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A single amino acid difference between *ether-a-go-go*-related gene channel subtypes determines differential sensitivity to a small molecule activator

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ABBREVIATIONS: ERG, *ether-a-go-go*-related gene; hERG1, human *ether-a-go-go*-related gene 1; I_{test} , current activated by membrane depolarization; I_{tail} , tail current; I-V, current-voltage; LQTS, long QT syndrome; rERG3, rat *ether-a-go-go*-related gene 3; RPR, (3R,4R)-4-[3-(6-methoxy-quinolin-4-yl)-3-oxo-propyl]-1-[3-(2,3,5-trifluoro-phenyl)-prop-2-ynyl]-piperidine-3-carboxylic acid; V_t , test potential; $V_{0.5}$, half-point of Boltzmann relationship; WT, wild-type

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Abstract

Activators of human *ether-a-go-go*-related gene 1 (hERG1) channels such as RPR260243 ((3R,4R)-4-[3-(6-methoxy-quinolin-4-yl)-3-oxo-propyl]-1-[3-(2,3,5-trifluoro-phenyl)-prop-2-ynyl]-piperidine-3-carboxylic acid) reverse the effect of hERG1 blockers and shorten the duration of cardiac action potentials. RPR260243 (RPR) slows the rate of deactivation and shifts the voltage dependence of channel inactivation to more positive potentials. We recently mapped the binding site for RPR to several residues located near the cytoplasmic ends of the S5 and S6 helices of the hERG1 subunit. These residues are conserved in the highly homologous *ether-a-go-go*-related gene 3 (ERG3) subunit; however, RPR blocks ERG3 channels. Here, we compare hERG1 and rat ERG3 (rERG3) channels to explore the molecular basis for differential channel sensitivity to RPR. Channels were heterologously expressed in *Xenopus* oocytes and currents were recorded using the two-electrode voltage clamp technique. Site-directed mutagenesis was used to swap the two residues within the putative binding domain that differed between hERG1 and rERG3. The differential sensitivity of hERG1 and rERG3 channels to the agonist effect of RPR could be accounted for by a single S5 residue (Thr556 in hERG1, Ile558 in rERG3). A Thr in this position favors agonist activity, whereas an Ile reveals a secondary blocking effect of RPR.

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Introduction

Ether-a-go-go-related gene (ERG) K⁺ channels are activated by voltage and exhibit inward rectification properties owing to a rapid voltage-dependent P-type inactivation process. Three ERG subfamily genes have been identified (Shi et al., 1997; Warmke and Ganetzky, 1994). In mammals, ERG1, ERG2 and ERG3 (Kv11.1-11.3) K⁺ channels share a high degree of amino acid sequence homology, although each differs subtly with respect to their voltage-dependence of activation and inactivation. ERG2 and ERG3 channels are expressed exclusively within the nervous system where they may play a role in the maintenance of resting membrane potential, control of excitability and frequency adaptation (Schwarz and Bauer, 2004). In contrast, ERG1 channels are expressed in many cell types, including cardiac myocytes where they conduct the rapid delayed rectifier K⁺ current (I_{Kr}) that mediates action potential repolarization (Sanguinetti et al., 1995; Trudeau et al., 1995).

Loss-of-function mutations in human ERG1 (hERG1) cause inherited long QT syndrome (LQTS) (Curran et al., 1995), a disorder characterized by delayed ventricular repolarization and a prolonged QT interval of the body surface electrocardiogram. Acquired LQTS is more common and is most often caused by unintended block of hERG1 channels by a plethora of common medications. Both forms of LQTS are associated with an increased risk of torsades de pointes, a ventricular arrhythmia that can degenerate into fibrillation and cause sudden death. Currently, the only drugs used commonly to treat LQTS are β -adrenergic receptor blockers (Schwartz, 2006). Theoretically, activators of hERG1 could provide an alternative and more specific pharmacological treatment for acquired or inherited LQTS.

Several compounds were recently identified that activate hERG1 channels and shorten cardiac action potentials. NS1643 (Casis et al., 2006; Hansen et al., 2005) and PD-118057 (Zhou et al., 2005) increase current magnitude without much effect on the rate of channel deactivation. RPR260243 (RPR) and primarily slows the rate of hERG1 deactivation (Kang et al., 2005), but also slows activation and enhances current magnitude by attenuation of P-type inactivation (Perry et al., 2007). PD-307243 affects hERG1 channels in a manner similar to RPR

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(Gordon et al., 2007). Site-directed mutagenesis and functional analysis of mutant channels suggests that RPR binds to several residues located near the cytoplasmic ends of the S5 and S6 helices of the hERG1 subunit (Perry et al., 2007). The amino acids that form the putative RPR binding site in hERG1 are conserved in hERG3 channels. However, RPR is not an activator of hERG3 channels and in fact, was reported to block these channels when heterologously expressed in CHO cells (Kang et al., 2005). Here, we compare the effects of RPR on hERG1 and rat ERG3 (rERG3) channels and explore the molecular basis for their differential sensitivity to this drug.

Materials and Methods

Molecular biology. cDNAs for wild-type (WT) human *ERG1* isoform 1a (GenBank accession number NM_000238) and rat *ERG3* (kindly provided by David McKinnon, GenBank accession number AF_016191) were cloned into the pSP64 oocyte expression vector. Sequence alignments were produced using Clustal W (www.ebi.ac.uk/clustalw/). Site-directed mutagenesis was performed with QuikChange (Stratagene, La Jolla, CA). S5-P linker (S5-PL) chimeras were constructed by using a 2-step domain swapping method (Kirsch and Joly, 1998). Oligonucleotide primers were designed to anneal and prime to the S5-PL of both hERG1 and rERG3 (Forward primer was 5', CCC TGA TTG CCC ACT GGC TGG CCT GCA TCT GGT ACG CCA TC, 3', Reverse primer was 5', GTA AAG TAA AGT GCC GTG ACA TAC TTG TCC TTG ATG GAG GG, 3'). PCR was performed to amplify only the S5-PL fragment of hERG1. After purification of this fragment, a second amplification of *rERG3* was performed using the amplified S5-PL DNA as megaprimers. cRNA was prepared by *in vitro* transcription with SP6 Cap-Scribe (Roche, Indianapolis, IN) after linearization of the vector plasmid with either *EcoR1* (for WT and mutant *hERG1*) or *AflIII* (for WT and mutant *rERG3*).

Voltage clamp of *Xenopus* oocytes. The isolation, culture and injection with cRNA of oocytes isolated from *Xenopus laevis* frogs were performed as described (Sanguinetti et al., 1995). Currents were recorded from oocytes 1-5 days after cRNA injection using the two-

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electrode voltage clamp technique as described (Stühmer, 1992). Agarose cushion electrodes were fabricated by filling pipette tips with 1% agarose dissolved in 3M KCl and then back-filling with 3M KCl (Schreibmayer et al., 1994).

Oocytes were voltage clamped to a holding potential of -90 mV and 5-s pulses to 0 mV were applied every 8 or 15 s until current magnitude reached a steady-state level. To determine standard current-voltage (I-V) relationships, test currents (I_{test}) were elicited with 2-s pulses applied from a negative holding potential (-90 or -110 mV). Pulses were applied in 10-mV increments to test potentials (V_t) that ranged from -80 to +50 mV. Each test pulse was followed by a 2-s repolarizing step to -70 mV to elicit tail currents (I_{tail}). The rate of I_{tail} decay at -70 mV was determined by fitting current traces to a two exponential function (Eq. 1) to obtain the fast and slow time constants (τ_f , τ_s) for deactivation and relative amplitude of the slow component of deactivation, $A_s/(A_s + A_f)$.

$$\text{Eq. 1} \quad I_{\text{tail}}(t) = A_f \exp^{-t/\tau_f} + A_s \exp^{-t/\tau_s} + C$$

The voltage dependence of hERG1 and rERG3 channel activation was determined by analysis of peak I_{tail} . The plot of normalized I_{tail} amplitude (I_n) versus V_t was fitted with a Boltzmann function (Eq. 2) to obtain the half-point ($V_{0.5}$) and slope factor (k) for channel activation.

$$\text{Eq. 2} \quad I_n = 1/(1 + \exp[(V_{0.5} - V_t)/k])$$

The fully activated I-V relationship was determined by applying 2-s prepulses to +40 mV, followed by measuring tail currents ($I_{\text{tail, FA}}$) upon repolarization to a variable V_t (-140 to +30 mV). The extent of inward rectification was quantified by the deviation of $I_{\text{tail, FA}}$ values from that predicted by linear extrapolation of $I_{\text{tail, FA}}$ measured at a V_t between -140 and -110 mV to more positive potentials. The deviation of the I-V relationships from linearity was corrected by the

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driving force for K^+ ($V_t - E_{rev}$) to obtain a rectification factor for each value of V_t . Finally, the plot of rectification factor versus V_t was fitted with a Boltzmann function to estimate the voltage dependence for inactivation (Sanguinetti et al., 1995).

After addition of RPR to the bathing solution, 5-s pulses to 0 mV were applied every 60 s for ~15 min. When a new steady state level of I_{test} was achieved, the voltage protocols described above were repeated. Each oocyte was treated with a single concentration of RPR. In the absence of drug, the increase in I_{test} for WT hERG1 during an equivalent 15 min period after switching between control solutions in the absence and presence of vehicle (0.03% dimethyl sulfoxide, DMSO) was $6 \pm 6\%$ ($n = 5$).

Data analysis. Digitized data were analyzed off-line with pCLAMP9 (Molecular Devices Corp., Sunnyvale, CA) and ORIGIN 7.5 (OriginLab Corp., Northhampton, MA) software. Results are reported as mean \pm SEM (n = number of oocytes). Statistical differences between control and drug treatment were evaluated by a Student's paired t-test. Statistical significance was assumed for $P < 0.05$.

Solutions and drugs. The extracellular solution contained (in mM): 96 NaCl, 2 KCl, 1 CaCl₂, 2 MgCl₂ and 5 Hepes, pH adjusted to 7.6 with NaOH. RPR260243 ((3R,4R)-4-[3-(6-methoxy-quinolin-4-yl)-3-oxo-propyl]-1-[3-(2,3,5-trifluoro-phenyl)-prop-2-ynyl]-piperidine-3-carboxylic acid) was kindly provided by David Rampe of Sanofi-Aventis Pharmaceuticals (Bridgewater, NJ). Drug solutions were prepared daily by dilution of a 10 mM DMSO stock solution.

Results

RPR enhances magnitude and slows deactivation of hERG1, but not rERG3 channel currents. As we previously reported (Perry et al., 2007), RPR (3 μ M) enhanced the magnitude and slowed the rate of activation and deactivation of WT hERG1 channel currents in *Xenopus* oocytes (Fig. 1A). In contrast, the same concentration of RPR had almost no effect on rERG3 currents elicited with the same pulse protocol (Fig. 1B, Tables 1 & 2). The effect of 3 μ M RPR

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on rERG3 current over an extended range of voltage was assessed with a fully activated I-V protocol in which peak tail currents ($I_{tail, FA}$) were measured upon repolarization to a variable V_t after a 2-s prepulse to +40 mV. RPR failed to enhance $I_{tail, FA}$ magnitude over this voltage range (Fig. 1C, Table 2). The deviation of the fully-activated I-V relationship from linearity was then used to estimate the voltage dependence for rERG3 inactivation. The $V_{0.5}$ for inactivation was not altered by 3 μ M RPR (Fig. 1D, Table 2). At 3 μ M and 10 μ M, RPR slightly slowed the rate of deactivation of rERG3, an effect that was more obvious with 30 μ M RPR (Fig. 1, E and F). However, the effect of 30 μ M RPR on rERG3 channel deactivation was much smaller than that observed for hERG1 channels exposed to 1 μ M RPR, suggesting at least a 30-fold difference in drug sensitivity. Similar to that previously reported for hERG3 expressed in CHO cells (Kang et al., 2005), 30 μ M RPR was an antagonist of rERG3, reducing current at 0 mV by $18 \pm 5\%$ ($n = 3$, Fig. 1F and G). The shift in $V_{0.5}$ for inactivation ($\Delta V_{0.5}$) induced by 30 μ M RPR was $+4.8 \pm 0.3$ mV, significantly greater than for the vehicle (0.3% DMSO) control ($\Delta V_{0.5} = +1.2 \pm 0.7$ mV, $n = 3$). However, for hERG1 channels, 30 μ M RPR shifted $V_{0.5}$ for inactivation by +37 mV and enhanced $I_{tail, FA}$ 3-fold (Perry et al., 2007). Thus, whereas RPR produced a concentration-dependent and marked activation of hERG1, rERG3 channels were insensitive (3 μ M) or blocked (30 μ M) by RPR.

A single S5 residue determines hERG1 and rERG3 sensitivity to RPR. We recently mapped the putative binding site for RPR to residues located within the cytoplasmic ends of the S5 and S6 helices of the hERG1 subunit (Perry et al., 2007). We hypothesized that amino acid differences within or nearby this binding site could account for the reduced sensitivity of rERG3 to RPR. Sequences from the C-terminal end of S4 to the end of the S6 for hERG1 and rERG3 subunits are aligned in Fig. 2, with non-conserved residues highlighted in bold text and the important RPR binding site residues in hERG1 denoted by asterisks. The C-terminal end of the S5 helix, together with the region that connects the S5 helix to the P-helix (S5-P linker), exhibit the most variation with differences in 16 residues. Only two non-conserved residues exist within the putative RPR binding domain: Phe (hERG1) to Met (rERG3) in the S4-S5 linker and Thr

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(hERG1) to Ile (rERG3) near the cytoplasmic end of the S5 helix. We therefore exchanged these two residues between hERG1 and rERG3 and examined the sensitivity of the altered channels to 3 μ M RPR.

First we examined the single variant residue of the S4-S5 linker. Residue Phe551 of hERG1 was mutated to Met, the equivalent residue in rERG3. RPR (3 μ M) enhanced F551M hERG1 current magnitude by an average of $64 \pm 17\%$ at 0 mV (Fig. 3A, Table 2) and caused a pronounced slowing of tail current deactivation (Fig. 3B, Table 1). RPR also reduced the extent of rectification of the fully-activated I-V relationship (Fig. 3C) and shifted the voltage dependence of inactivation by 11 ± 1 mV ($n = 5$, Fig. 3D). All these effects of RPR were reversible upon wash-out and were not significantly different from those observed for WT hERG1 (Tables 1 and 2). Similar to WT hERG (Perry et al., 2007), the rate of onset of, and recovery from, RPR action on F551M hERG channels was slow ($\tau_{on} = 156$ s; $\tau_{recov} = 196$ s; $n = 5$). The reverse mutation, Met553 to Phe in rERG3 produced a mutant channel which had only a very weak response to 3 μ M RPR (Fig. 3, E-H), similar to that observed for WT rERG3 channels (Fig. 1B). Taken together, these results indicate that the equivalent residues Phe551 in hERG1 and Met553 in rERG3 are not important determinants of RPR sensitivity.

Next we examined the single variant residue in the S5 helix. Thr556 of hERG1 was mutated to Ile, the equivalent residue in rERG3. Unlike WT hERG1, T556I hERG1 did not exhibit any appreciable change in current magnitude, rate of current deactivation or voltage dependence of inactivation in response to 3 μ M RPR (Fig. 4A-D). Thus, T556I hERG1 resembled WT rERG3 with respect to RPR sensitivity. To determine if WT hERG1-like sensitivity to RPR could be engineered into rERG3, we produced the reverse mutation in which Ile558 in rERG3 was substituted with a Thr. RPR (3 μ M) slowed the rate of I558T rERG3 deactivation to a similar extent as WT hERG1, but caused a smaller increase in current magnitude and smaller shift ($\Delta V_{0.5} = 4 \pm 1$ mV) in the voltage dependence of inactivation (Fig. 4 E-H, Table 2). At 30 μ M, RPR greatly slowed the rate of I558T rERG3 activation and deactivation (Fig. 4, I and J) to a similar extent as in hERG1 for this concentration (Perry et al.,

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2007). However, the enhanced current magnitude ($92 \pm 39\%$, Fig. 4K) and shift in the voltage dependence of inactivation ($\Delta V_{0.5} = +18 \pm 1$ mV, Fig. 4L) was more comparable to 3 μ M RPR on hERG1 (Table 2).

The double mutant, M553F/I558T rERG3, was created and tested for RPR sensitivity. Fig. 5 demonstrates that the response of this double mutant to 3 μ M RPR was almost identical to I558T rERG3 (Table 2), further confirming that Phe551 is not an important residue for RPR sensitivity. Thus, the I558T mutation in rERG3 endowed sensitivity to RPR that rivaled hERG1 with regards to slowed activation and deactivation, but the mutation did not as effectively enhance the drug's effect on inactivation gating.

The S5-P linker is not directly involved in conferring sensitivity to RPR. A

A comparison of hERG1 and rERG3 amino acid sequences indicates 16 non-conserved residues in the extracellular end of the S5 helix and the S5-P linker (Fig. 2). Given that the S5-P linker has a proposed role in hERG1 inactivation (Liu et al., 2002), we hypothesized that it might also be responsible for the observed differences in the ability of RPR to attenuate inactivation of I558T rERG3 to the same extent observed for WT hERG1 channels. To investigate this further, we constructed a chimeric channel (rERG3-S5PL1) in which the extracellular end of the S5 helix and the entire S5-P linker (boxed region in Fig. 2) of rERG3 was exchanged for the equivalent region of hERG1. Similar to that observed for WT rERG3 channels (Fig. 1C-F), the chimeric channel was relatively insensitive to 3 μ M RPR (Fig. 6A-D). RPR produced a small, but statistically significant increase ($12 \pm 4\%$, $n = 6$) in the magnitude, and a small shift in the voltage-dependence of inactivation ($\Delta V_{0.5} = +4 \pm 1$ mV, $n = 6$) of rERG3-S5PL1 channel current (Table 2). Introduction of the mutation I558T into rERG3-S5PL1 did not enhance drug sensitivity beyond that observed with I558T rERG3 (compare Fig. 4E-H with 6E-H, Tables 1 and 2). Thus, differences in the sequence of the S5-P linker cannot explain how RPR attenuates inactivation of hERG1 more than rERG3.

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Discussion

RPR activates hERG1 by dramatically slowing the rate of channel deactivation (Kang et al., 2005) and shifting the voltage dependence of inactivation to more positive potentials (Perry et al., 2007). It is conceivable that two distinct binding sites on the channel protein could mediate the deactivation and inactivation effects of RPR. Functional analysis and drug sensitivity of mutant channels was utilized to locate the putative binding site for RPR to a specific region of the hERG1 subunit (Perry et al., 2007). Point mutations of two residues in S5 (L553 and F557) and two residues in an adjacent region of S6 (N658, V659) attenuated both the deactivation and inactivation effects of RPR. In a homology model of the hERG1 pore region (Fig. 7), it is apparent that L553, F557, N658 and V659 are located close together, forming a cluster of residues (colored red) that we proposed constitutes a putative binding site for RPR (Perry et al., 2007). However, this interpretation is muddled by analysis of the effects of RPR on other mutant channels. Point mutations of two residues in the S4-S5 linker (V549, L550) and three residues in an adjacent region of S6 (I662, L666, Y667) prevented the slowing of deactivation but did not alter the shift in the voltage dependence of inactivation by RPR. These 5 residues form a second cluster (colored blue in Fig. 7) that either defines a second binding site for RPR or a region of allosteric modulation that mediates the effects of RPR on activation/deactivation gating. In either model, binding of RPR to hERG1 presumably slows channel deactivation by interfering with normal electromechanical coupling, the bending of the S6 helices towards the central axis of the pore in response to repositioning of the S4 voltage sensors and S4-S5 linkers (Long et al., 2005). P-type inactivation of K⁺ channels involves more subtle conformational changes, specifically within the selectivity filter (Loots and Isacoff, 1998), a structure that is distant from the proposed binding site for the drug. Thus, binding of RPR alter inactivation gating by a long-range allosteric effect.

The critical residues in hERG1 required for RPR activity are also present in its close homologs hERG3 and rERG3. However, RPR acts as an antagonist of hERG3 channels that

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are heterologously expressed in CHO cells (Kang et al., 2005) and rERG3 channels in oocytes (Fig. 1). We employed a cross-mutational approach between hERG1 and rERG3 to examine the molecular basis for this difference and identified a single Thr residue within the cytoplasmic end of the S5 helix that confers the differential sensitivity to RPR. In our previous Ala scan of hERG1, the T556A mutation reduced, but did not eliminate the effect of 3 μ M RPR. For this reason, it seems unlikely that Thr556 (shown in space fill mode in Fig. 7) contributes to the binding site for RPR, but instead mutations at this site may alter RPR binding by steric hindrance or by acting allosterically on nearby binding residues. Based on the functional consequences of mutation, Phe557 is a key component of the binding site for RPR. Thus, mutation of Thr556 to the larger and more hydrophobic Ile might interfere with interaction of RPR with Phe557.

PD-307243 affects hERG1 channels in a manner similar to RPR and based on simulated docking, it was proposed that this compound binds to residues located on the extracellular side of the channel (Gordon et al., 2007). However, the docking was restricted to the extracellular portion of the hERG1 channel and was not supported by any mutagenesis data. In contrast, the slow onset and recovery from RPR effects and our present and previous (Perry et al., 2007) mutational analyses indicate that RPR accesses its binding site from the cytoplasmic side of the membrane.

RPR decreases the magnitude of rERG3 channel current. A similar antagonist effect of RPR on hERG1 channels may be masked by its more marked agonist effect. A dual activity on hERG1 was previously described for NS1643, where the blocking effect was mediated by interaction with the well-described binding site located within the central cavity of the channel (Casis et al., 2006).

RPR slowed the deactivation rate of I558T rERG3 channels with a potency similar to hERG1. However, the rightward shift in the voltage dependence of inactivation caused by 3 μ M RPR was much smaller for I558T rERG3 channels (~4 mV) than for hERG1 channels (~13 mV). Several studies have suggested a role for the S5-P linker in the inactivation process of ERG

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channels based on the finding that specific mutations in this region perturb inactivation (Liu et al., 2002). Since the S5-P linker regions of hERG1 and rERG3 channels contain several non-conserved residues, we hypothesized that the S5-P linker of hERG1 could be required for complete RPR attenuated inactivation in I558T rERG3 channels. However, replacing the S5-P linker of I558T rERG3 channels, with that of hERG1 channels, did not fully confer the same RPR-attenuated inactivation. Another unidentified region of these ERG proteins may account for the differential allosteric effect of RPR on inactivation. Identification of the structural basis for this difference may provide new insights into the molecular mechanisms of ERG channel inactivation.

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Footnotes:

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Legends for Figures

Fig. 1. Comparison of RPR effects on hERG1 and rERG3 channel currents recorded in *Xenopus* oocytes. A, 3 μ M RPR increases the magnitude and slows deactivation of hERG1 current. Currents shown were elicited with 2 s pulses to 0 mV from a holding potential of -110 mV and tail currents were measured at -70 mV. B - H, Effect of RPR on rERG3 channel current. B, At 3 μ M, RPR failed to enhance the magnitude of rERG3 channel current and produced only a very small but significant slowing of deactivation. Protocol as in A. C, Fully-activated I-V relationships, normalized relative to peak outward current in control, before (control, ■), during (●) and after washout of (△) 3 μ M RPR. D, Voltage dependence of rERG3 inactivation determined from rectification of the fully activated I-V relationship. Symbols as in C. E, RPR affect on rERG3 deactivation at 3 μ M, 10 μ M and 30 μ M. Superimposed mean tail currents recorded at -70 mV were normalized relative to their peak value before (Control) and after addition of 3 ($n = 5$), 10 ($n = 4$) and 30 μ M ($n = 3$) RPR. F, At 30 μ M, RPR produced a partially reversible antagonist effect on rERG3 current. Protocol as in A. G, Fully-activated I-V relationships, normalized relative to peak outward current in control, before (control, ■), and after (●) 30 μ M RPR. H, Voltage dependence of rERG3 inactivation determined from rectification of the fully activated I-V relationship. Symbols as in G. 30 μ M RPR shifted the $V_{0.5}$ by 4.8 ± 0.3 mV ($n = 3$) compared with 1.2 ± 0.7 mV ($n = 3$) using 0.3% DMSO as control. Quantitative analysis of the effects of RPR on rates of deactivation and voltage dependence of inactivation of wild-type channels are summarized in Tables 1 & 2, respectively.

Fig. 2. Alignment of hERG1 and rERG3 amino acid sequences within the putative binding region for RPR. Shaded bars indicate the proposed S5, P (pore) and S6 helices. Residues that differ between sequences are highlighted in bold. One residue (Phe551 hERG1 \approx Met553 rERG3) is in the S4-S5 linker and another (Thr556 hERG1 \approx Ile558 rERG3) is in the cytoplasmic end of the S5 helix. The residues proposed to be important for RPR binding to hERG1 (Perry et

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al., 2007) are indicated by asterisks (*). An additional 16 non-conserved residues are in the extracellular end of the S5 helix and the S5-P linker. Boxed region indicates the sequence of the extracellular end of the S5 helix and the entire S5-P linker that was exchanged from hERG1 into rERG3 to produce the rERG3-S5PL1 chimera.

Fig. 3. The variant residue in the S4-S5 linker does not confer differential sensitivity of hERG1 and rERG3 channel currents to RPR. A, Effect of 3 μ M RPR on F551M hERG1 mutant channel currents expressed in a *Xenopus* oocyte. RPR reversibly enhanced the magnitude and slowed the deactivation rate of F551M hERG1 currents elicited with 2-s pulses to 0 mV from a holding potential of -110 mV. Tail currents were measured at -70 mV. B, RPR significantly slowed F551M hERG1 deactivation. Superimposed tail currents recorded at -70 mV ($n = 5$) were averaged, normalized relative to their peak value before (Control), during and after washout of 3 μ M RPR. C, Fully-activated I-V relationships, normalized relative to peak outward current in control, before (control, ■), during (●) and after washout (△) of 3 μ M RPR ($n = 5$). D, Voltage dependence of rERG3 inactivation determined from rectification of the fully activated I-V relationship (same symbols as in C). E - H, 3 μ M RPR had minimal effect on M553F rERG3 mutant channel currents ($n = 6$). Protocols and analysis were the same as that described in panels A-D. Quantitative analysis of the effects of RPR on rates of deactivation and voltage dependence of inactivation of mutant channels are summarized in Tables 1 & 2, respectively.

Fig. 4. A variant residue in the S5 domain partially confers differential sensitivity of hERG1 and rERG3 channel currents to RPR. A - D, Lack of effect of 3 μ M RPR on T556I hERG1 mutant channel currents expressed in *Xenopus* oocytes. A, Currents were elicited with 2-s pulses to 0 mV from a holding potential of -110 mV. Tail currents were measured at -70 mV. B, RPR does not slow T556I hERG1 current deactivation. Superimposed tail currents recorded at -70 mV ($n = 5$) were averaged and then normalized relative to their peak value before (Control), in the presence of 3 μ M RPR and after washout. C, Fully-activated I-V relationships, normalized

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relative to peak outward current in control, before (control, ■), during (●) and after washout (△) of 3 μ M RPR ($n = 5$). D, Voltage dependence of T556I hERG1 inactivation determined from rectification of the fully activated I-V relationship. E - H, 3 μ M RPR slows deactivation of I558T rERG3 channels ($n = 7$). Protocols and analysis were the same as that described in panels A-D. Quantitative analysis of the effects of RPR on rates of deactivation and voltage dependence of inactivation of mutant channels are summarized in Tables 1 & 2, respectively. I – L, 30 μ M RPR enhances current magnitude and slows deactivation and activation of I558T rERG3 channels ($n = 3$). Protocols and analysis were the same as that described in panels A-D.

Fig. 5. Effect of 3 μ M RPR on M553F/I558T rERG3 channel currents. A, Currents were elicited with 2-s pulses to 0 mV from a holding potential of -110 mV. Tail currents were measured at -70 mV. B, RPR slows M553F/I558T rERG3 current deactivation. Superimposed tail currents recorded at -70 mV ($n = 4$) were averaged and then normalized relative to their peak value before (Control), during and after washout of 3 μ M RPR. C, Fully-activated I-V relationships, normalized relative to peak outward current in control ($n = 4$). D, Voltage dependence of inactivation determined from rectification of the fully activated I-V relationship. Quantitative analysis of the effects of RPR on rates of deactivation and voltage dependence of inactivation of M553F/I558T rERG3 channels are summarized in Tables 1 & 2, respectively.

Fig. 6. S5-P linker of ERG channels is not a structural determinant of RPR sensitivity. A-D, Effects of 3 μ M RPR on rERG3-S5PL1 channels (rERG3 with S5-P domain from hERG1). A, Currents were elicited with 2-s pulses to 0 mV from a holding potential of -110 mV. B, RPR does not slow rERG3-S5PL1 channel current deactivation. Superimposed tail currents recorded at -70 mV ($n = 6$) were averaged and then normalized relative to their peak value before (Control), during and after washout of 3 μ M RPR. C, Fully-activated I-V relationships, normalized relative to peak outward current in control, before (control, ■), during (●) and after washout (△) of 3 μ M RPR ($n = 6$). D, Voltage dependence of inactivation determined from rectification of the fully

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activated I-V relationship. E-H, 3 μ M RPR slightly enhances current magnitude and slows deactivation and activation of I558T rERG3-S5PL1 channels ($n = 7$). Protocols and analysis were the same as that described in panels A-D. Quantitative analysis of the effects of RPR on rates of deactivation and voltage dependence of inactivation of mutant channels are summarized in Tables 1 & 2, respectively.

Fig. 7. Homology model for a single hERG1 subunit, highlighting the residues (shown in stick mode) previously identified (Perry et al., 2007) as important for interaction with RPR. Mutation of residues colored in red reduced or eliminated the effects of 3 μ M RPR on deactivation and inactivation gating. Mutation of residues colored in blue reduced or eliminated effects of 3 μ M RPR on deactivation only. Thr556 is shown in space-fill mode.

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TABLES

Table 1. Summary of effects of RPR on rates of deactivation for hERG1 and rERG3 channels.

		τ_{slow} (ms)	τ_{fast} (ms)	As/(As+Af)	n
WT-hERG1	Control	1210 \pm 81	271 \pm 12	0.49 \pm 0.01	10
	3 μ M RPR	6710 \pm 1450**	265 \pm 13	0.54 \pm 0.02*	
	Control DMSO	1040 \pm 65 1090 \pm 83	254 \pm 13 261 \pm 16	0.46 \pm 0.03 0.48 \pm 0.03	5
F551M hERG1	Control	1310 \pm 114	271 \pm 15	0.49 \pm 0.03	5
	3 μ M RPR	7140 \pm 743**	256 \pm 19*	0.60 \pm 0.04*	
T556I hERG1	Control	881 \pm 50	225 \pm 11	0.39 \pm 0.02	5
	3 μ M RPR	898 \pm 57	216 \pm 9	0.41 \pm 0.02	
WT-erg3	Control	356 \pm 39	92 \pm 5	0.13 \pm 0.03	5
	3 μ M RPR	458 \pm 22*	98 \pm 4*	0.13 \pm 0.02	
	Control DMSO	543 \pm 19 536 \pm 33	86 \pm 1 78 \pm 3	0.05 \pm 0.01 0.04 \pm 0.01	3
M553F erg3	Control	233 \pm 29	57 \pm 7	0.16 \pm 0.03	6
	3 μ M RPR	272 \pm 30*	60 \pm 7	0.16 \pm 0.03	
I558T erg3	Control	472 \pm 41	116 \pm 6	0.20 \pm 0.02	5
	3 μ M RPR	5020 \pm 615**	119 \pm 5	0.39 \pm 0.03**	
M553F/I558T erg3	Control	504 \pm 13	101 \pm 8	0.16 \pm 0.03	4
	3 μ M RPR	3550 \pm 92**	107 \pm 11	0.37 \pm 0.04*	
erg3-S5PL1	Control	300 \pm 24.	80 \pm 3	0.17 \pm 0.02	6
	3 μ M RPR	413 \pm 25**	78 \pm 0.01	0.14 \pm 0.01**	
I558T erg3- S5PL1	Control	430 \pm 24	106 \pm 5	0.24 \pm 0.02	7
	3 μ M RPR	4000 \pm 269**	109 \pm 5	0.25 \pm 0.01	

* P <0.05; ** P <0.005

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Table 2. Effects of RPR on voltage dependence of inactivation for hERG1 and rERG3 channels.

	Inactivation $V_{0.5}$ (mV)			$\Delta V_{0.5}$ RPR	% change $I_{tail, FA}$ (0mV)	<i>n</i>
	Control	3 μ M RPR	Washout			
WT-hERG1	-46.3 \pm 1.8	-33.9 \pm 2.2	-43.3 \pm 2.0	12.5 \pm 1.0**	63 \pm 9	10
<i>DMSO Control</i>	-43.2 \pm 1.9	-43.1 \pm 2.4	NA	0.1 \pm 0.9	6 \pm 6	5
F551M hERG1	-39.6 \pm 2.2	-28.9 \pm 2.9	-39.4 \pm 1.9	10.6 \pm 1.0**	64 \pm 17	5
T556I hERG1	-47.5 \pm 2.1	-45.8 \pm 2.4	-48.1 \pm 2.6	1.8 \pm 0.5*	4 \pm 1	5
WT-erg3	-36.9 \pm 2.7	-36.8 \pm 2.8	-37.5 \pm 2.8	0.1 \pm 0.4	-4 \pm 2	5
<i>DMSO Control</i>	-35.4 \pm 2.2	-34.2 \pm 2.9	-36.5 \pm 1.8	1.2 \pm 0.7	-4 \pm 3	3
M553F erg3	-30.5 \pm 4.4	-27.8 \pm 4.3	-28.8 \pm 4.8	2.8 \pm 0.9	9 \pm 6	6
I558T erg3	-40.8 \pm 1.6	-36.4 \pm 1.9	-41.7 \pm 2.0	4.4 \pm 0.9*	9 \pm 4	5
M553F/I558T erg3	-44.1 \pm 3.7	-38.0 \pm 4.4	-44.3 \pm 3.4	6.1 \pm 1.4*	14 \pm 8	4
erg3-S5PL1	-43.9 \pm 1.9	-40.1 \pm 2.1	-44.1 \pm 2.1	3.8 \pm 1.1*	12 \pm 4	6
I558T erg3-S5PL1	-51.9 \pm 3.4	-45.2 \pm 3.5	-51.5 \pm 3.1	6.7 \pm 0.8**	22 \pm 3	7

* P <0.05; ** P <0.005

Figure 1

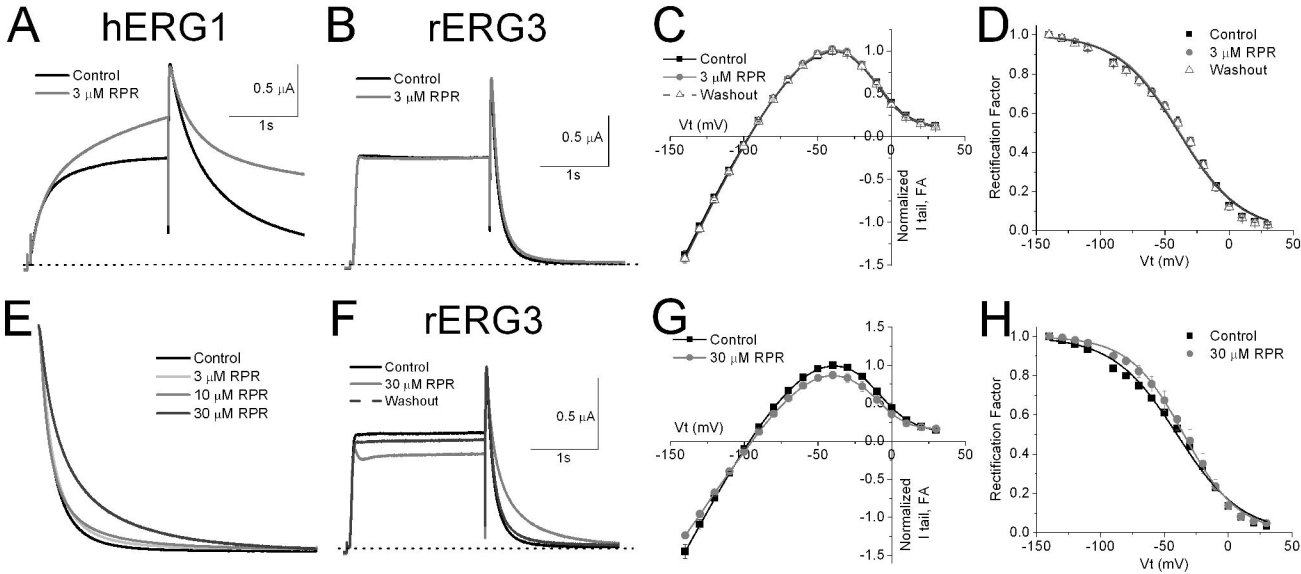


Figure 2

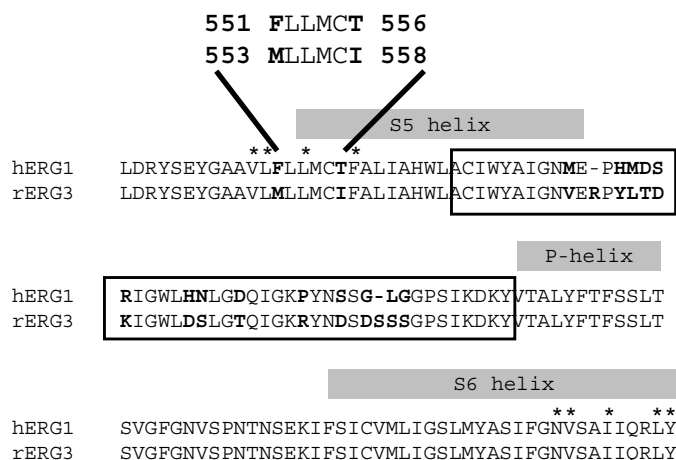


Figure 3

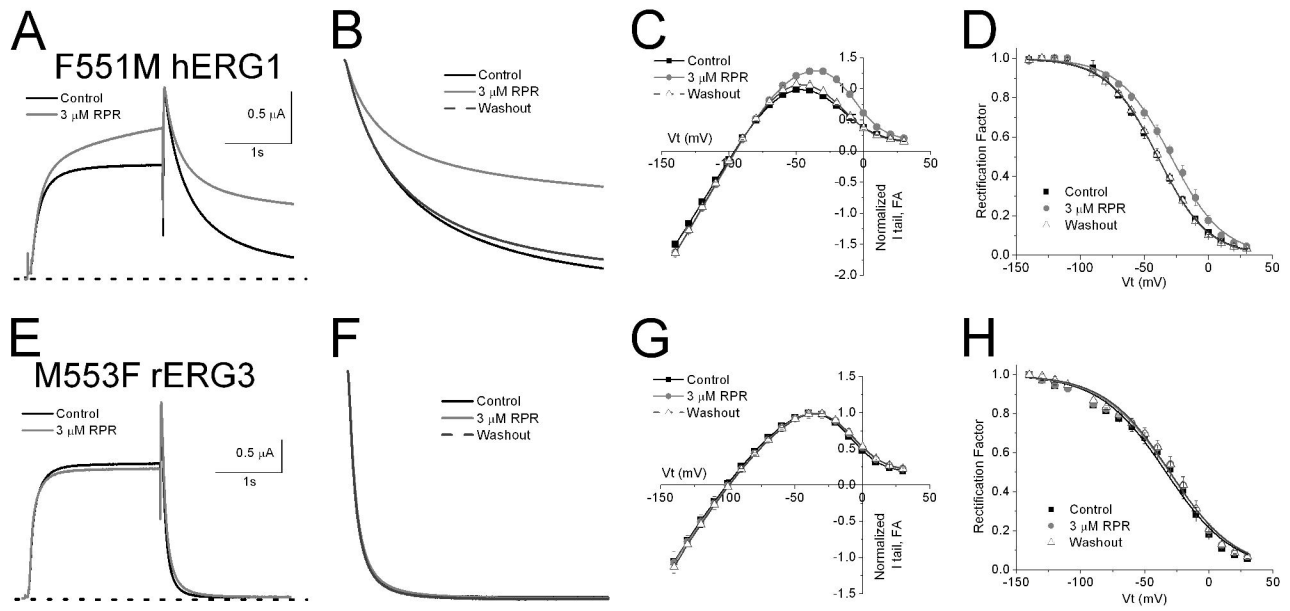


Figure 4

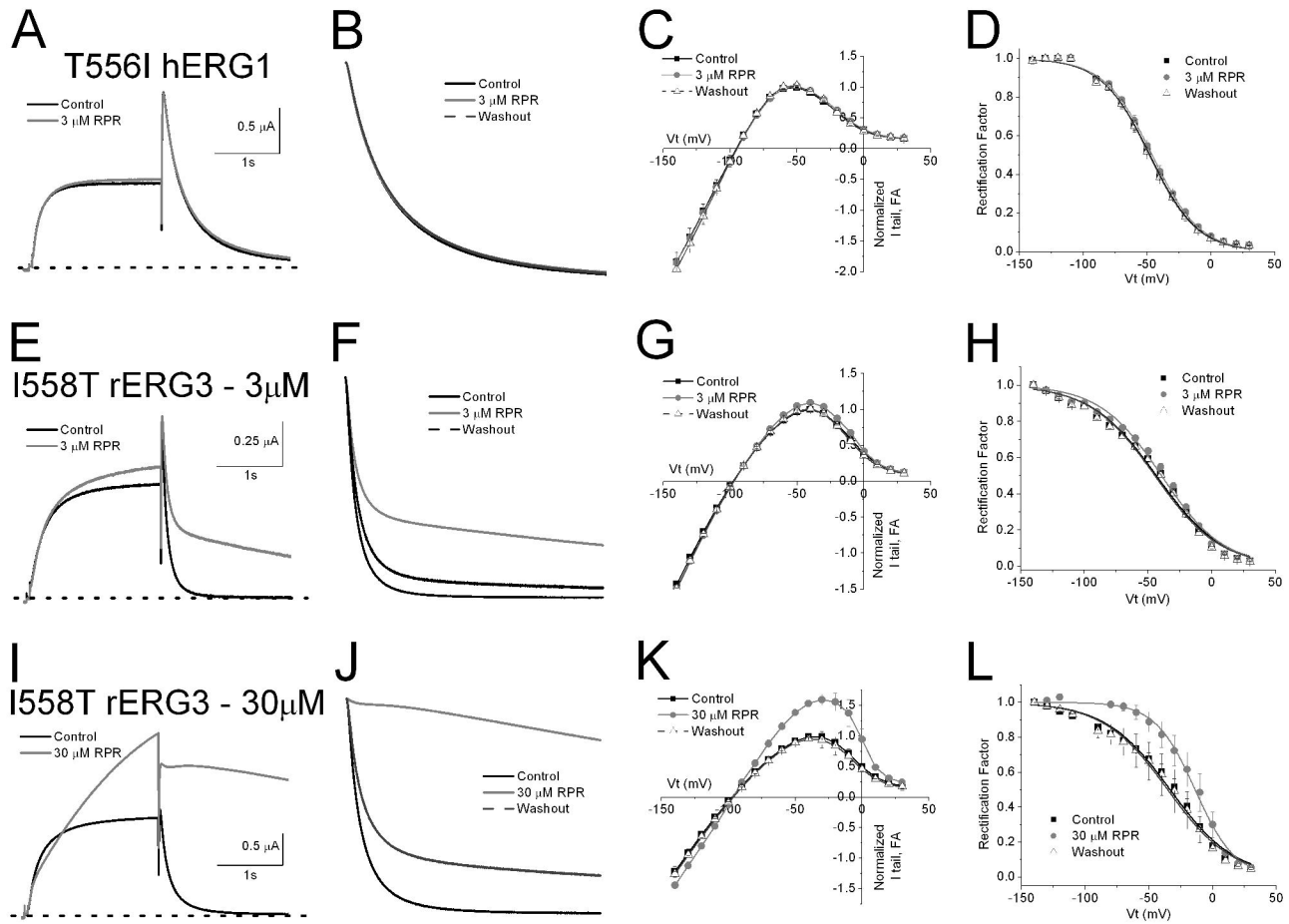


Figure 5

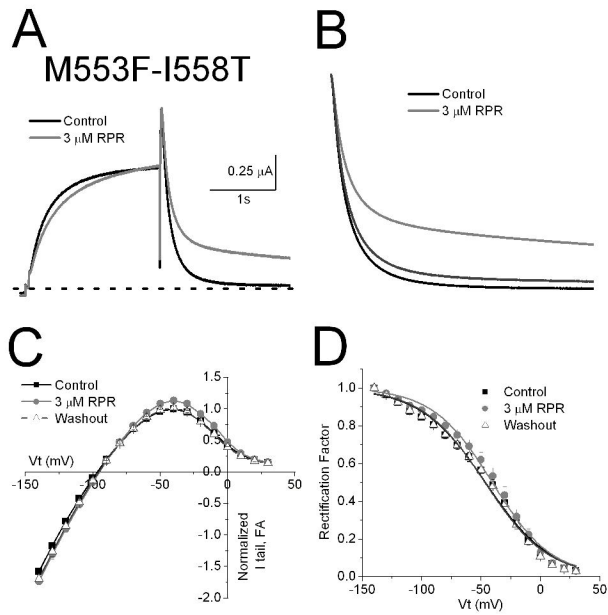


Figure 6

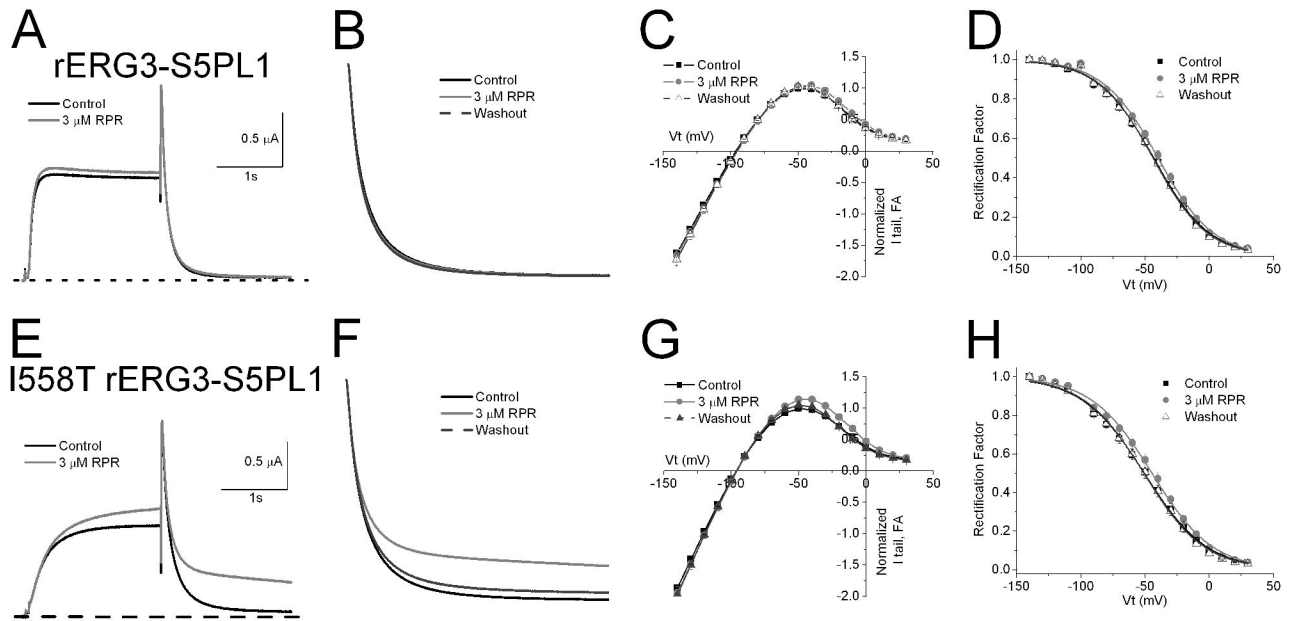


Figure 7

