine, 1 mM stocks in DMSO) at room temp.
2. Prepare Predictor™ hERG membrane by sonification, dounce homogenization, or trituration to ensure a uniform suspension with no precipitate or aggregates.
3. A dilution series of both astemizole and terfenadine was prepared as follows:
  - 3.1. 1:10 pre-dilution: In a 96-well plate, add 40  $\mu$ l of DMSO to wells B1-H1 and A2-H2 (for Astemizole pre-dilution) and B7-H7 and A8-H8 (for Terfenadine pre-dilution).
  - 3.2. Add 40  $\mu$ l of 1 mM astemizole to well A1 and 40  $\mu$ l of 1 mM terfenadine to well A7.
  - 3.3. Prepare Master Dilutions: Transfer 20  $\mu$ l from well A1 to well A2, mix by pipetting up and down several times, then transfer 20  $\mu$ l from well A2 to well B1, mix again, then transfer 20  $\mu$ l from well B1 mixed into well B2, mix again, then transfer 20  $\mu$ l from well B2 to well C1, repeating this pattern downwards to the bottom of plate (see Figure 1). Repeat this same procedure with terfenadine starting in well A7,
  - 3.4. From well A1, remove 4  $\mu$ l compound in DMSO and add to well A3 in column 3. From well A2, remove 4  $\mu$ l compound in DMSO and add to well A4 in column 4. Repeat to bottom of plate, transferring 4  $\mu$ l aliquots from initial dilution to new column. This step can be simplified by use of a multichannel pipette.

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3.5. Repeat for terfenadine, adding 4  $\mu$ l to columns 9 and 10. When finished, add 96  $\mu$ l Predictor™ hERG FP Assay Buffer to each new well to create the intermediate dilution series.

4. Add 5  $\mu$ l compound per well to a nontreated Corning 3677 black polystyrene 384-well plate, as follows:

4.1. From well A3 of the 96-well plate (maximum astemizole, conc. 40  $\mu$ M) transfer 6 replicates of 5  $\mu$ l compound to wells A1-A6 of the 384 assay plate. In the end, column 3 of the 96-well plate will fill the first 6 wells in rows A, C, E, G, I, K, M, and O while column 4 will fill the first 6 wells of rows B, D, F, H, J, L, N, and P. Repeat same procedure was repeated for the next 6 wells in each row on the 384-well assay plate using columns 9 and 10. This step can be simplified by use of a multichannel pipette.



**Figure 1: Schematic for Preparation of Compound Dilution.** Schematic demonstrates progression of dilutions and final transfer to assay plate. Note in this case 2 compounds were titrated and each compound was finally used in replicates of 6 (3 with and 3 without saturating E-4031).

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5. Once compound titration wells are prepared (columns 1-12 of assay plate), also prepare the following controls:
  - **Negative Control, Tracer fully Bound (Column 13, B).** Add 5  $\mu$ l Predictor<sup>TM</sup> buffer containing 4% DMSO to all wells in Column 13 (to control for DMSO in titration wells). Tracer and membrane will be added later. This control will show full tracer binding activity of assay.
  - **Positive control, Tracer Displaced (Column 14, D).** Add 8  $\mu$ l E-4031 to 192  $\mu$ l Predictor<sup>TM</sup> hERG FP Assay Buffer also containing 4% DMSO. Add 5  $\mu$ l per well of this solution to each well of Column 14 of assay plate. Tracer and membrane will be added later. This control will show maximum displacement of tracer from the hERG channels in the membrane prep.
  - **Free Tracer Control (Column 15, F).** Add 15  $\mu$ l Predictor<sup>TM</sup> hERG FP Assay Buffer to each well of column 15. Tracer will be added later. This control will show fully unbound tracer in solution, as well as being used to calibrate the instrument.
  - **Assay Blank (Column 16, Blank).** Add 10  $\mu$ l of Predictor<sup>TM</sup> hERG FP Assay Buffer per well to Column 16. 10  $\mu$ l membrane will be added later. This control will show background fluorescence and to background subtract the rest of the assay wells prior to calculating FP values.
6. Sonicated/triturated membrane split: remove 2 ml Predictor membrane and add 40  $\mu$ l E-4031 to these membranes.
7. Add 10  $\mu$ l E-4031-containing membrane per well to columns 4-6 (astemizole with E-4031 baseline subtraction) and 10-12 (terfenadine with E-4031 baseline subtraction).
8. Add 10  $\mu$ l untreated membrane per well to columns 1-3 (astemizole), 7-9 (terfenadine), 13 (Bound Tracer), 14 (Displaced Tracer), and 16 (Blank).
9. Tracer added last: Add 32  $\mu$ l tracer to 1968  $\mu$ l Predictor hERG FP Assay Buffer and add 5  $\mu$ l/well of this solution to all wells in Columns 1-15.

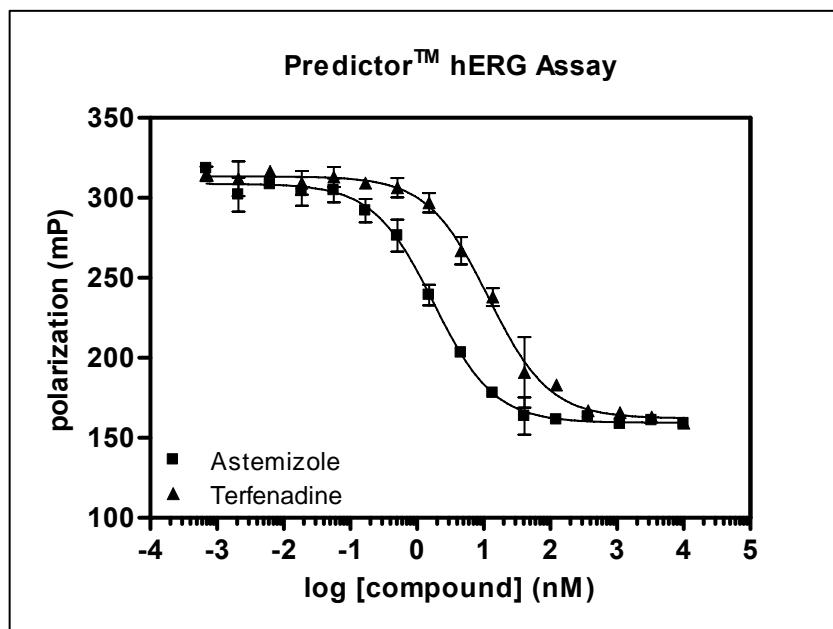
**NOTE:** Predictor<sup>TM</sup> is a highly sensitive assay. Furthermore, the baseline correction wells use saturating levels of E-4031. **It is extremely important to ensure that pipetting steps are done carefully and there is no carryover/contamination of neighboring wells in Predictor.** The order of wells in the plate layout may be changed to best facilitate your pipetting and minimizing carryover if desired, but all the controls above are highly recommended.

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10. Shake plate on an orbital plate shaker for 30 seconds, cover with foil, and incubate 2 hours before reading.
11. Read plate on plate reader as outlined above.

	1	2	3	4	5	6	7	8	9	10	11	12	B	D	F	Blank	17	18	19	20	21	22	23	24
A																								
B																								
C																								
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**Figure 2: Schematic of Assay Plate Layout.** Predictor™ hERG assay was set up with a dose-response curve of Astemizole and Terfenadine (1:3 dilution series from a starting concentration in the top row of 10  $\mu$ M) prepared in replicates of 6. From this, 3 replicates were assayed for ability to disrupt Predictor™ Tracer binding and 3 were assayed in saturating levels of E-4031 in order to prepare a baseline correction for each data point.

**D. Results:**

**Figure 3: Predictor™ hERG FP Assay.** Dose-Response Curves read on the Tecan Infinite® F500 using the Predictor™ assay and 1:3 dilution series prepared for Astemizole and Terfenadine from a starting concentration of 10  $\mu$ M. Curve calculations were baseline-corrected against duplicate dilution series prepared in saturating E-4031.

	avg	std dev
<b>free tracer</b>	54.13	4.05
<b>no inhibitor</b>	312.85	8.65
<b>30 <math>\mu</math>M E-4031</b>	159.13	3.04
<b><math>\Delta</math>mP</b>	153.72	
<b>Z'-factor</b>	0.77	
<b>Astemizole IC50</b>	1.79 nM	
<b>Terfenadine IC50</b>	11.78 nM	

**Table 1. Predictor™ Assay Results on the Tecan Infinite® F500.**